## REDBOOK 1971 PART III

SELECTED PAPERS FROM THE 1971 ANNUAL MEETING

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## Note

REDBOOK 1971 appears in 3 books. The first book contains Part 1, Proceedings of the Standing Committee on Research and Statistics. The second book contains Part II, Reports on Researches in the ICNAF Area in 1970. The third book contains Part III, Selected Papers from the 1971 Annual Meeting.

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1971

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SECTION A OCEANOGRAPHY

1. Temperatures and salinities in the eastern Newfoundland area in $1970^{1}$
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## Introduction

The 6 standard monitoring sections, taken each year at approximately the same dates in July-August across the Labrador Current east of Newfoundland, were occupied by the Cape Freels. Station 27, off Cape Spear, was occupied monthly or more of ten during the year. The section temperatures are compared with the lowest, average, and highest temperatures at each depth at each station in 1951-65 in July-August (unpublished) and also with temperatures in 1969 (Templeman, 1970) and for the St. John's-Flemish Cap section for 196669 (Templeman, 1967, 1968, 1969, 1970). Salinities are also compared with those in the above years.

Sections across the Labrador Current in July-August

## Temperatures

In the Seal Islands, Labrador, section across Hamilton Inlet Bank (Fig.1), surface temperatures were similar to the long-term average. The lowest temperatures at mid-depths near the coast were similar to the lowest of the 1951-65 period and were slightly lower than in 1969. The volume of water with temperatures below $-1^{\circ} \mathrm{C}$ was above average. Temperatures in the more offshore portion of the cold central area of the Labrador Current were lower than the average of the 1951-65 period but higher than the lowest recorded and the 1969 temperatures. Temperatures of the bottom water in Hawke Channel were slightly higher than the long-term average and higher than in 1969. The temperature at the crest of Hamilton Inlet Bank was the highest recorded. Temperatures of the deep water seaward of Hamilton Inlet Bank at Stations 55A and 56 were similar to the highest recorded averages and considerably higher than in 1969, and at Station 57, the most seaward station, deep-water temperatures from 200 to $1,000 \mathrm{~m}$ were higher than any previously encountered, and between 200 and 600 m considerably higher than in 1969.

In the section off Cape Bonavista and southward to the northern Grand Bank (Fig. 2), surface temperatures of the Cape Bonavista portion were slightly below the long-term average and below those of 1969 , except at the most coastal station, and mainly slightly below 1969 in the Grand Bank section. Temperatures in the inshore central core of the Labrador Current were slightly lower than the 1951-65 average and approximately similar to those of 1969, and the volumes of water with temperatures of below $-1^{\circ} \mathrm{C}$ and $0^{\circ} \mathrm{C}$ were close to the average. The lowest temperature of the shoreward cold water and the lowest temperatures of the eastward-flowing branch of the Labrador Current near the Grand Bank were slightly below those of 1969. Except for the bottom temperatures at Stations 46 and 47 , temperatures from 300 to 1,000 m were the highest recorded and considerably higher than in 1969.

In the St. John's-Flemish Cap section (Fig. 3), surface temperatures were slightly below average and approximately similar to those of 1969, except at

[^0]the inshore Station 27 where the surface temperature was high. The lowest temperatures of the shoreward central core of the Labrador Current were similar to the 1951-65 average and below those of 1969. The amount of water below $0^{\circ} \mathrm{C}$ was also close to the average but the temperature at the bottom at Station 27 in the Avalon Channel was the second highest recorded. Temperatures at the cold-water core of the eastern branch of the Labrador Current were close to the average and the extent of water with temperatures below $0^{\circ} \mathrm{C}$ a little greater than the average. Water temperatures in the upper layers above 150 m were average. Temperatures from 400 m downward in the Flemish Channel and those from 200 to $1,000 \mathrm{~m}$ seaward of Flemish Cap were among the highest and sometimes, including the $200-800-\mathrm{m}$ levels at Station 42A, the highest recorded in the period 1951-70.

In the section from St. John's to the southeast slope of the Grand Bank (Fig. 4), the lowest temperatures in the coldest part of the Labrador Current, near shore, were similar to the lowest recorded. The temperature near bottom at Station 27 and temperatures over the surface of the Grand Bank were near but mainly slightly above the average. The distribution of the coldest water of the eastern branch of the Labrador Current, which usually hugs the eastern slope of the Grand Bank, was highly unusual, swinging away from the bank and with a warmer water border fringing the bank. Temperatures at the centre of this cold water were average. Temperatures at the mast eastern station (33F) were above average but lower than those of 1969.

In the section extending along the southwestern edge of the Grand Bank at about 75 m (Fig. 5), surface temperatures were close to the 1951-65 average and higher than in 1969. Temperatures in the cold water near bottom in the Haddock Channel were close to the highest of the 1951-65 period but lower than in 1969. Bottom temperatures over the surface of the Grand Bank were close to the average for 1951-65 and somewhat similar to those of 1969. In the cold eastern branch of the Labrador Current at the eastern edge of the bank, temperatures were higher and the volume and depth of water with temperatures below $0^{\circ} \mathrm{C}$ and below $2^{\circ} \mathrm{C}$ much less than usual. The temperatures of the water fringing the Grand Bank on its eastern slope were by far the highest recorded and this warm water intrusion separated the low temperature and the median temperature water of the Labrador Current.

In the section at 275 m along the southwestern slope of the Grand Bank and extending to St. Pierre Bank (Fig. 6), surface temperatures were above the average for the 1951-65 period and considerably higher than in 1969. Temperatures of the intermediate cold Labrador Current water from the Haddock Channel were below average and, in the coldest water of the eastern branch of the Labrador Current, above average and almost as high as the highest recorded. Temperatures of the higher temperature deep slope water between these two branches of the Labrador Current were well above average. The bottom temperatures were mainly above average, all at Stations 10, 15, and 19 higher than any previously encountered in the 1951-65, 1969 period. As in the previous section (and at the same station, 26F) a branch of the Labrador Current with intermediate temperatures, which could be a return current passing northward, was apparent to the east of the slope separated from the main eastern branch by a warm water intrusion.

## Salinities

In these comparisons, salinities from 20 m to the surface are omitted
because they are too much influenced by local and temporary precipitation and runoff to be very useful in year to year comparisons.

In the Seal Island section of $f$ southern Labrador (Fig. 1), at all stations except 55A, where there was some upwelling, salinities in the upper 100 m , at Station 53 the upper 150 m , and at Station 57 the upper 200 m were either the lowest or generally among the lowest 2 to 5 out of 16 or 17 years of records. Salinities were also in the lower half of records for the period at Station 55 for $150-250 \mathrm{~m}$, at Station 55 A for 50 m to 300 m , at Station 56 for 200 m , and at Station 57 for $250-500 \mathrm{~m}$. The salinities at $200-250 \mathrm{~m}$ at Station 53 were close to the average. Seaward from this station, salinities were above average and usually in the upper quarter for the period at Station $54,100 \mathrm{~m}$ and deeper; Station $55,300 \mathrm{~m}$ and deeper; Station $55 \mathrm{~A}, 400 \mathrm{~m}$ and deeper; at Stations 56 and 57 at 250 m and deeper. Salinities at Station 55 A ( 500 m ), at Station $56(300-500 \mathrm{~m})$ and at Station $57(600-1,000 \mathrm{~m})$ were the highest in the records for 6-17 years.

In the section off Cape Bonavista and southward to the northern Grand Bank (Fig. 2), at all depths from 30 m and greater at Stations 43 and 44, depths down to 150 m at Stations 45 and 49, to 250 m at Stations 46 and 47, to 100 m at Station 48, and to 200 m at Station 50 were either the lowest encountered in the period under comparison ( 18 of 40 ) or were in the lowest quarter. Salinities in the deep water: bottom at Station 45 and 47, 250 m to bottom at Stations 48 and 49, and 400 to $1,000 \mathrm{~m}$ at Station 50 , were either the highest or equal to the highest on record ( 9 of 12) or in the highest quarter. Similarly high deep-water salinities and low upper-water salinities prevailed in the northern Grand Bank portion of the section for which longterm comparisons are not available.

In the section from St. John's to Flemish Cap (Fig. 3), salinities at Station 27 ( $30-125 \mathrm{~m}$ ), Station 28 ( $30-100 \mathrm{~m}$ ), Stations 29, 36 and 37 ( $30-50 \mathrm{~m}$ ), Station 35 ( 30 m to bottom), Station $37 \mathrm{~A}(30 \mathrm{~m}$ ), Stations 38 and 40 ( $30-150 \mathrm{~m}$ ), Stations 39 and 42A ( $30-100 \mathrm{~m}$ ), and Stations 41 and 42 ( $30-75 \mathrm{~m}$ ) were all below average; 9 ( 8 at 30 m ) of 44 the lowest encountered for the period 1951-70, and 33 of 44 in the lowest quarter. Deep-water salinities at Station 27 ( 150 m to bottom), Station 28 (bottom), Stations 34, 36 and 37 ( 75 m to bottom), Station 37A ( 50 m to bottom), Station 38 ( $200-800 \mathrm{~m}$ ), Station 39 ( $150-750 \mathrm{~m}$ ), Station 40 ( 200 m to bottom), Stations 41 and 42 ( 100 m to bottom), and Station 42A ( 150 m to bottom) were above average, 33 of 53 the highest or equal to the highest encountered for the period $1951-70$ and 46 of 53 in the highest quarter. The deep-water salinities above $35 \%$ at Stations 38-42A, in Flemish Channel and on the eastern slope of Flemish Cap, were the highest in our records.

In the section from St. John's to the southeast slope of the Grand Bank (Fig. 4), salinities at Stations 27-32A (all depths from 30 m downard), Stations $33(30-75 \mathrm{~m}), 33 \mathrm{~A}(30-250 \mathrm{~m}), 33 \mathrm{~B}(30-400 \mathrm{~m}), 33 \mathrm{D}(30-250 \mathrm{~m})$ and 33 F ( $30-150 \mathrm{~m}$ ) were below average, 34 of 55 the lowest encountered for the period and 49 of 55 in the lowest quarter. Salinities in the deeper water of the eastern slope: Station 33 ( 100 m to bottom), Stations 33A and 33D ( 300 m to bottom), Station 33B (bottom), and Station 33F ( 200 m to bottom) were mostly above average, 6 of 20 the highest or equal to the highest encountered for the period and 10 of 20 in the highest quarter.

In the section at about 75 m extending along the southwestern slope of
the Grand Bank (Fig. 5), salinities at all depths from 30 m downward at Stations 20B-26A and at depths of $30-200 \mathrm{~m}$ at Station 26 F were the lowest (12 of 38) or among the lowest ( $1 / 8$ to $1 / 4$ ) of the salinities encountered in the period under comparison. At depths of from 30 m to bottom at Station 26 B , 50 m to bottom at Station 26D, and from 300 to 600 m at Station 26 F , salinities were equal to the highest or were the highest recorded (13 of 20) or were in the highest quarter recorded.

In the section at 275 m along the southwestern slope of the Grand Bank and extending to St. Pierre Bank (Fig. 6), salinities down to 100 m at Station 10 , to 150 m at Station 13 , to 75 m at Station 18 and at 30 m at Stations 16 , 17 and 19 were usually either the lowest ( 7 of 15) or in the lowest quarter of salinities in the period under comparison. Salinities from 150 m to bottom at Stations 10, 15 and $17,200 \mathrm{~m}$ to bottom at Stations 13 and 16, and from 50 m to bottom at Station 19 were either the highest or equal to the highest on record for the period under comparison (11 of 24) or usually among the highest quarter of the records for the period. Data for Stations 26 D and 26 F of this section are the same as in Fig. 5.

The most noteworthy features of the 1970 sections were the warmer water and especially the unusually high salinities in the deeper water at the eastern slope of the continental shelf and banks. Judging from the location of occurrence of high salinities in the sections (Figs. 1-6), and from Dietrich (1965, fig. 11) there was an unusual degree of intrusion of Atlantic water into the eastern and deeper parts of the Labrador Current on the northeastern and eastern slopes of the Grand Bank. Also noteworthy were the unusually low salinities of the upper water layers and the separation of the cold core of the Labrador Current from the Grand Bank to the east of the Southeast Shoal. The latter must be highly unusual as it is the first time that it has occurred in 17 years of records since 1951.

Station 27, 1970
In Station 27 off Cape Spear (Fig. 7), except for January, surface temperatures throughout the year were above the 1950-62 average (Templeman, 1965). Also, the highest surface temperatures in July-August were well above those in 1969 (Templeman, 1970). Bottom temperatures except for February and December were above the 1950-62 average and were mostly similar to those of 1969. At intermediate levels, the lowest water temperatures, below $-1^{\circ} \mathrm{C}$, were below the 1950-62 average and below those of 1969 and were evidently not produced locally but were brought in by the Labrador Current. The coldest water came through Station 27 in August and there was a much greater than usual persistence of a large volume of water with very low temperatures up to the beginning of December.

Upper layer salinities in late summer and in autumn were well below and surface salinities throughout the year usually below those of 1969. Otherwise, they were generally similar to those of 1969 , except that water with a salinity below $33 \%$ o sometimes extended to the bottom in 1970 but not in 1969.

## Acknowledgements

I am grateful to Mr A.G. Kelland, hydrographic technician at the St. John's Station and to Mr L.N. Cluett for their interest in gathering data for
this paper, also to the scientists and technicians of the St. John's Station who have taken hydrographic observations on Station 27 and on the various sections.

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Fig. 1. Temperature ( ${ }^{\circ} \mathrm{C}$ ) above and salinity ( ${ }^{\circ} \%$ ) below, Seal IslandHamilton Inlet Bank section, 2-3 August 1970.


Fig. 2. Temperature ( ${ }^{\circ} \mathrm{C}$ ) above and salinity ( $\%$ oo) below, for section off Cape Bonavista, and southward to northern Grand Bank, 28 July-1 August 1970.


Fig. 3. Temperature ( ${ }^{\circ} \mathrm{C}$ ) above and salinity ( ${ }^{\circ} / \%$ ) below, St. John's-Flemish Cap section, 27-29 July 1970.


Fig. 4. Temperature ( ${ }^{\circ} \mathrm{C}$ ) above and salinity ( ${ }^{\circ} / \circ \circ$ ) below, St. John's-SE slope Grand Bank, 17-19 August 1970.


Fig. 5. Temperature ( ${ }^{\circ} \mathrm{C}$ ) above and salinity ( $\% / \%$ ) below, Green Bank-SE Grand Bank, 20-23 August 1970.


Fig. 6. Temperature ( ${ }^{\circ} \mathrm{C}$ ) above and salinity ( $\%$ ) below, SW slope Grand Bank-St. Pierre Bank, 20-23 August 1970.


Fig. 7. Temperature ( ${ }^{\circ} \mathrm{C}$ ) above and salinity ( $\%$ 。) below, January 1970 to January 1971, from surface to bottom at Station 27 (see Figs. 3, 4 inset), 2 nautical miles of $f$ Cape Spear near St. John's.
2. Hydrographic conditions off West Greenland during $1970^{1}$
by Frede Hermann
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A number of standard sections off West Greenland (Fig. 1) were worked from the R/V Adolf Jensen from May to December 1970. Section I off Frederikshaab was worked in May and October. Section II across Fyllas Bank was worked in May, June, twice in July and in September, October, November and December. Sections III and IV over Lille Hellefiske Bank and Store Hellefiske Bank were worked in May and July.

Ice conditions were rather severe in 1970. In both April and in July/ August, the "storis" reached from Cape Farewell to a position north of Godthaab during its maximum extension. In April, the "storis" was relatively scattered and in a period with northerly wind in May, it disappeared from the bank area off the middle part of West Greenland.

Temperature sections are shown in Figs. 2-8. As in the two preceding years, temperatures were unusually low in the upper 100 m . In the Fyllas Bank section, negative temperatures were measured as late as September off the western slope of the Bank.

The temperature deviation from the mean value for the period 1950 to 1966 (Hermann, 1967) for the station at $63^{\circ} 53^{\prime} \mathrm{N}-53^{\circ} 22^{\prime} \mathrm{W}$ west of the slope of Fyllas Bank in July is given in Table 1 and the corresponding salinity anomaly is given in Table 2.

Table 1. Temperature anomalies west of Fyllas Bank in July.

| Depth Interval <br> $(\mathrm{m})$ | Mean Temperature <br> $(1950-66)$ | $\Delta \mathrm{T}$ <br> (July 1970) |
| :---: | :---: | :---: |
| $0-50$ | 2.07 | -1.31 |
| $50-100$ | 1.33 | -1.12 |
| $100-200$ | 1.85 | -1.73 |
| $200-300$ | 2.88 | -2.06 |
| $300-400$ | 3.79 | -1.19 |
| $400-500$ | 4.22 | -0.08 |
| $0-500$ | 2.89 | -1.2 |

[^1]Table 2. Salinity anomalies west of Fyllas Bank in July.

| Depth Interval <br> $(\mathrm{m})$ | Mean Salinity <br> $(1950-66)$ | $\Delta \mathrm{S}$ <br> (July 1970) |
| :---: | :---: | :---: |
| $0-50$ | 33.29 | -0.46 |
| $50-100$ | 33.65 | -0.50 |
| $100-200$ | 34.00 | -0.56 |
| $200-300$ | 34.39 | -0.60 |
| $300-400$ | 34.67 | -0.30 |
| $400-500$ | 34.81 | +0.03 |
| $0-500$ | 34.27 | -0.38 |

In the upper 400 m the temperature was between $1^{\circ}$ and $2^{\circ}$ lower than the mean value and the salinity was between $0.30^{\circ} \%$ and $0.6 \%$ lower than the mean value. The low temperature and salinity indicate a strong inflow of polar water to the West Greenland area.

Figure 9 shows the 5-year running averages of the surface temperatures in the West Greenland area ( $A_{1}$ ) and South Greenland area (B) based on the surface anomalies by Jens Smed for the years up to 1969 (Smed, up to 1968).

The strong decrease in temperature in the last half of the sixties is rather alarming.

The low temperature in the surface layers will probably affect the survival of the cod larvae. The 1970 cod year-class will probably be very small in the northwest Greenland area.

The author is grateful to Jens Smed for placing the unpublished temperature anomalies for 1969 at his disposal.

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Frederiksháb 8/10 70 TO
Fig. 2. Temperature sections off Frederikshaab (I) for May and October 1970.


Fig. 3. Temperature sections across Fyllas Bank (II) in May and June 1970.


Fig. 4. Temperature sections across Fyllas Bank (II) in July 1970.


Fig. 5. Temperature sections across Fyllas Bank (II) in September and October 1970.


Fig. 6. Temperature sections across Fyllas Bank (II) in November and December 1970.





Fig. 8. Temperature sections across Store Hellefiskebanke (IV) in May and July 1970.

Fig. 9. Sea surface temperature anomalies, 5-year running mean. $\begin{array}{ll}A_{1} \text { (solid line): } & \text { West Greenland area. } \\ \text { B (dotted line): } & \text { South Greenland area. }\end{array}$
3. Environmental studies in ICNAF Div. 3P and 4 V in spring $1970^{1}$
by R. L'Herrou and J.P. Minet
ISTPM, Saint-Pierre et Miquelon
In spring 1970, R/V Thalassa completed a cruise in ICNAF Div. 3P and 4V to study the stocks of fish. During the cruise, various hydrographic observations were made. Temperatures were taken with bathythermograph from $0-300 \mathrm{~m}$ after each trawl haul. From 13-17 April, temperature and salinity measurements were made over a network of 28 stations. Conjointly, vertical plankton hauls were made.

## A. Hydrographic conditions

I. Distribution of surface and near bottom temperature

At the surface (Fig. 1), cold waters of $1-2^{\circ} \mathrm{C}$ flow out of the Gulf of St. Lawrence and extend along the coasts of Cape Breton Island. A lobe of $2-3^{\circ} \mathrm{C}$ water spreads to the centre of the Laurentian Channel with an overlay of $5-6^{\circ} \mathrm{C}$ water rising from the bottom to the western slopes of Saint Pier re Bank. In addition, on the eastern plateau, $2-3^{\circ} \mathrm{C}$ water extends from the Newfoundland banks.

Near bottom (Fig. 2), as at the surface, there is a progression of Gulf of St. Lawrence waters along the shores and banks of Nova Scotia. South of Burin Peninsula, two cold lobes $\left(0^{\circ} \mathrm{C}\right)$, resulting from mixing of waters from Labrador and Newfoundland, are found near the eastern part of Saint Pierre Bank and give rise to a very strong thermal gradient ( $1^{\circ}-7^{\circ} \mathrm{C}$ ) along the western slopes of the Bank.

## II. Hydrographic sections - temperatures and salinities

Five sections containing a total of 28 hydrographic stations were occupied at the positions listed in Table 1 and referred to in Fig. 3.

At each station, Nansen reversing bottles were lowered to standard depths. Two hundred and fifty-eight temperature samples were taken, using Richter $\&$ Wiese protected and non-protected thermometers. Temperature corrections were made on board, using the Culbertson method. For salinity studies, 258 sea water samples, collected in duplicate, were analyzed in the laboratory using the Auto-Lab induction salinometer (Sydney, Australia).

1. Section I (Stations 04-08) - Burin Peninsula to the southern tip of

Continental shelf waters consist of a surface layer which is slightly warmer in early spring ( $1.15-1.25^{\circ} \mathrm{C}$ ) and an intermediate cold water layer of Labrador origin ( $0.88-0.95^{\circ} \mathrm{C}$ and $32.00-32.20^{\circ} \%$ ) which spreads to Saint Pierre Bank without overflowing from the slope. In the channel separating the Newfoundland coast and the northern slope of Saint Pierre Bank, there is a residue of water which was formed in the winter with below $0^{\circ}$ temperatures $\left(-0.13\right.$ to $\left.-0.26^{\circ} \mathrm{C}\right)$ and higher salinities (32.20-32.50\% o).

[^2]At approximately 140 m ，a lobe of water of warmer and more saline gradient than surface and underlying waters $\left(3.78^{\circ} \mathrm{C}\right.$ and $33.59 \%$ ）appears on the south shores of Saint Pierre Bank．Moreover，this water，arising from a mixture of intermediate slope and surface waters，sends a second lobe to the $75-\mathrm{m}$ level at Station 08.

From $200-280 \mathrm{~m}$ ，there is a very strong temperature gradient（ 5 degrees in 80 m ）and salinity gradient（ $33.04-34.11 \%$ ）which becomes slope water clearly influenced by warm open sea waters $\left(7.0^{\circ} \mathrm{C}\right.$ and $34.11^{\circ} \%$ at 300 m ）．

2．Section II（Stations 10－14－South Banquereau Bank to South Saint Pierre Bank（Figs．5A and B）

Again，a cold layer extends over Saint Pierre Bank（Section I）and here， overflows to the southeast slope of the Bank（ -0.11 to $0.99^{\circ} \mathrm{C}$ at 75 m ）． Parallel to this are two lobes of intermediate slope water influenced by the Atlantic：the first is on the southwest slopes of Banquereau Bank $\left(7.0^{\circ} \mathrm{C}\right.$ and $34.72 \%$ ）at 200 m ；the second is on the southeast slopes of Saint Pierre Bank $\left(7.84^{\circ} \mathrm{C}\right.$ and $34.76^{\circ} \%$ ）at 220 m and extends to the centre of the Laurentian Channel（ $8.07^{\circ} \mathrm{C}$ and $34.60^{\circ} \%$ ）at 100 m ．This warm water，therefore，enters the Laurentian Channel on the left and especially on the right of the section， thus isolating a mixture of colder water between 250 m and the bottom（6．00－ $4.50^{\circ} \mathrm{C}$ and $34.65-34.84^{\circ} \%$ 。）．

3．Section III（Stations 15－24）－Cape Breton to South Banquereau Bank （Figs．6A and B）

Lying near the Cape Breton coast（Stations 23 and 24）is very cold and much less saline water（ -1.00 to $0.0^{\circ} \mathrm{C}$ and $31.00-31.20^{\circ} \%$ 。）produced from the Gulf of St．Lawrence and spreading southeast．An intermediate cold layer， warm at the surface，extends over the shelf to the shores at 100 m ，without overflowing the slope（ $2.00-4.00^{\circ} \mathrm{C}$ and $32.50-32.70^{\circ} \%$ 。）．Between 150 and 250 m warmer intermediate slope water is observed，originating from the open sea and washing the southern slopes of Banquereau Bank $\left(6.00-7.00^{\circ} \mathrm{C}\right.$ and $34.00-$ $34.50 \%$ o）；at $300-2,500 \mathrm{~m}$ ，this water gives way to deep slope water whose temperature $\left(4.50^{\circ} \mathrm{C}\right.$ at $300 \mathrm{~m}, 3.50^{\circ} \mathrm{C}$ at $2,000 \mathrm{~m}$ ）decreases and salinity $\left(34.60^{\circ} \%\right.$ at $250 \mathrm{~m}, 34.92 \%$ at $300-2,500 \mathrm{~m}$ ）increases with depth．

4．Section IV（Stations 25－29）－Across Laurentian Channel，north of Misaine Bank to north of Saint Pierre Bank（Figs．7A and B）

As in Section III，a cold lobe $\left(1.00^{\circ} \mathrm{C}\right.$ and $32.00^{\circ} \%$ ）produced by the Gulf of St．Lawrence from a mixture of Labrador water and St．Lawrence estuary water spreads to the shores of Nova Scotia．In addition，water originating from the Newfoundland Banks is found on Saint Pierre Bank and in the middle of the Laurentian Channel，at 100 m ．Above this cold intermediate water resulting from a successive mixture，there is a slightly warmer surface layer originating from the Newfoundland coast．Beneath this cold intermediate layer， the intermediate slope water is clearly evident on Saint Pierre slopes，at 200－ $250 \mathrm{~m}\left(6.95^{\circ} \mathrm{C}\right.$ and $34.50^{\circ} \%$ 。）This advance of water clinging to Saint Pierre Bank draws an accumulation of overlying waters（ $2.00-5.00^{\circ} \mathrm{C}$ and $33.00-34.00^{\circ} \%$ ） from the Nova Scotia shores．From 300 m to the bottom of the Laurentian Channel，deep slope waters are always present $\left(5.14^{\circ} \mathrm{C}\right.$ at 300 m and $4.54^{\circ} \mathrm{C}$ at bottom； $34.66 \%$ os 300 m and $34.88 \%$ 。 at bottom）．
5. Section V (Stations 30-34) - Cabot Strait (Figs. 8A and B)

This section shows more clearly again the relative position of the different formations of water. As in Section IV, the outflow of Gulf of St. Lawrence water along the coast of Cape Breton Island is clearly apparent $\left(-1.00\right.$ to $0.0^{\circ} \mathrm{C}$ and $3.00-31.50^{\circ} \%$ ). Along the south coast of Newfoundland, to the right of the section, the $1.50^{\circ} \mathrm{C}$ isotherm and $31.80^{\circ} \%$ and $32.00^{\circ} \%$ isohalines show an accumulation of mixed water (Labrador and Newfoundland) which progresses towards the Gulf of St. Lawrence. Under the cold intermediate layer ( $1.50-3.00^{\circ} \mathrm{C}$ and $32.50-33.50^{\circ} \%$ ), the advance of the intermediate slope water makes its influence felt up to this level, and also along the shores of Newfoundland (5.00-6.00 ${ }^{\circ} \mathrm{C}$ and $34.00-34.50^{\circ} \%$ ). Below 300 m , the deep slope waters (5.11-4.57 ${ }^{\circ} \mathrm{C}$ and $34.72-34.89^{\circ} / \circ 0$ ) are again found.

## B. Zooplankton distribution

Vertical plankton hauls were made with a 72-cm diameter Hensen net which is raised from the bottom to the surface at $1 \mathrm{~m} / 3 \mathrm{sec}$. A total of 133 stations were occupied. The volume of total plankton collected in each sample along with an inventory of eggs and larvae of fish collected were recorded.

## I. Quantitative zooplankton distribution

Plankton volume was measured by displacement of filtered sea water (in $\mathrm{cm}^{3} / 10 \mathrm{~m}^{3}$ ) using a distribution card (Fig. 9) which shows that, in the spring, all sections studied showed light concentrations of plankton. However, concentrations are heavier in some zones of the southern part of Saint Pierre, Banquereau and Misaine Banks.

In comparing plankton distribution with surface (Fig. 1) and near bottom (Fig. 2) temperatures, it is noted that zones with heavier concentrations are those with $0^{\circ}-3^{\circ} \mathrm{C}$ temperatures, while zones with lighter concentrations are those with $3-6^{\circ} \mathrm{C}$ temperatures on the bottom and $3-4^{\circ} \mathrm{C}$ at the surface.
II. Inventory of fish eggs and larvae

The following table lists the eggs and larvae of fish which have been identified in the plankton samples:


No distribution table is given for the larvae of American plaice (Hippoglossoides $p l$. platessoides) as their number is too small. The distribution of eggs (Fig. 10) shows that the spawning, although of little importance, is found spread along the bank areas on either side of the Laurentian Channel in which there are few, if any, eggs found. The limited number of larvae taken confirms that the spawning from March-May, with a maximum in April, is of small magnitude.

Cod eggs are also scarce. This is explained by the fact that April is the end of the spawning period. On the other hand, only three larvae were taken which suggests that recruitment was very weak in the area studied.

Redfish larvae are fairly well represented if one considers that it is only the beginning of the reproduction period for both the marinus and mentella types of redfish.

## Conclusion

Although these observations contribute to a better environmental knowledge of the area studied, they are nevertheless insufficient since they are limited to one time of the year.

In order to define all phenomena and to follow their development in time, a seasonal study will be undertaken beginning in July 1971 by R/V Cryos which is attached to the laboratory of 1'Institut des Peches Maritimes at Saint Pierre and Miquelon. The 28 stations presented here (Table 1) will provide a background for the study which will be complemented by 69 bathythermographs taken every 20 nautical miles. A plankton haul will also be made at each station.

Table 1. Positions of hydrographic stations.



Fig. 1. Distribution of surface temperature.


Fig. 2. Distribution of near bottom temperaturt.



Fig. 4A. Temperatures - Section I - Burin Peninsula/South of Saint Pierre Bank.


Fig. 4B. Salinities - Section I - Burin Peninsula/South of Saint


500

Fig. 5A. Temperatures - Section II - South of Banquereau/South of Saint Pierre Bank.


500

Fig. 5B. Salinities - Section II - South of Banquereau/South of Saint Pierre Bank.


Fig. 6A. Temperatures - Section III - Cape Breton/South of Banquereau.


Fig. 6B. Salinities - Section III - Cape Breton/South of Banquereau.


Fig. 7A. Temperatures - Section IV - North of Misaine Bank/ North of Saint Pierre Bank.


Fig. 7B. Salinities - Section IV - North of Misaine Bank/ North of Saint Pierre Bank.


Fig. 8A. Temperatures - Section V - Cabot Strait.


Fig. 8B. Salinities - Section V - Cabot Strait.


4. Hydrographic conditions in the Northwest Atlantic in $1970^{1}$

by V.V. Burmakin and B.P. Kudlo PINRO, Murmansk

In 1970 R/V Perseus III, Rossiya and Protsion conducted oceanographic research for PINRO in Subareas 1, 2 and 3. More than 1,100 hydrographic stations were occupied over standard sections during six cruises. The main portion of the hydrographic work was completed by R/V Protsion. Temperatures and salinities, as well as oxygen and phosphorus concentrations and other hydrochemical elements, were determined at most of the stations. Mean estimates of temperature and salinity on the sections for different water masses and branches of currents were calculated in the boundaries determined by Elizarov (1962) and Burmakin (1971).

## Subarea 1

In August 1970, hydrographic investigations were conducted by R/V Perseus III in the northern part of Davis Strait between $62^{\circ} 30^{\prime} \mathrm{N}$ and $65^{\circ} 30^{\prime} \mathrm{N}$. (Div. 1C1D). Treatment of the data obtained by means of the dynamic method showed that a secondary cyclonic vortex with low current velocities was located in the central part of the area investigated. This vortex developed on the northern periphery of the vortex of the Labrador Sea.

Comparison of the dynamic charts with charts of isotherms and isohalines obtained during the same survey shows good agreement between circulation, and temperature and salinity. The central part of the cyclonic vortex is characterized by the highest temperatures at all depths. The same result is obtained, if we compare the dynamic chart and distribution of salinity. At the surface in the area of the vortex it is above $33 \%$ (the maximum is $33.28 \%$ 。). The water temperature fluctuated here from $3^{\circ}$ to $4.4^{\circ}$. Especially great is the similarity between the surface currents and the temperature at 100 m , where the influence of summer warming is insignificant. The $1^{\circ}$ isotherm at this depth is nearly the same in pattern as the isolines of dynamic heights in the area of the vortex, with below zero temperatures over Tovqussaq Bank. It seems that the circulation field can be judged indirectly from observations of water temperature, and best from those at the 100 -m level.

Estimates of discharge through Davis Strait on Sections 11-A and 10-A were also obtained by means of the dynamic method (Table 1).

The results seem to contradict the present point of view on water transport through Davis Strait from the Baffin Sea into the Atlantic. It is quite possible that the results of calculations of water discharge were affected by the fact that no observations were made in coastal waters. Nevertheless, the predominance of discharge through the Strait in the northern direction in August 1970 allows us to make a supposition concerning the availability of great fluctuations of water exchange through Davis Strait.

It is possible to estimate the temperature conditions on Lille Hellefiske and Fyllas Banks in August from observations made on Sections 10-A and 11-A.

[^3]Table 1. Water discharge through Davis Strait in August 1970.

| Section | Date | Position | Discharge <br> to the <br> North | Discharge <br> to the <br> South | Total <br> Discharge <br> (to the <br> North) |
| :---: | :---: | :---: | :---: | :---: | :---: |
| $10-\mathrm{A}$ | $14-16$ August | $65^{\circ} 27^{\prime}, 64^{\circ} 59^{\prime}, 65^{\circ} 18^{\prime}$ <br> $54^{\circ} 69^{\prime}-57^{\circ} 36^{\prime}-60^{\circ} 47^{\prime}$ | 3.59 | 2.36 | 1.23 |
| $4-5$ August | $63^{\circ} 57^{\circ} 63^{\circ} 03^{\prime}-63^{\circ} 41^{\prime}$ <br> $53^{\circ} 08^{\prime}-58^{\circ} 56^{\prime}-62^{\circ} 30^{\prime}$ | 8.88 | 6.98 | 1.90 |  |

In Table 2, prepared by courtesy of I.I. Svetlov, these data are compared with data from observations made in previous years (though the data are not given for one and the same date).

Table 2 shows that the temperature of both components of the West Greenland Current was lower in Div. 1C-1D (Sections 10-A and 11-A) in August 1970, the $0-50-m$ layer being coldest in the years under consideration.

Table 2. Mean water temperature of the warm and cold components of the West Greenland Current by depth layers in some years and months.

| Date | Section | Depths (m) |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | 0-50 |  | 0-200 |  | 50-200 |  | $\begin{array}{\|l\|} \hline 200-500 \\ \hline \text { warm } \\ \text { comp. } \\ \hline \end{array}$ |
|  |  | warm comp. | $\begin{aligned} & \text { cold } \\ & \text { comp. } \end{aligned}$ | warm comp. | $\begin{aligned} & \text { cold } \\ & \text { comp. } \end{aligned}$ | warm comp. | $\begin{aligned} & \text { cold } \\ & \text { comp. } \end{aligned}$ |  |
| 3 August 1959 | 10-A | 2.40 | - | 1.52 | - | 1.11 | - | 3.40 |
| 19 August 1963 |  | 3.82 | - | 2.62 | - | 2.20 | - | 3.32 |
| 8-9 August 1964 |  | 3.31 | - | 2.01 | _ | 1.48 | - | 3.51 |
| 10-12 August 1967 |  | 3.08 | - | 1.38 | - | 0.56 |  | 4.14 |
| 14-16 August 1970 |  | 2.57 | - | 1.62 | - | 1.30 | - | 4.03 |
| 3 August 1964 | 11-A | 4.09 | 3.96 | 4.24 |  |  | 3.36 |  |
| 16-18 August 1969 |  | 2.20 | 1.31 | 2.00 | 1.71 | 1.90 | 1.70 | 4.36 |
| 4-5 August 1970 |  | 1.95 | 1.48 | 1.48 | 1.05 | 1.29 | 0.98 | 4.06 |
| 13 September 1961 | 8-A | 7.52 | 4.44 | 6.85 | 5.36 | 6.66 | 5.66 | 5.52 |
| 23-24 September 1963 |  | 4.26 | 2.02 | 5.63 | 3.24 | 6.14 | 3.76 | 5.59 |
| 1-2 September 1970 |  | 3.86 | 0.64 | 4.24 | 1.42 | 4.37 | 3.76 1.70 | 4.73 |
| 19-21 October 1962 | 8-A | 6.32 | 3.44 | 6.12 | 4.16 | 6.06 | 4.52 | 5.04 |
| 30-31 October 1963 |  | 4.96 | - | 5.45 | - | 5.62 | 4.52 | 5.26 |
| 27-29 October 1964 |  | 6.61 | 4.84 | 6.53 | 5.21 | 6.52 | 5.38 | 5.39 |
| 16-19 October 1966 |  | 6.59 | 2.60 | 6.27 | 3.25 | 6.17 | 3.50 | 5.60 |
| 25-26 October 1970 |  | 2.82 | 0.13 | 4.07 | 1.18 | 4.49 | 1.55 | 5.13 |

According to data obtained from Section 8-A. in September and October, temperatures $1-4^{\circ}$ lower than in some previous years were also registered in Div. $1 F$.

As far as one can judge from such fragmentary data, water temperature in August-October 1970 was lowest on the West Greenland Shelf, as compared with the temperature for the same months in the years under comparison. Cooling was especially marked in the surface layer and tended to increase from August to the end of the year.

Subareas 2 and 3
Temperature
The comparative characteristics of temperature conditions for Subareas 2 and 3 in 1970 was shown by the mean temperature in the $0-200 \mathrm{~m}$ layer on standard hydrologic Sections 8-A, 7-A, 6-A, 4-A, 3-A, 2-A, 1-A, and 44-A. In Table 3, the temperature is given for each month when the observations were made. The deviation of each mean temperature from its annual mean estimate on the date of observation is also given. Curves of seasonal variations of water temperature of the $0-200 \mathrm{~m}$ layer for each section ( jet , water mass) were constructed from all available data obtained by the research vessels of the Polar Institute and the International Ice Patrol (see the paper by V.V. Burmakin, ICNAF, 1971). These curves allow us to estimate the anomaly of the mean temperature on each section on the date of observation.

Table 3 shows that in January 1970 temperature anomalies in the $0-200 \mathrm{~m}$ layer were positive on all sections made on the Grand Bank and Cabot Strait and fluctuated from 0.06 to $1.94^{\circ}$. The highest anomalies were recorded on Section l-A on the southwestern slope of the Grand Bank. A negative anomaly $\left(-0.44^{\circ}\right)$ on Section 2-A recorded on the southern slope of the Grand Bank was the only exception to these positive anomalies.

Beginning in February, the anomalies were negative or near the norm in the whole of the Subarea 3. This was observed to the end of August.

In Subarea 2 observations were made in May and showed a positive anomaly of $+0.20^{\circ}$ (Section $8-\mathrm{A}$, intercept B). In September positive anomalies were still found, but by the end of October negative anomalies ( $-0.04^{\circ}$ ) were observed in the main branch of the Labrador Current (B).

In autumn in Subarea 3 the anomalies were negative on the northeastern (Section 7-A) and southeastern (Sections 4-A and 3-A) slopes of the Grand Bank, whereas in the south (Section 2-A) and in Flemish Cap Channel they were slightly positive (+0.06).

In September positive anomalies were observed in Cabot Strait and on the southwestern slope of the Grand Bank (Sections 44-A and 1-A), but they were about 3 times smaller than those in January.

Thus, at the beginning of 1970, waters in Subareas 2 and 3 were warm, in spring and summer temperatures were below the norm and at the end of the year an insignificant warming was observed in the western part of Subarea 3, whereas the waters in the east of the Grand Bank became considerably colder. Evidently, this can be explained by the simultaneous increase of intensity of the Labrador Current, which caused the cooling on the eastern slope of the Grand Bank, and
of the Gulf Stream, that warmed the western part of the Bank.

Table 3. Mean temperature of the $0-200 \mathrm{~m}$ layer and its anomalies on standard sections in the areas of Labrador and Newfoundland in 1970.

| Month | Section |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 8-A (B) | 7-A | 6-A (G) | 4-A | 3-A | 2-A | 1-A | 44-A |
| JANUARY Date t $\Delta \mathrm{t}$ | - | - | $\begin{array}{r} 3 \\ 4.01 \\ +0.25 \end{array}$ | $\begin{array}{r} 7 \\ 3.95 \\ +0.83 \end{array}$ | $\begin{array}{r} 9 \\ 0.79 \\ +0.06 \end{array}$ | $\begin{array}{r} 25 \\ 1.43 \\ -0.44 \end{array}$ | $\begin{array}{r} 12 \\ 6.39 \\ +1.94 \end{array}$ | $\begin{array}{r} 17-18 \\ 3.50 \\ +0.72 \end{array}$ |
| FEBRUARY <br> Date $t$ $\Delta t$ | - | - | - | - | 11 0.32 -0.52 | - | - | - |
| $\begin{array}{lr} \text { MAY } & \\ & \text { Date } \\ & t \\ & \Delta t \end{array}$ | $\begin{array}{r} 4-5 \\ 0.18 \\ +0.20 \end{array}$ | $\begin{array}{r} 17 \\ 0.90 \\ -0.20 \end{array}$ | $\begin{array}{r} 19 \\ 1.39 \\ -0.21 \end{array}$ | $\begin{array}{r} 15 \\ 2.05 \\ +0.04 \end{array}$ | $\begin{array}{r} 16 \\ 0.48 \\ -0.28 \end{array}$ | - | - | - |
| JUNE <br> Date <br> $t$ <br> $\Delta t$ | - | - | - | - | - | - | - | 19 4.08 -0.33 |
| $\begin{array}{ll} \text { JULY } \\ & \text { Date } \\ & \mathrm{t} \\ & \mathrm{t} \end{array}$ | - | - | - | - | - | 14 2.95 -0.09 | 20 6.00 -0.20 | - |
| ```AUGUST Date t \Delta t``` | - | 4 0.99 -0.80 | 9 1.96 -1.29 | - | - | - | - | - |
| SEPTEMBER Date $t$ $\Delta t$ | $\begin{array}{r} 4-5 \\ 1.32 \\ +0.50 \end{array}$ | $\begin{array}{r} 22-23 \\ 1.70 \\ -0.67 \end{array}$ | - | - | - | - | $\begin{array}{r} 24 \\ 7.45 \\ +0.69 \end{array}$ | $\begin{array}{r} 27 \\ 4.95 \\ +0.53 \end{array}$ |
| $\begin{array}{r} \text { OCTOBER } \\ \text { Date } \\ \mathrm{t} \\ \Delta \mathrm{t} \end{array}$ | $\begin{array}{r} 30 \\ 1.36 \\ -0.04 \end{array}$ | - | $\begin{array}{r} 6 \\ 4.39 \\ +0.06 \end{array}$ | $\begin{array}{r} 10-11 \\ 2.94 \\ -0.77 \end{array}$ | $\begin{array}{r} 12 \\ 1.49 \\ -0.48 \end{array}$ | 20 3.06 +0.06 | - | - |

The temperature of the $0-200 \mathrm{~m}$ layer, on Sections 4-A and 3-A, adjusted for 15 May, and on Section 8-A adjusted for 1 November, was compared with the norm (Table 4). On Sections 4-A and 3-A the norms were calculated for the
period 1936-41, 1949-54, 1956-59, 1961-63, 1968-69, and on Section 8-A(B), for 1958, 1962, 1964-68, In our previous reports to ICNAF, the norms for Sections 4-A and 3-A were those accepted by Elizarov (1962).

Table 4. Mean temperature of the $0-200 \mathrm{~m}$ layer on Sections 3-A, 4-A on 15 May and 8-A on 1 November for 1968-70, as compared with the norm.

| Year | 3-A, 15 May |  |  | 4-A, 15 May |  |  | 8-A, 1 November |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\mathrm{t}^{\circ} \mathrm{C}$ | norm | $\Delta t$ | $t{ }^{\circ} \mathrm{C}$ | norm | $\Delta \mathrm{t}$ | $t{ }^{\circ} \mathrm{C}$ | norm | $\Delta t$ |
| 1968 | 1.85 | 0.49 | +1.36 | 2.25 | 1.71 | +0.54 | 0.50 | 1.00 | -0.50 |
| 1969 | 0.80 | - | +0.31 | 3.46 | - | +1.75 | 0.50 | - | -0.50 |
| 1970 | 0.44 | - | -0.05 | 2.05 | - | +0.34 | 0.60 | - | -0.40 |

On Sections 3-A and 8-A the mean temperature was calculated for the stations situated within the cold component of the Labrador Current, and on Section 4-A, for the stations situated in the frontal zone of the waters of the Banks, of Labrador and North Atlantic waters.

Table 4 shows that, in spring and autumn 1970 on Sections 3-A and 8-A in cold Labrador waters, the temperature was below the norm, whereas on Section 4-A in the Frontal Zone it was above the norm. Signs of the anomalies, determined by two methods, coincide (Tables 3 and 4).

Concerning the three-year period, 1968-70, in 1970 the temperature on the southeastern slope of the Grand Bank was lower by $0.20-1.41^{\circ}$ and insignificantly higher in the area of South Labrador, than in 1968 and 1969 (Table 4).

A comparison of charts of water temperature distribution in the near-bottom layer on the Grand, Green and St. Plerre Banks for the last three years shows that the temperature at the bottom in 1970 was higher than in 1969 and considerably lower than in 1968.

## Salinity

Observations carried out in the intercept B of Section 89A give information on salinity of the main branch of the Labrador Current on the Hamilton Bank (Elizarov, 1962). Average values of salinity in the $0-200 \mathrm{~m}$ layer in the intercept B , calculated from data collected by the International Ice Patrol and USSR research vessels for the period 1936-57 ( 15 sections) are taken from Elizarov (1963), and for the period 1958-70 ( 45 sections) are calculated by us. The results of these calculations are inserted into the graph (Fig. 1) in accordance with the date of each observation on the section. The figure shows that the many years' data used by us do not give complete information on the yearly variation of salinity, due to the fact that no observations were made in some months. However, the curve of the yearly variation of salinity can be drawn with a certain degree of approximation. If we accept the given curve for the "norm", it is possible to obtain the anomaly of salinity for every case observed on the section for the actual date of observation. Such a method
of calculation of the value of salinity anomalies (as well as anomalies of other hydrological parameters) makes it possible to define the hydrological conditions of the year, even if the results from few observations made at different times are available in the given year. In Fig. 1 the upper and lower curves show the yearly variations of the extreme values of the mean salinity of the Labrador Current.

The method described was used to eatimate the salinity of the Labrador Current in 1969 and 1970. Table 5 shows that in June 1969 the salinity of the Labrador Current was somewhat increased, and by the end of the year the negative anomaly reached $1 / 4^{\circ} / \circ 0$. As noted by Templeman (1970) on 4 August 1969, the salinity on the Seal Islands-Hamilton Bank section at a depth of 30 m and more was almost always below the mean, being measured at different stations, but close to values obtained in the last four years, pertaining to the period 195165.

Table 5. Mean salinity value and its anomaly of polar waters of the Labrador Current in the $0-200 \mathrm{~m}$ layer in accordance with the observations in 1969 and 1970 on Section 8-A.

| Date | $\%$ | $\%$ |
| :--- | :---: | :---: |
| 27-31 July 1969 | 33.49 | +0.10 |
| 17-19 December 1969 | 33.50 | -0.25 |
| 4-5 May 1970 | 33.85 | +0.07 |
| 4-5 September 1970 | 33.58 | +0.15 |
| 30 October 1970 | 33.59 | +0.04 |

Observations in 1970 show that in the spring to autumn period the salinity of the Labrador Current was somewhat above the norm or near it.

## Conclusions

1. The secondary cyclonic vortex, which developed on the northern periphery of the circulation of the Labrador Sea, separated the West Greenland Current from the Baffin Island Current in August 1970. High temperatures and salinities are characteristic of the zone of the vortex.
2. From August to October water temperature on the West Greenland Shelf was the lowest for the last ten years. The cooling was most marked in the surface layer and evidently increased from August to the end of the year.
3. In the beginning of 1970 positive anomalies predominated in Subareas 2 and 3, in spring and summer of this year negative anomalies prevailed; by the end of the year some warming was observed in the western part of the Grand Bank and some cooling in the eastern part, and slight cooling at South Labrador.
4. As compared with the norm, the mean temperature of the $0-200 \mathrm{~m}$ layer on the eastern Grand Bank adjusted for 15 May , and at South Labrador adjusted for

1 November, had anomalies of the same signs when compared with the curve of the yearly temperature variation. However, the values of the anomalies differed considerably.
5. In May-August 1970 in near-bottom layers of Subarea 3, the temperature was higher than in the same months of 1969 and insignificantly lower than in 1968.
6. From May to October 1970, salinity in the core of the Labrador Current was somewhat higher than the norm.

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SECTION B COD
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5. Selection of cod by polyamide trawl codends in ICNAF Div. $4 \mathrm{Vn}^{1}$
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## Introduction

Recently the ICES/ICNAF Joint Working Group on Selectivity Analysis made an attempt to assess the effect of the physical properties of the netting yarns on the selectivity (ICNAF, $1970 a$ and $b$ ). This attempt failed because until now very little data on the properties of the netting yarns used in selection experiments have been published. However, the Group thought that the elongation (extensibility) of the netting yarns is possibly one of the most important properties by which the selectivity might be influenced. Since this assumption was not yet substantiated scientifically, the Group proposed an international experiment to find out whether high elongation and high selectivity are positively correlated or not (ICNAF, 1970b, p. 4). Realizing that, in general, considerable time is needed for the preparation of such an International test, the Institut fur Fangtechnik decided to start its own study of the relationship between elongation and selectivity already in the spring of 1970.

The cruise program also included the collection of selection data for a codend made of a definite polyamide netting yarn $R 6,484$ tex. According to a recommendation of the above-mentioned Working Group (ICNAF, 1970a, p. 49), this netting yarn has been introduced to replace manila as a new standard for selectivity purposes ${ }^{2}$. It was suggested "that, whenever possible, research vessels should use codends made of the new standard polyamide and collect selectivity data so that a large number of selection factors covering all species, seasons, areas, etc., be obtained as quickly as possible, ..." (ICNAF, i970b, p. 4).

Another important item of the cruise program was the repetition of an experiment carried out with an extra strong codend in 1969 (Bohl, 1970). This experiment had shown that there is no significant difference between the selectivities of netting yarns of normal diameter and those which are unusually thick. Therefore, it was concluded that extra strong codends can be used to obviate the need for topside chafers. Since this is a matter of great consequence, it was thought advisable to confirm the result of the previous experiment by a second test.

## Material and methods

The experiments were carried out during April 1970 on the Scotian Shelf between Cape•Breton and Cape Smoky in depths ranging from 105 to 180 m (IGNAF Div. 4Vn). FRB Walther Herwig - a diesel-electric stern trawler of 83.23 m

[^4]total length, 1,987 gross tons, capable of developing $2,000 \mathrm{~h} . \mathrm{p} . \mathrm{e}$. at 190 rpm - used the German standard roundfish bottom trawl (140-ft groundrope).

The selectivities of four polyamide multifilament codends were studied. Two of these codends have already been mentioned above, viz. those which are made of the standard polyamide ( $\mathrm{R} 6,484 \mathrm{tex}$ ) and the extra strong netting yarn ( $\mathrm{R} 17,465$ tex $^{3}$ ). The remaining two codends were procured especially to evaluate the relationship between elongation and selectivity. The netting yarns of these codends are made of one and the same kind and number of single yarns (article no. $N_{t} 3 / 500$ ) and differ inasmuch as the yarn of the one codend is extremely hard twisted (high twist factor) and untreated to produce a high elongation, while the yarn of the other is medium twisted (low twist factor) and thermofixed to produce a low elongation. In consequence of the different netting yarn construction, there are also inevitable differences in some other physical properties, e.g., fineness, diameter and flexibility.

The properties of the netting yarns and codend nettings used in the experiments were determined according to the ISO standards for testing methods. The results are given in Table 1. The relation between load and elongation, which is based on new netting yarns, is shown in Fig. 1.

During the experiments the covered codend technique was used. The topside covers used were in accordance with ICES specifications. They were made of single polyamide netting yarn ( 25 tex $\times 16 \times 3$, twisted), and the mesh opening was about 60 mm . The inner underside of the codends was lined with smallmeshed netting similar to that of the cover.

The length composition of the catches was determined by measuring the total fish length to the centimeter below. The mesh measurements were taken immediately after each haul using an ICES gauge with an operating pressure of 4 kg 。

In order to study the girth/length relationship of cod, the unconstricted maximum body girth was measured to the nearest millimeter.

Results
During the course of the experiments a total of 51 successful hauls was made. The catches, ranging from 0.6 to 6.1 metric tons, were of rather uniform composition. Cod were always predominant; they represented in 32 hauls more than $90 \%$, in 12 hauls $80-90 \%$ and only in 7 hauls less than $80 \%$ of the catch weights. The duration of tow varied from 30 to 120 minutes, but in most cases (42 hauls), the trawl was towed for one hour.

The relative length composition of the total cod catch which consisted of 66,258 specimens caught in the codends and 31,479 specimens caught in the covers, is shown in Fig. 2. Fish of the $45.0-47.9 \mathrm{~cm}$ length-class were most abundant. This fact proved extremely favourable for the experiments, because the $50 \%$ retention length of the four codends used were found to be close to the modal length of the fish aggregation. Thus, unusually large numbers of specimens could be recorded within the selection ranges.

The selection data obtained from combined hauls are compiled in Table 2.

[^5]- 59 -
Table 1. Information about codends and netting yarns used.

| Codend no. | 54 | $\stackrel{55}{ } \stackrel{56}{\text { Polyamide multifilament }}$ |  | 57 |
| :---: | :---: | :---: | :---: | :---: |
| Material and type of fibre |  |  |  |  |
| Construction of netting yarn | Twisted | Plaited | Twisted | Twisted |
| of codend |  | Double braided |  |  |
| Method of manufacture of codend |  | Hand-made |  |  |
| Treatment of netting yarn | Thermo-fixed | Untreated Untreated |  | Thermo-fixed |
| of netting |  |  |  |  |
| Age of codend in fishing hours | 55 | 35 | 0 | 0 |
| Rtex (g/ $1,000 \mathrm{~m}$ ) | 6,484 | 17,465 | 6,588 | 5,655 |
| Runnage (m/kg) | 154 | 57 | 152 | 177 |
| Diameter (mm) | 2.9 | $\simeq 7.0$ | 3.1 | 2.7 |
| Flexibility ${ }^{1}$, wet (g) | 48 | 246 | 375 | 30 |
| Amount of twist in turns per meter (t/m) | 74 | - | 100 | 62 |
| Twist factor ${ }^{2}=\propto$ tex $=t / m \times \sqrt{\text { Rtex } / 1,000}$ | 188 | - | 256 | 147 |
| Weaver-knot breaking load, wet (kp) | 299 | 735 | 257 | 292 |
| Breaking load, without knot, dry (kp) | 355 | 854 | 254 | 306 |
| without knot, wet (kp) | 315 | 752 | 194 | 267 |
| Breaking length, dry (km) | 54.9 | 48.9 | 38.6 | 54.1 |
| wet (km) | 48.6 | 43.1 | 29.4 | 47.2 |
| Knot breaking length, wet, $1 / 2$ (km) | 23.1 | 21.0 | 19.5 | 25.8 |
| Elongation (\%) at a load of |  |  |  |  |
| 5 kp | 2.2 | 0.3 | 5.6 | 1.8 |
| 10 kp | 4.3 | 1.1 | 9.1 | 3.3 |
| 30 kp | 10.3 | 3.8 | 20.1 | 9.6 |
| 50 kp | 13.9 | 6.2 | 25.7 | 13.1 |
| 1/2 weaver-knot breaking load, wet (kp) | ) 23.8 | 21.0 | 38.8 | 21.5 |

The flexibility (resistance against deformation) has been determined by means of the "Lbtzener Method"
2 The twist factor makes it possible to compare the amount of twist of netting yarns of different fineness.

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Table 2. Compilation of cod selection data for grouped hauls.


The selection curves for combined hauls are shown in Fig. 3. They are based on smoothed percentages of retained fish (three-point moving averages). The curves are fitted by eye.

From Table 2 it can be seen that the extra strong codend gave a selection factor of 3.51 , while the codend made of the standard polyamide yielded a factor of 3.41. Hence, the results obtained from the same codends in 1969 (Boh1, 1970) could be confirmed: there is no significant difference between the selectivity properties of extra strong and "normal" polyamide codends. This implies that the widespread use of such a robust codend as tested in the German experiments could be considered as a possible step towards the elimination of topside chafers.

As to the codends with different elongations (codend nos. 56 and 57), Table 2 appears to show that a low elongation is associated with a low selection factor, and vice versa: the selection factor of the netting yarn with $21.5 \%$ elongation was 3.15 and that of the netting yarn with $38.8 \%$ elongation was 3.49. However, taking into account that the two other codends used in the experiments were also characterized by low elongations (comp. codend nos. 54 and 55 in Table 1 and Fig. 1), and realizing that these codends gave selection factors of the same order of magnitude as the codend with the high elongation, it was rather doubtful whether the elongation was really the causative agent for the selectivity differences observed. Looking at all the data given in Table 2, it becomes obvious that the lowest selection factor obtained (3.15, codend no. 57, "low elongation") was based on the largest catches made during the tests. This suggests that this low selection factor could probably be attributed to catch size.

To look into this matter in more detail, an analysis of single hauls had to be made. Since cod were sufficiently numerous in most of the catches, reliable selection factors could be obtained for 44 individual hauls. Only with 7 hauls was it not possible to determine the selection factors quite precisely. These factors are queried in Tables 3-6, in which all the data are presented on a haul-by-haul basis.

In the following the unweighted mean selection factors calculated from single hauls are compared with the selection factors calculated from combined hauls.

| Codend no. | 54 | 55 | 56 | 57 |
| :--- | :---: | :---: | :---: | :---: |
| Selection factor based <br> on grouped hauls | 3.41 | 3.51 | 3.49 | 3.15 |
| Unweighted mean <br> selection factor $\pm$ s.e. <br> based on single hauls | $3.44 \pm 0.03$ | $3.50 \pm 0.05$ | $3.50 \pm 0.05$ | $3.16 \pm 0.02$ |
| Range of selection <br> factors | $3.28-3.65$ | $3.16-3.78$ | $3.27-3.80$ | $3.10-3.25$ |
| Number of hauls | 18 | 15 | 12 | 6 |

Table 3. Cod selection data for individual hauls; codend no. 54.

Eaul Date Poaition Depth Duration Mesh 50\% Selection No. of codin Total no. of Yuantity (bagketa) of 1970 (min) (mm) (mm) Codend Cover Codend Cover Codend Cover Codend Cover $\begin{array}{lllllllll}165 & 1,064 & 256 & 22 & 1 / 4 & 3 & 1 / 2 & 1\end{array}$ $\begin{array}{llllllll}602 & 1,989 & 782 & 38 & 3 / 4 & 91 / 2 & 21 / 2 & +\end{array}$ $\left.\begin{array}{llllllllllllllll}20 & 13 & 486^{\circ} 071 \mathrm{~N} & 155 & 60 & 132.2 & 445 & 3.37 & 113 & 442 & 427 & 1,087 & 498 & 22 & 51 / 2 & 2\end{array}\right] 1 / 4$. \begin{tabular}{llllllllllllllllll}
\hline 22 \& 13 \& 460081 N \& 175 \& 60 \& 131.3 \& 479 \& $3.65 ?$ \& $97 ?$ \& $77 ?$ \& $73 ?$ \& 389 \& 111 \& 12 \& $3 / 4$ \& 2 \& $11 / 2$ \& $1 / 2$

 

7 \& 1,231 \& 286 \& 28 \& $31 / 2$ \& $11 / 2$ \& $1 / 2$ <br>
\hline

 $\begin{array}{lllllllll}90 & 1,046 & 161 & 27 & 1 / 2 & 21 / 2 & 1 & 1 / 4 & 1 / 4\end{array}$ $\begin{array}{lllllll}3 & 2,579 & 1,262 & 53 & 15 & 31 / 4 & 3 / 4\end{array}$ 

3909 \& 1,661 \& 665 \& 39 \& $1 / 2$ \& 8 \& 3 <br>
\hline

 $\begin{array}{lllllllll}180 & 1,318 & 278 & 40 & 3 / 4 & 3 & 1 / 4 & 21 / 2 & 1 / 4\end{array}$ 

204 ? \& 936 \& 378 \& 21 \& $1 / 4$ \& $31 / 2$ \& $33 / 4$ \& $1 / 4$ <br>
\hline

 

287 \& 738 \& 403 \& 13 \& $1 / 4$ \& 4 \& $1 / 4$ \& 3 \& $3 / 4$ <br>
\hline

 

202 \& 700 \& 299 \& 21 \& 3 \& $1 / 4$ \& 3 <br>
\hline

 $\begin{array}{lllllll}217 & 553 & 371 & 133 / 4 & 4 & 31 / 2 & 1 / 4\end{array}$ $\begin{array}{llllllll}11^{\Gamma} & 318 & 311 & 8 & 31 / 2 & 21 / 4 & 1 / 2\end{array}$ 

\hline 78 \& 20 \& 59 \& <br>
\hline
\end{tabular}


Table 4. Cod selection data for individual hauls; codend no. 55.



 | 47 | 17 | $46^{\circ}{ }^{\circ} 5^{\prime} \mathrm{N}$ | $130-$ | 60 | 122.4 | 387 | 3.169 | 116 | 226 | 172 | 794 | 238 | 14 | 2 | 5 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |



 $\begin{array}{llllllllllllllllllllllllll}59 & 19 & 46^{\circ}{ }^{\circ} \mathbf{4 N}^{\prime} \mathrm{N} & 140- & 60 & 122.7 & 458 & 3.73 & 85 & 791 & 852 & 1,975 & 1,152 & 42 & 1 / 4 & 13 & 1 / 2 & 3 & 1 / 2 & 1 / 4\end{array}$ | 901 | 60 | 11 | $1 / 2$ | $51 / 2$ | $1 / 4$ |
| :--- | :--- | :--- | :--- | :--- | :--- |


Table 5. Cod selection data for individual hauls; codend no. 56.

| $\begin{aligned} & \text { Haul } \\ & \text { No. } \end{aligned}$ | $\begin{aligned} & \text { Date } \\ & \text { Apri1 } \\ & 1970 \end{aligned}$ | Position | $\begin{gathered} \text { Depth } \\ \text { (m) } \end{gathered}$ | Duration of haul (min) | $\begin{aligned} & \text { Meah } \\ & \text { size } \\ & \text { (mmi) } \end{aligned}$ | 50\% Selection |  |  | No, of cod in Selection range |  | Total no. of cod caught |  | Quantity (baskets) of |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  | $\underset{(\mathrm{mm})}{\mathrm{Length}} \underset{(\mathrm{mm})}{\text { Factor }}$ |  |  |  |  |  |  | By-catch |  |
|  |  |  |  |  |  |  |  |  | Codend | Cover |  |  | Codend | Cover | Codend | Cover | Codend | Cover |


| 25 | 13 | $\begin{aligned} & 46^{\circ} 08^{\prime} \mathrm{N} \\ & 58^{\circ} \frac{491 \mathrm{~W}}{} \\ & \hline \end{aligned}$ | 160 | 60 | 132.0 | 447 | 3.39 | 84 | 221 | 186 | 1,293 | 295 | 31 3/4 | 4 | $13 / 4$ | 1/2 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 26 | 14 | $\begin{array}{r} 46^{\circ} 08^{\prime} \mathrm{N} \\ 58^{\circ} 501 \mathrm{~W} \\ \hline \end{array}$ | $\begin{array}{r} 160- \\ 140 \\ \hline \end{array}$ | 60 | 128.7 | 444 | 3.45 | 79 | 312 | 306 | 1,382 | 470 | 32 1/2 | 5 | $11 / 2$ | 1/2 |
| 27 | 14 | $\begin{array}{r} 46^{\circ} 091 \mathrm{~N} \\ 58^{\circ} 50^{1 \%} \\ \hline \end{array}$ | 140 | 60 | 129.0 | 440 | 3.41 | ? | ? | ? | 1,550 | 424 | $361 / 2$ | 5 | $21 / 2$ | 1/2 |
| 28 | 14 | $\begin{array}{r} 46^{\circ} 09^{\prime} \mathrm{N} \\ 58^{\circ} 51^{\prime} \mathrm{W} \\ \hline \end{array}$ | 160 | 60 | 128.1 | 419 | 3.27 | 96 | 186 | 144 | 958 | 230 | 20 | $21 / 2$ | $11 / 4$ | 1/4 |
| 29 | 14 | $\begin{aligned} & 46^{\circ} 091 \mathrm{~N} \\ & 58^{\circ} 51^{\prime} \mathrm{W} \\ & \hline \end{aligned}$ | 160 | 60 | 127.4 | 4819 | 3.78? | $65 ?$ | $66 ?$ | 92? | 211 | 238 | $41 / 2$ | $21 / 2$ | 1 | 1/2 |
| 32 | 14 | $\begin{aligned} & 46^{\circ} 077^{\prime} \mathrm{N} \\ & 58^{\circ} 51^{\prime} \mathrm{W} \\ & \hline \end{aligned}$ | 155 | 90 | 127.0 | 430 | 3.39 | ? | $?$ | $?$ | 1,819 | 550 | 40 1/2 | 6 | $31 / 4$ | 1/2 |
| 33 | 14 | $\begin{aligned} & 46^{\circ} 131 \mathrm{~N} \\ & 59^{\circ} 00^{\prime} \mathrm{W} \\ & \hline \end{aligned}$ | 150 | 60 | 127.1 | 450 | 3.54 | 97 | 553 | 500 | 1,558 | 695 | $313 / 4$ | 8 | $31 / 4$ | 1/4 |
| 34 | 15 | $\begin{aligned} & 46^{\circ}{ }^{13 \prime N} \\ & 59^{\circ} 00^{\prime} \\ & \hline \end{aligned}$ | 150 | 60 | 127.5 | 485 | 3.80 | 106 | 357 | 547 | 873 | 765 | 19 | $81 / 2$ | $11 / 4$ | 1/4 |
| 55 | 19 | $\begin{aligned} & 46^{\circ}{ }^{\circ} 45^{\prime} \mathrm{N} \\ & 59^{\circ} 46^{\prime} \mathrm{VI} \\ & \hline \end{aligned}$ | $\begin{aligned} & 110- \\ & 120 \\ & \hline \end{aligned}$ | 60 | 123.4 | 418 | 3.39 | 68 | 411 | 388 | 1,248 | 627 | 19 1/2 | $51 / 2$ | $21 / 2$ | + |
| 56 | 19 | $\begin{aligned} & 46^{\circ}{ }^{44} 1 \mathrm{~N} \\ & 599^{\circ} 31 \mathrm{~W} \\ & \hline \end{aligned}$ | $\begin{aligned} & 115- \\ & 105 \\ & \hline \end{aligned}$ | 60 | 124.2 | 425 | 3.42 | 84 | 443 | 432 | 1,042 | 623 | $181 / 2$ | $51 / 2$ | 3 | 1/2 |
| 57 | 19 | $\begin{aligned} & 46^{\circ} 42^{\prime} \mathrm{N} \\ & 59^{\circ} 40^{\prime} \mathrm{II} \\ & \hline \end{aligned}$ | $\begin{aligned} & 115- \\ & 110 \\ & \hline \end{aligned}$ | 60 | 124.8 | 439 | 3.52 | 84 | 871 | 867 | 1,786 | 1,376 | $311 / 4$ | 14 | $41 / 4$ | 1/2 |
| 58 | 19 | $\begin{aligned} & 46^{\circ} 421 \mathrm{~N} \\ & 59^{9} 41^{1} \mathrm{~W} \\ & \hline \end{aligned}$ | $\begin{aligned} & 125- \\ & 110 \\ & \hline \end{aligned}$ | 60 | 125.2 | 450 | 3.59 | 77 | 790 | 902 | 1,692 | 1,239 | $291 / 4$ | 12 | 3 | 1/2 |

Table 6. Cod selection data for individual hauls; codend no. 57.

| Eaul | Date | Position | ${ }_{\text {(m) }}^{\text {Depth }}$ | Duration | Mesh |  | $\frac{\text { Selec }}{\text { Factor }}$ | $\frac{\text { tion }}{\text { Mange }}$ |  | cod in | Tot | no. of | Quan | tity (b | $\frac{\text { keta) }}{\text { By-c }}$ |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| No. | $\begin{aligned} & \text { April } \\ & { }_{1970} \end{aligned}$ |  |  | $\begin{gathered} \text { of haul } \\ \text { (min) } \end{gathered}$ | $\begin{gathered} \text { size } \\ (\mathrm{mm}) \end{gathered}$ | $\underset{(\mathrm{mm})}{\text { Length }}$ | Factor | $\begin{array}{r} \text { Mange } \\ (\operatorname{man}) \end{array}$ | $\frac{\text { Select }}{\text { Codend }}$ | $\frac{\text { n range }}{\text { Cover }}$ | Codend | $\frac{\text { aught }}{\text { Cover }}$ | $\frac{C_{0}}{\text { Codend }}$ | $\frac{\mathrm{d}_{\text {Cover }}}{}$ | ${ }_{\text {Codend }}$ | $\frac{\text { cover }}{\text { Corer }}$ |
| 37 | 15 | $\begin{aligned} & 46^{\circ} 111 \mathrm{~N} \\ & 58^{\circ} 57 \mathrm{~W} \\ & \hline \end{aligned}$ | 150 | 90 | 144.9 | 471 | 3.25 | 88 | 198 | 164 | 780 | 272 | 22 | 4 | 2 | 1/4 |
| 38 | 15 | $\begin{aligned} & 46^{\circ}{ }^{\circ} 11 \mathrm{~N} \\ & 58^{\circ} 54^{\prime} 17 \\ & \hline \end{aligned}$ | $\begin{aligned} & 160- \\ & 150 \\ & \hline \end{aligned}$ | 120 | 144.3 | 453 | 3.14 | 113 | 540 | 501 | 1,420 | 701 | $311 / 4$ | 9 | 3 | 1/2 |
| 39 | 16 | $\begin{aligned} & 46^{\circ} 17{ }^{\prime \prime N} \\ & 59^{\circ} 17 \text { 'W } \\ & \hline \end{aligned}$ | $\begin{aligned} & 105- \\ & 120 \\ & \hline \end{aligned}$ | 60 | 142.8 | 454 | 3.18 | 120 | 1,730 | 2,102 | 3,290 | 3.173 | 56 3/4 | $291 / 4$ | $21 / 4$ | 1 |
| 40 | 16 | $\begin{aligned} & 46^{\circ}{ }^{\circ} 17{ }^{\prime \prime} \mathrm{N} \\ & 59^{\circ} 15 \mathrm{w} \\ & \hline \end{aligned}$ | 120 | 45 | 142.7 | 448 | 3.14 | 107 | 969 | 1,158 | 1,895 | 1,614 | $321 / 4$ | 15 | 3 | 1 |
| 41 | 16 | $\begin{array}{r} 46^{\circ}{ }^{\circ} 71 \mathrm{~N} \\ 59^{\circ} 191 \mathrm{~W} \\ \hline \end{array}$ | 120 | 50 | 142.6 | 451 | 3.16 | 98 | 1,130 | 1,275 | 2,327 | 1,959 | 42 | 19 | $31 / 4$ | 1 |
| 42 | 16 | $\begin{aligned} & 46^{\circ}{ }^{181 \mathrm{~N}} \\ & 59^{\circ} 2211 \end{aligned}$ | 120 | 55 | 142.6 | 442 | 3.10 | 91 | 1,231 | 1,284 | 2,614 | 2,173 | $441 / 2$ | $203 / 4$ | 4 | 1 |

For each codend separately, it can be seen that the selection factors calculated in two ways do not differ to any appreciable extent. This indicates that the selectivity was not negatively correlated with the catch size. Otherwise, provided that a sufficiently wide range of catch sizes was covered by each codend, the selection factors for grouped hauls would have been expected to be markedly lower than the unweighted mean selection factors.

The absence of a reciprocal relation between selectivity and catch size is also shown in Fig. 4, in which the selection factors derived from individual hauls have been plotted against the corresponding quantities of cod caught in the codends.

Thus, it remains an open question why the selection factor for the codend no. 57 was found to be significantly lower than the selection factors for the three other codends. The difference can neither be attributed to the elongation properties of the netting yarns nor to the catch sizes. Further experiments are needed to find out which property or properties of a netting yarn are of primary importance in determining its selectivity.

The result of 959 girth measurements taken during the course of the selection experiments is shown graphically in Fig. 5. The relationship between maximum body girth (G) and total length (L) is described by the regression equation $G=0.538 \mathrm{~L}-3.80 \mathrm{~cm}$.

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Fig. 3. Cod selection curves for combined hauls.


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6. Water temperature as an index of cod distribution
in Subarea 1 (West Greenland) ${ }^{1}$
by I.I. Svetlov
PINRO, Murmansk
The present paper is a continuation of the author's work on determination of the relationships between water temperature and cod distribution on the West Greenland banks (Svetlov, 1970). Asynchronous relationships by which it is possible to forecast in advance the areas of the most probable cod concentrations are considered in the paper. A method for such forecasting was suggested by K.G. Konstantinov and used for forecasting the fishery importance of separate areas in the Barents Sea (Konstantinov, 1964; Konstantinov and Mukhin, 1964; Konstantinov, 1967a).

Some relationships between water temperature and fishery importance of ICNAF Div. 1B are considered in the paper; Store Hellefiske Bank is in this division where an intensive cod feeding is observed usually in summer and early autumn (Rasmussen, 1957; Sidorenko, 1964).

Water temperature anomaly of the $0-50,50-100,100-200 \mathrm{~m}$ layers on the station at $66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W}$ in July (Hermann, 1967) and in the area off Godthaab ( $64^{\circ} 11^{\prime} \mathrm{N}, 51^{\circ} 43^{\prime} \mathrm{W}$ ) was taken as an index of the heat state of the sea (Bulletin Hydrographique, 1959-1962). The fishery importance of Div. 1B as a percentage, calculated as a ratio of cod yield in Div. $1 B$ in the given month to the total yield in all divisions of Subarea 1 in the same month was taken as an index of cod distribution. While calculating the fishery importance of Div. 1B the data on cod catches, taken by all fishing gears, were used. Data on catches were obtained from Statistical Bulletins of ICNAF for the period 1954 to 1966 (ICNAF Statistical Bulletin, 1956-1968).

As no data on water temperature were available from the Godthaab station since 1962 and from the station at $66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W}$ since 1966 , the data on catches for the last years were not included in the statistical analysis. Some relationships that are considered below were obtained from the data available by means of correlation analysis.
A. Water temperature anomalies in the $50-100 \mathrm{~m}$ layer in July (Fig. $1 a$ and the fishery importance of Div. IB in November

The relationship between the temperature anomaly ( $\Delta t$ ) in the $50-100 \mathrm{~m}$ layer on the station at $66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W}$ in July and the fishery importance (F) of Div. 1B in November is shown in Fig. 1b. The correlation coefficient $(r)$ is 0.851 , whereas the range length $N=13$, the error of correlation coefficient (E) is $0.051 ; \frac{r}{E}=16.6$, i.e., the relationship is real. The dependence is characterized by the following equation:

$$
\begin{equation*}
\mathrm{F}_{\overline{\mathrm{XI}}}=40.2 \Delta t \overline{\mathrm{VII}} 50-100+27.5 \tag{1}
\end{equation*}
$$

[^6]where:
\[

$$
\begin{aligned}
\Delta t \overline{\mathrm{VII}} 50-100= & \text { temperature anomaly in the } 50-100 \mathrm{~m} \text { layer on } \\
& \text { the station at } 66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W} \text { in July. } \\
& =\text { fishery importance of Div. } 1 \mathrm{~B} \text { in November. }
\end{aligned}
$$
\]

The relationship shows that if the temperature increases, then the fishery importance of Div. 1B will also generally increase. Table 1 illustrates the fitness of the equation (1).

Table 1. Actual and calculated values of fishery importance (F) of Div. 1B, anomalies and errors in percentage from the longterm amplitude $\left(A=F_{\max }-F_{\min }=60.7\right)$.

| Year | $\mathrm{F}_{\text {act }}$ | $\mathrm{F}_{\text {calc }}$ | $\mathrm{E}=\mathrm{F}_{\text {act }}-\mathrm{F}_{\text {calc }}$ | $\frac{\mathrm{E}}{\mathrm{A}}$ | $\frac{\mathrm{E}}{\mathbf{A}} \%$ |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1954 | 11.9 | 19.5 | - 7.6 | 0.12 | 12 |
| 1955 | 18.0 | 32.5 | -14.5 | 0.24 | 24 |
| 1956 | 15.2 | 11.4 | + 3.8 | 0.06 | 6 |
| 1957 | 4.6 | 8.7 | - 4.1 | 0.07 | 7 |
| 1958 | 2.5 | 11.4 | -8.9 | 0.14 | 14 |
| 1959 | 3.7 | 7.4 | - 3.7 | 0.06 | 6 |
| 1960 | 4.6 | 11.4 | - 6.8 | 0.11 | 11 |
| 1961 | 28.2 | 31.5 | - 3.3 | 0.05 | 5 |
| 1962 | 31.6 | 31.5 | $+0.1$ | 0.00 | 0 |
| 1963 | 50.5 | 47.6 | $+2.9$ | 0.05 | 5 |
| 1964 | 63.2 | 35.5 | +27.7 | 0.46 | 46 |
| 1965 | 41.7 | 43.6 | - 1.9 | 0.03 | 3 |
| 1966 | 53.7 | 51.6 | + 2.9 | 0.03 | 3 |

Equation (1) shows that in $84 \%$ of cases the error was not greater than $20 \%$ of the amplitude of the forecasting index.
B. Water temperature anomalies in the $100-200 \mathrm{~m}$ layer in July and cod diatribution in Div. 1B in November

Water temperature anomalies in the $100-200 \mathrm{~m}$ layer on the above-mentioned station $\left(66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W}\right)$ in July and fishery importance of Div. 1B in November (Fig. 2) were compared. The relationship between the comparative values (the length of the range $N=13$ ) is expressed in terms of the correlation coefficient $r=0.800 ;$ error $E=0.067 ; \quad \frac{r}{E}=11.9$, the equation is represented as:

$$
\begin{equation*}
F_{\overline{\mathrm{XI}}}=26.8 \Delta t_{\underline{\text { VIII }}} 100-200+23.7 \tag{2}
\end{equation*}
$$

where:

$$
\begin{aligned}
& \Delta t_{\overline{V I I} 100-200=} \begin{array}{l}
\text { water temperature anomaly in the } 100-200 \mathrm{~m} \text { layer } \\
\text { on the station at } 66^{\circ} 37^{\circ} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W} \text { in July }
\end{array} \\
& \mathrm{F}_{\overline{\mathrm{XI}}} \quad \begin{array}{l}
\text { fishery importance of } \mathrm{Div} .1 B \text { in November. }
\end{array}
\end{aligned}
$$

Table 2 illustrates the fitness of equation (2).

Table 2. Actual and calculated valucs of fishery importance (F) of Div. 1B, anomalies and errors in percentage from the longterm amplitude $\left(A=F_{\text {max }}-F_{\text {min }}=60.7\right)$.

| Year | $F_{\text {act }}$ | $F_{\text {calc }}$ | $E=F_{\text {act }}-F_{c a l c}$ | $\frac{E}{A}$ | $\frac{E_{\%}}{A}$ |
| :--- | ---: | ---: | :---: | ---: | ---: |
| 1954 | 11.9 | 10.3 | +1.6 | 0.03 | 3 |
| 1955 | 18.0 | 18.3 | -0.3 | 0.00 | 0 |
| 1956 | 15.2 | 18.3 | -3.1 | 0.05 | 5 |
| 1957 | 4.6 | 15.7 | -11.1 | 0.18 | 18 |
| 1958 | 2.5 | 15.7 | -13.2 | 0.22 | 22 |
| 1959 | 3.7 | 13.0 | -9.3 | 0.15 | 15 |
| 1960 | 4.6 | 13.0 | -8.4 | 0.14 | 14 |
| 1961 | 28.2 | 18.0 | +9.9 | 0.16 | 16 |
| 1962 | 31.6 | 21.0 | +10.6 | 0.17 | 17 |
| 1963 | 50.5 | 37.1 | +13.4 | 0.22 | 22 |
| 1964 | 63.2 | 34.4 | +28.8 | 0.49 | 49 |
| 1965 | 41.7 | 45.1 | -3.4 | 0.06 | 6 |
| 1966 | 53.7 | 69.3 | -15.6 | 0.26 | 26 |

Equation (2) shows that in $69 \%$ of cases the error was not greater than $20 \%$ of the amplitude of the forecasting index.

From Figs. 1 and 2 it may be seen that in separate years some anomalies in general regularity were registered. An especially large anomaly was observed in 1964: when water temperature in July decreased, the fishery importance of Div. IB in November decreased.

In accordance with the hydrological observations (Svetlov, 1966), water temperature in 1964 along almost the whole of the west coast of Greenland was above the norm. But in July on the station at $66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W}$, it was 0.1 $0.3^{\circ}$ below the average temperature of the $50-100 \mathrm{~m}, 100-200 \mathrm{~m}$ layers. It is supposed that the lower water temperature could be the result of the expansion of the cold Canadian Current which, in July 1964, came near this station; whereas, the heat level of 1964 for the whole Subarea 1 was fairly high (Bratberg, 1965; Hansen and Hermann, 1965; Meyer, 1965). Perhaps, therefore, the cod concentrations observed in Div. 1B to November caused a sharp increase of the fishery importance of this division.
C. Water temperature anomalies in the $50-100 \mathrm{~m}$ layer in July and fishery importance of Div. 1B in December

Comparison of the water temperature anomaly in the 50-100 m layer in July on the station at $66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W}$ and fishery importance of Div. 1B in December showed a direct relationship. The correlation coefficient $r=0.828$; $n=13 ; E=0.063$ (Fig. 3); $\frac{r}{E}=13$. The relationship is expressed by the
following formula: following formula:

$$
\begin{equation*}
F_{\overline{\text { XIII }}}=32 \Delta t \underline{\overline{\text { VII }} 50-100+23} \tag{3}
\end{equation*}
$$

where:

$$
\begin{aligned}
& \Delta t_{\overline{V I I}} 50-100= \\
& \text { water temperature anomaly in the } 50-100 \mathrm{~m} \text { layer } \\
& \text { on station at } 66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W} \text { in July }
\end{aligned} \quad \begin{aligned}
& \mathrm{F}_{\overline{\mathrm{XII}}} \quad=\text { fishery importance of Div. } 1 \mathrm{~B} \text { in December. }
\end{aligned}
$$

The values of the fishery importance (Table 3) were calculated for the assessment of the fitness of equation (3).

Table 3. Actual and calculated values of fishery importance ( $F$ ) of Div. 1B, anomalies and errors in percentage from the longterm amplitude $\left(A=F_{\max }-F_{\text {min }}=44.4\right)$.

| Year | $F_{\text {act }}$ | $F_{\text {calc }}$ | $E=F_{\text {act }}-F_{\text {calc }}$ | $\frac{E}{A}$ | $\frac{E_{q}}{A}$ |
| :--- | ---: | ---: | :---: | ---: | ---: |
| 1956 | 2.9 | 10.2 | -7.3 | 0.16 | 16 |
| 1957 | 0.6 | 5.8 | -5.2 | 0.12 | 12 |
| 1958 | 0.4 | 10.2 | -9.8 | 0.22 | 22 |
| 1959 | 0.1 | 7.0 | -6.9 | 0.16 | 16 |
| 1960 | 16.0 | 10.2 | +5.8 | 0.13 | 13 |
| 1961 | 28.7 | 26.2 | +2.5 | 0.06 | 6 |
| 1962 | 42.0 | 26.2 | +25.8 | 0.58 | 58 |
| 1963 | 41.1 | 39.0 | +2.1 | 0.05 | 5 |
| 1964 | 39.4 | 35.4 | +4.0 | 0.09 | 9 |
| 1965 | 14.5 | 35.8 | -21.3 | 0.48 | 48 |
| 1966 | 44.5 | 42.2 | +2.3 | 0.05 | 5 |

D. Water temperature of the $0-50 \mathrm{~m}$ layer on the Godthaab station in October and fishery importance of Div. 1B in June of the next calendar year

One more type of dependence between water temperature of the $0-50 \mathrm{~m}$ layer in October on the Godthaab station and fishery importance of Div. 1B in June of the following calendar year is represented in Fig. 4. The correlation coefficient ( $r$ ) of this relationship is $0.70,(N=10), E=0.11, \frac{r}{E}=6.4$, the
regression is:

$$
\begin{equation*}
F_{\overline{V_{I}}+1}=23.3 t_{\underline{\bar{x}} 0-50}-43.5 \tag{4}
\end{equation*}
$$

where:

$$
\begin{aligned}
\mathrm{t}_{\underline{\bar{x}} 0-50}= & \begin{array}{l}
\text { water temperature of the } 0-50 \mathrm{~m} \text { layer on the Godthaab } \\
\\
\text { station in October }
\end{array} \\
\mathrm{F}_{\overline{\mathrm{VI}} \mathrm{n}+1}= & \begin{array}{l}
\text { fishery importance of Div. } 1 \mathrm{~B} \text { in June of the following } \\
\\
\text { calendar year }(n-\text { year of temperature observations) } .
\end{array}
\end{aligned}
$$

The values of fishery importance calculated due to equation (4) are represented in Table 4.

In the calculations of the fishery importance due to equations (3) and (4) in $80 \%$ of cases the error was not greater than $20 \%$ of the amplitude of forecasting index. The estimate of the fishery importance value due to equations

Table 4. Actual and calculated values of fishery importance (F) of Div. 1B, anomalies and errors in percentage from the longterm amplitude $\left(A=F_{\max }-F_{\min }=46.9\right)$.

| Year | $F_{\text {act }}$ | $F_{c a l c}$ | $E=F_{a c t}-F_{c a l c}$ | $\frac{E}{A}$ | $\frac{E_{\%}}{A}$ |
| :--- | ---: | ---: | ---: | ---: | ---: |
| 1954 | 0.4 | 2.7 | -2.3 | 0.05 | 5 |
| 1955 | 1.8 | 3.1 | -1.3 | 0.03 | 3 |
| 1956 | 0.5 | 0.9 | -0.4 | 0.00 | 0 |
| 1957 | 4.8 | 8.0 | -3.2 | 0.07 | 7 |
| 1958 | 1.8 | 20.1 | -18.3 | 0.39 | 39 |
| 1959 | 47.3 | 23.4 | +23.9 | 0.51 | 51 |
| 1960 | 12.1 | 16.1 | -4.0 | 0.08 | 8 |
| 1961 | 18.8 | 17.3 | +1.5 | 0.03 | 3 |
| 1962 | 19.8 | 24.8 | -5.0 | 0.11 | 11 |
| 1963 | 7.1 | 5.2 | +1.9 | 0.04 | 4 |

(1-4) was made for those years which are indicated in Tables 1-4.
Asynchronous character of the above considered relationships is a peculiar feature. This fact is of great importance for fishery forecasting. It is known that fish do not react immediately; also, their migration from one area to another takes time (Konstantinov, 1967b). The equations of relationships obtained above provide a means of forecasting the fishery importance of Div. 1B from 4 to 8 months in advance.

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Fig. 1. Comparison between the fishery importance (F) of Div.iB in November and temperature anomaly in the $50-100 \mathrm{~m}$ layer on the station at $66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W}$ in July:
a) water temperature anomaly (1) and fishery importance (2),
b) the relationship between them.


Fig. 2. Comparison between the fishery importance ( $F$ ) of Div.lB in November and water temperature anomaly in the 100-200 m layer on the station at $66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W}$ in July:
a) water temperature anomaly (1) and fishery importance (2),
b) the relationship between them


Fig. 3. Comparison between the fishery importance ( $F$ ) of Div.1B in December and water temperature anomaly in the $50-100 \mathrm{~m}$ layer on the station at $66^{\circ} 37^{\prime} \mathrm{N}, 57^{\circ} 05^{\prime} \mathrm{W}$ in July:
a) water temperature anomaly (1) and fishery importance (2),
b) the relationship between them


Fig. 4. Comparison between the water temperature of the $0-50$ $m$ layer in October and the fishery importance ( $\overline{\mathrm{VI}}_{\underline{n}+1}$ ) of Div. $1 B$ in June of the following calendar year ( n - year of temperature observations):
a) water temperature (1) and fishery importance (2),
b) the relationship between them.

SECTION C
HADDOCK
7. On haddock spawning in the Northwest Atlantic area ${ }^{1}$
by V.P. Serebryakov VNIRO, Moscow

In the Northwest Atlantic area haddock are distributed from Greenland to Cape Hatteras. But, more or less detailed studies of haddock spawning have only been made for the areas of New England and Georges Bank. For the other areas there are some data on distribution of mature haddock and only scrappy data on distribution of eggs and larvae (Walford, 1938; Bigelow and Schroeder, 1953; Chase, 1955; Marak, 1960; Colton and Temple, 1961; Miller, Colton and Marak, 1963; Colton, 1965).

The author has tried to study the location of spawning areas as well as the drift and habitats of young haddock off Newfoundland and the Scotian Shelf. Data on maturity stages obtained by research vessels of the Polar Research Institute of Fisheries and Oceanography (PINRO) in field conditions in 1954 and in 1957-1963 were used in this paper. The maturity stages were determined as recommended for cod by Sorokin (1957). Ichthyoplankton samples were also collected in 1959-1963 by the research vessels of PINRO and by those of the Prospective Fish Scouting Service. Conical egg nets with $80-\mathrm{cm}$ diameter opening were mainly used during the experiments. Usually, a vertical haul was made from the bottom to the surface and a 15 -minute oblique haul while drifting or circling. A total of 2,000 stations were completed during the period of investigations in the Northwest Atlantic from Greenland to Georges Bank. Some haddock were caught in the cod and redfish fisheries near West Greenland, but they were scarce in number, and it was, therefore, impossible to obtain representative data on seasonal changes in gonad condition. Haddock eggs were observed in Greenland waters in April, May and June. The numbers are not high, no more than 11 per vertical haul.

In Labrador waters eggs were not observed, although the adult fish are found in trawl catches.

Off Grand Bank, the haddock fishery may be very effective in certain years. The possibility of haddock spawning in this area was suggested earlier by some scientists (Needler, 1930; Thompson, 1939; Grosslein, 1962). Haddock with gonads at the pre-spawning and spawning maturity stages were observed in this area from March to September. In June, the largest number of haddock with running eggs were observed in the southern part of the Grand Bank and on its southwestern slope and in May in the areas of Saint Pierre and Green Banks. Distribution of haddock eggs in the waters of Grand Bank indicate spawning in this area, but, as few eggs are observed, the spawning is not heavy. According to data on maturity stages and egg distribution, a mass spawning takes place in the areas of 3aint Pierre and Green Banks. The spawning peak is observed in late May-early June. Shestov (1967) considers that the spawning takes place mainly on the southwestern slope, at the border of ICNAF Div. 30 and 3 N , at depths of $60-160 \mathrm{~m}$ and in temperatures of $3.0^{\circ}-6.0^{\circ}$. According to Shestov, the western part of the southwestern slope of the Grand Bank is the second spawning area. None of the authors pays any great attention to spawning areas located on the southern slopes of Saint Pierre and Green Banks. But, from data on egg distribution, one can confirm that spawning in these areas is much heavier than in the areas of the Grand Bank. Main spawning areas near Saint

[^7]Pierre and Green Banks can be observed at depths of $80-250 \mathrm{~m}$, and spawning takes place in slope waters in temperatures of $3.0^{\circ}-6.0^{\circ}$. The location of haddock spawning areas in the Grand, Saint Pierre and Green Banks is given in Fig. 2.

In the period 1961-1966, PINRO conducted regular investigations on the distribution of young haddock off Newfoundland Banks. These investigations showed that the young haddock are observed on the southwestern slope of the Grand Bank (Nevinsky, 1962) and also that young haddock more than 15 cm in length were distributed over the area. Bulatova (1962) analyzed the catches of young haddock for 1954-1961 and concluded that the young specimens were distributed only on the southern slopes of Grand Bank and on its southern part and on the southern slopes of Saint Pierre Bank. The young, up to 25 cm in length, are distributed at lesser depths than fish $26-35 \mathrm{~cm}$ in length. Investigators, thus, concluded that the young haddock inhabit the southern part of Grand Bank and its southern slope and the southwestern slope of Saint Pierre Bank. The fingerlings were distributed mainly on the southern and southwestern slopes of Saint Pierre and Green Banks, with the greatest concentrations being observed in the southern part of Grand Bank (Fig. 3).

Thus, the largest concentrations of eggs, larvae and fingerlings of haddock are found on the southern and southwestern slopes of Saint Pierre and Green Banks.

Analysis of the age composition of the young haddock showed that from Saint Pierre and Green Banks to the eastward, i.e., to the southern part of the Grand Bank, the numbers of older specimens of the young increase in the catches while the one-year-olds decrease (Shestov, personal communication).

A total scheme of haddock spawning is suggested as follows. A single stock of Newfoundland Bank haddock spawns on the southern and southwestern slopes of Saint Pierre and Green Banks and on southern part of the southwestern slope of the Grand Bank at depths of $80-25 \mathrm{~m}$ and at near-bottom temperatures, $3.0^{\circ}-6.0^{\circ}$. Apparently, the eggs and larvae do either not drift from the spawning areas, or they drift to the southwest in the low velocity current and then are carried out to the areas adjacent to the spawning areas.

Correspondingly, the fingerlings settle to the bottom either in the spawning areas or in areas adjacent to them. The young move to Grand Bank, mainly its southwestern slope.

Some scientists consider that two independent haddock stocks inhabit the Newfoundland Banks (Grosslein, 1962), i.e., a Saint Pierre and Green Bank stock and a Grand Bank stock.

Sometimes Grand Bank haddock are found in good numbers. An insignificant spawning within the Bank waters can hardly ensure a great abundance for an independent stock. In addition, in the case of the availability of an independent stock, a great number of haddock fingerlings should be observed in this area.

Shestov (1967) has suggested that a single stock of haddock inhabits the waters of Grand, Saint Pierre and Green Banks, whose origin is from the spawning areas of the southern and southwestern slopes of Saint Pierre and Green Banks. Data obtained by the author confirm this suggestion.

Two stocks inhabit the waters of the Scotian Shelf, one of them is located in the northeastern part of the Shelf, north of the Scotia Deep, the other one on the southwestern part of the Shelf.

Spawing haddock are observed on the Scotian Shelf in April, May, June (Fig. 1). Dannevig (1919) noted the distribution of larvae in the Sable Island area in May. Thompson (1939) showed the distribution of haddock fingerlings on Banquereau Bank. Young haddock are distributed over Misaine and Banquereau Banks, southeast of Sable Island over the Shelf slope and on Emerald Bank (Fig. 3).

Grosslein (1962) showed that the peak of spawning on the Scotian Shelf is in April. The distribution of haddock eggs in May testifies to the fact that spawing probably takes place mainly in May. It should be said here that in April the survey program took place only partially on the Scotian Shelf.

Usually, haddock spawning takes place sometime later than that for cod in the same areas, and it is unlikely that the Scotian cod make an exception.

The drift of haddock eggs and larvae is determined by the direction and the speed of the current, flowing out of the Gulf of St. Lawrence and dominatIng the Scotian Shelf. The speed of this current is high - up to $94 \mathrm{~cm} / \mathrm{sec}$ over Misaine and Banquereau Banks (Klimenkov and Pochorukov, 1962). In the case where the eggs are drifting in the upper 20 -m water layer, they will be carried in the water masses to the southwest along the Shelf (Walford, 1938; Miller, Colton, and Marak, 1963; Colton, 1965). But, sometimes the upper water layer (up to 100 m in depth) on the Scotian Shelf is freshened ( $<31 \%$ 。) and becomes cool (Klimenkov and Pochorukov, 1962). The density of this layer is, naturally, decreasing. Kisljakov (1958) showed that the cod eggs are drifting with the deeper layers when the upper layer is freshened. The behaviour of haddock eggs should be the same in similar environmental conditions. Thus, it can be suggested that haddock eggs and larvae are drifting In the deep water layers. The rate of speed is less at great depths (Colton and Temple, 1961) and the temperature higher. Correspondingly, the rate of development is greater and the duration of the period of the pelagic development is less. This means that, in spite of a rather high current speed and low temperatures in the upper water layers over the Scotian Shelf, the eggs and larvae are not carried of $f$ the limits of the Shelf. In any case, such conclusions can be made for ICNAF Div. 4V, 4W. It is possible that they are not carried far away from the spawning areas. This can be confirmed from the distribution of haddock fingerlings off Emerald Bank, in the areas where the greatest catches of eggs were taken.

Discussion of results
In the northwest part of the Atlantic Ocean haddock are spread from Labrador to Cape Hatteras, they can also be observed near Greenland. In the ICNAF Area, haddock are most abundant off Nova Scotia and New England. Some local stocks of haddock are observed in these areas, they are characterized by different number of vertebrae. Previously, scientists supposed that two haddock stocks existed off Newfoundland. From the above data it is possible to say that only one haddock stock Inhabits the Newfoundland Banks area. This stock originates from the spawning areas located in the southwestern and southern slopes of Saint Pierre and Green Banks and in the western part of the southwestern slope of the Grand Bank. Because of drift conditions, haddock of

Newfoundland Banks are isolated from the Scotian Shelf. Stocks of adult fish of these areas probably mix very little, as the number of vertebrae differs greatly enough. The location of spawning areas on the one hand, and the drift pattern of eggs and larvae on the other hand, suggest that there are two haddock stocks inhabiting the northeastern and southwestern parts of the Shelf respectively. The number of vertebrae for haddock of these two stocks differs, though not greatly, a fact which suggests there is considerable mixing of these two stocks. It is characteristic that the greateat number of vertebrae (54.19) are from haddock inhabiting the shores of the Cape Breton Island, the area of the lowest water temperature during the spawning period. An independent haddock stock inhabits the area of Georges Bank, the number of their vertebrae is almost the same as for haddock of the Newfoundland Banks.

In the Northwest Atlantic haddock spawn from February to July. The spawning peak does not coincide in time for different areas. Off Newfoundland Banks, spawning is observed late in May-early in June. In the Nova Scotia area, spawning takes place in May. On Georges Bank, spawning reaches its apogee in May-April, though haddock spawn there in January and even in December, and some spawning individuals and eggs are observed in July and in August.

Thus, in the Northwest Atlantic the length of the spawning period for haddock is greatest in southern areas, on Georges Bank. In the northern areas, the spawning period begins earlier and is shorter. Haddock spawning is observed at depths from 40 m to 250 m in $3^{\circ}-6^{\circ}$ temperature. As noted above, the lowest temperatures were registered for the Scotian Shelf. Comparing the location of the spawning areas to the distribution of near-bottom temperatures during the spawning or pre-spawning periods, it can be concluded that the time of haddock spawning in the various areas is determined by the meandering of the warm currents. This is especially apparent on the southern slopes of Saint Pierre, Green Banks and on the western part of the southwestern slope of Grand Bank.

The haddock eggs and larvae drift is unique for each different area of the northwestern Atlantic. In the areas off Newfoundland Banks, eggs and larvae are carried off the spawning grounds to the southeast. Their drift pattern is like that of an incomplete circle. The shorter the distance from the spawning area to the centre of the circular water motion, the shorter is the drift of eggs and larvae from their spawning area. The centre of the current is probably displaced in different years and, in this connection, the picture of eggs and larvae drift changes as well. But, those changes probably do not affect the general character of the drift.

In the Nova Scotia area, the drift may be determined by a strong Cabot Current, as was shown earlier. In this case, eggs and larvae of the Scotian Shelf are carried from the northeastern part of spawning areas to the southwest, and from the spawning grounds of $f$ Browns and LaHave Banks northwesterly, i.e., to the eastern part of the Gulf of Maine and Passamaquoddy Bay. When eggs and larvae are not in low salinity surface water layers, but in deeper layers with an oceanic salinity, they will not be carried a great distance. In all cases, the extent of drift in space and time is not too great, being much less than that for cod in the northern areas.

During the passive pelagic period of life, survival depends greatly on the whole complex of different conditions and on the areas, where larvae have been carried at the fingerling stage, when the bathypelagic and near-bottom mode of life begins. It can be supposed that this process is determined by
the current intensity, when the eggs and larvae are being carried by powerful currents like that in the North Atlantic.

The drift of eggs and larvae may depend on wind currents. This is observed, when they are drifting in waters with slow current speed, i.e., in turbulences and rotations, which, in their turn, depend on wind conditions. Carruthers (1951) investigated the influence of the wind conditions on the abundance of haddock year-classes in the North Sea for the period 1929 to 1949. It appeared that relatively rich year-classes prevailed in the years when northerly and earterly winds dominated, but they were poor in the years with mainly northerly and westerly winds. Of 20 year-classes, only two (1946 and 1947) were exceptions to this regularity.

In the Northwest Atlantic, analogous investigations were made for Georges Bank haddock. Chase (1955) obtained a correlation coefficient of $0.766 \pm 0.057$ between the size of haddock year-classes and favourable winds for the period of investigations 1928 to 1952.

Of course, the strength of a year-class is determined not only by the wind regime and this factor is of great significance only under certain conditions.

It can be considered that the survival of eggs, larvae and fingerlings is more closely related to wind when the spawning grounds are adjacent to areas with environmental conditions unfavourable to their development, and if they are brought there they will die. This conclusion can be related to the haddock spawning areas off Faroes (Saville, 1965) and Georges Bank. The spawning areas located on Saint Pierre, Green Banks and on the southwestern slope of Grand Bank are affected by the factors described above.

The winds from the south and southwest provide favourable conditions for haddock development on Newfoundland Banks during their spawning and drifting periods, and those from the north and northeast - the unfavourable ones. Thus, it is possible to make some preliminary conclusions. If the winds blow from the south, the year-classes will be abundant in April-October; if they blow from the north, they will be poor.

In conclusion, it may be said that some other conditions for survival are overlapping the effect of wind, especially those of extremely warm or cold years.

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Fig. 2. Distribution of haddock spawning areas. Conventional signs:

1. Intensive spawning
2. Not intensive spawning
3. Spawning areas, preliminary determination


Fig. 3. Distribution of young haddock Conventional signs:

1. Distribution of fingerlings and one-year-olds
2. Distribution of young specimens


Fig. 4. Distribution of spawning grounds, eggs, larvae, fingerlings and migration paths of haddock Conventional signs:

1. Spawning grounds
2. Egg distribution
3. Larvae distribution
4. Fingerling distribution
5. Drift paths
6. The retention inside the Bay of Fundy of herring larvae spawned off the southwest coast of Nova Scotia
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## Introduction

The seasonal distribution of larval herring in relation to the physical and biological environment in the Bay of Fundy-Gulf of Maine region for the years 1967 to 1969 as shown by cruises at three-month intervals has been described by Tibbo and Lauzier (1970). Three additional short cruises at approximately fortnightly intervals were carried out in the fall of 1969 to follow in more detail the dispersal of larvae from spawning grounds off the southwest Nova Scotia coast, this being one of the major sources of larvae in the region (Tibbo, et al., 1958). The area studied is now the scene of a large fishery for pre-spawning and spawning adults (Iles and Tibbo, 1970) but the extent to which the stock contributes to the juvenile fisheries for herring in the Bay of Fundy and Gulf of Maine is not known with certainty and more detailed knowledge of larval movement from the spawning area was thought likely to make a contribution towards the solution of this problem.

## Methods

The cruises were undertaken from the St. Andrews Biological Station of the Fisheries Research Board of Canada on M/V E.E. Prince during the periods 7-10 October, 28-30 October, and 14-17 November. The two October cruises covered the entrance to the Bay of Fundy and an area off the Nova Scotia coast north of latitude $44^{\circ} 32^{\prime}$; the November cruise surveyed the southern side of the entrance and much of the Bay of Fundy itself.

As far as possible sampling was based on a rectangular grid, station distances varying with circumstances from about five to about ten miles. An Isaacs-Kidd $2-\mathrm{m}$ traw1 ( 1.6 mm codend mesh) was towed at 5 knots and most of the water column was sampled.

A telemetering device on the trawl allowed the calibration of gear depth to warp length so as to estimate the leeway required to avoid bottom contact. About 5 m was allowed in shallow water and a little more than 10 m for deeper water. It was assumed that the water column above these depths was adequately sampled. The gear was lowered rapidly to the maximum depth of the tow and hauled step-wise during the 30 -minute towing period to approximate an oblique tow.

Bathythermograph records were taken at each station and surface temperatures and salinities measured.

Plankton samples were sorted and analyzed at the St. Andrews laboratory. Fish larvae were identified and counted and total lengths of herring larvae were measured on the whole of small samples and on at least 100 of larger samples. Estimates of numbers of invertebrates in broad systematic categories were made for each station.

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Larval escapement from gear
The high towing speed was designed to reduce larval avoidance of gear to a minimum. Such avoidance is usually demonstrated by markedly increased catches at night compared with day-time catches (Bridger, 1956). Table l lists day and night catches (numbers per tow) for each of the three cruises and for all taken together. There were no systematic differences that cannot be easily explained on the basis of differences in larval abundance in areas sampled at day and at night time.

Table 1. Comparison of day and night catches of herring larvae.

|  | Day |  | Night |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Cruise | Tows | Average <br> No/Tow | TowsAverage <br> No/Tow | Total Number <br> of Larvae |  |
| $7-10$ October <br> $27-30$ October <br> $14-17$ November | 28 | 465 | 22 | 283 | 19,242 <br> 18 |
| 126 | 16 | 468 | 10,318 <br> 6,307 |  |  |
| All cruises <br> together | 76 | 266 | 48 | 324 | 35,867 |

That larval escapement was certainly reduced for the Isaacs-Kidd trawl is indicated by comparison of catches made at the same day-time station by a 1 m ( 60 mesh/inch) net hauled vertically (and thus more slowly) and the IsaacsKidd trawl hauled at 5 knots. Only larvae less than 14 mm total length were caught in the $1-m$ net, whereas a full size range up to $26+m m$ was caught in the Isaacs-Kidd.

Herring larvae develop considerable short-burst swimming power at 15 m as the caudal fin develops (Blaxter, 1962).

## Results

Larval distribution and abundance

Larval numbers were plotted for each station and contour lines drawn at different levels of abundance to give results shown in Figs. 1-3 for the three cruises separately.

The results for the first cruise showed two areas of concentration of larvae, a small one at the southwest end of the Grand Manan Channel and a much larger and more dense one off the southwest coast of Nova Scotia which blankets the known spawning grounds of the Nova Scotia fall spawners (Tibbo, et al., 1958; Das, 1968). Within this area the highest concentrations occurred a few miles to the northwest of Trinity Ledge and a similar distance from Lurcher Shoal.

The westerly and southerly limits of this larval concentration were clearly delimited; if, as is thought likely, the main spawnings occur on

Trinity Ledges and Lurcher Shoals, then there was a general dispersion from this area but movement out of the area was restricted to the north and east; towards St. Mary's Bay and, on the outside of this Bay, along the north shore of Digby Neck into the Bay of Fundy proper. There is little evidence of larval movement across the entrance of the Bay of Fundy towards Grand Manan, nor on the northern side, of larvae moving out of the bay along the New Brunswick coast towards the Grand Manan channel. It should be noticed that relatively few larvae were found at the head of St. Mary's Bay itself.

By the end of October larval numbers had declined sharply (Fig. 2) both because of mortality and movement out of the area, but it is not possible to estimate the effect of each of these separately. The areas of high concentration had contracted, and the one immediately off the Nova Scotia coast had split up into northerly and southerly components. The reduction in larval numbers was much more marked in the Lurcher Shoals area and the western section of the survey area. An overall increase in density at the head of St. Mary's Bay towards its north side was indicated, however, suggesting a net accumulation of larvae there.

Again, there was little sign of transport across the entrance of the Bay of Fundy and the concentration of larvae at the southwest entrance to the Grand Manan channel was still evident and had moved only a very short distance.

The November cruise, extending into the Bay of Fundy, completed the pattern suggested by the two earlier cruises. The path of transport of larvae along the northern shore of Digby Neck could be seen as confluent with a large area of concentration centered at a point 15 miles southwest of Saint John and extending northwards and eastwards, up to Chignecto Bay and Minas Basin (Fig. 3). Densities were relatively high, fmplying an accumulation and retention of larvae in the area. There was also indication of a more even spatial dispersal of larvae in this area which is demonstrated in Fig. 4. This campares, for the three cruises, the frequency distribution of numbers of larvae/tow, divided into conveniently grouped cells. A higher proportion of the tows for the November cruise has high numbers of larvae and the frequency distribution was less asymmetric than for the two October cruises.

The westward limit of distribution was a little west of Saint John; no larvae were recorded west of Point Lepreau, so that again, there was no indication of transport of larvae out of the Bay of Fundy along the northern side.

Larval length distributions
Inspection of length-frequency distributions of larvae from individual tows for the first October cruise indicated size differences between areas. Five areas were separated; around Grand Manan, the southern part of the area off the southwest Nova Scotia coast, the entrance to St. Mary's Bay, the southern entrance to the Bay of Fundy, and lastly, the head of St. Mary's Bay, i.e., above the passages connecting the bay to the Bay of Fundy. Similar area differences were found for the results of the second October cruise and data for both cruises are presented for each area in Fig. 5.

The smallest larvae were found near the entrance to St. Mary's Bay, i.e., near Trinity Ledges; most were below 13 mm total length at the beginning of October and represent, probably, the products of a recent spawning in the vicinity. Larvae to the south and west of St. Mary's Bay were larger and those
nearer the entrance to the Bay of Fundy and around Digby Neck larger still. Considering these three groups together, there was a suggestion, at least, of three modes each, possibly, representing larvae spawned at different times during the season. For the first October cruise larvae at the head of St. Mary's Bay were distinctly larger than those at the mouth; very few were smaller than 15 mm total length.

Finally larvae in the Grand Manan area were larger on average than in the other areas and showed a much wider range of size, from 11 mm to 34 mm total leng th.

By the end of October larvae in each of the areas were larger, although only marginally so for those at the head of St. Mary's Bay. The differences in average size for the other areas varied from about 3 mm to about 7 mm which is consistent with larval growth rates given by Tibbo, et al. (1958) and Das (1968) for larvae from the same general region.

There was no indication of an influx of smaller larvae in any area so that it is highly improbable that any new spawning had occurred.

The length distribution for the November cruise had a wide range, from about 10 mm to about 40 mm and had also a more even spread of length (Fig. 5); this is consistent with the hypothesis that these larvae represented accumulation of larvae transported into the Bay from Nova Scotia spawnings over the spawning season as a whole, or over a large part of it.

## Discussion

Evidence has been steadily accumulating that larvae from fall spawnings off the southwest Nova Scotia coast are transported northwards and thence northeast along the southern side of the Bay of Fundy along a non-tidal inflow associated with the Gulf of Maine eddy (Tibbo, et al. 1958; Tibbo and Legare, 1960; Das, 1968; Tibbo and Lauzier, 1970).

This study demonstrates such a transport and indicates that the path of transport might be quite narrow (Fig. 2). However, a corresponding transport out of the bay along the northern New Brunswick side, which would be associated with the same current system (Tibbo and Lauzier, 1970) is not indicated by these results. Instead, there is evidence, both from the distribution and abundance of larvae and from length frequency distributions, that larvae transported into the bay on the southern side are retained in the shallow areas of the upper part of the bay. This could be the result of a slackening off of the current system which begins in the fall (Bumpus and Lauzier, 1965), and results in almost a closed circulation within the Bay of Fundy during the winter. It is now suggeated that larvae retention inside the Bay of Fundy may involve larval behavioural and activity patterns of the kind described by Graham (in press) to account for overwintering larval concentrations in estuaries along the Maine coast.

That such mechanisms may exist is suggested by the retention of larvae at the head of St. Mary's Bay. It was shown above that in early October most of the larvae at the head of this bay exceeded 15 mm in length, the same length at which the development of the caudal fin confers a marked increase in swimming ability, whereas, larvae at the entrance to the bay were smaller, implying that maintenance of position is a function of swimming ability. This does not
imply that St. Mary's Bay is itself a major overwintering area for the larvae of the Nova Scotia stock, neither its size nor the density of larvae found there match up with the size of the stock or of the fishery it supports (Iles and Tibbo, 1970), although it is not excluded as a possibility that larger densities of larvae may occur in the bottom 5 m , not sampled by these surveys.

The occurrence of larval concentrations during the fall, at the southwest end of the Grand Manan channel might be construed as evidence of transport out of the Bay of Fundy but the similarity in length distributions between early and late October samples (and allowing for intervening growth) noted above makes it far more likely that these represent a relatively stationary population originating in the area itself. Das (1968) cites instances of spawning being recorded in this area.

On the other hand, larvae have been reported well inside the Bay of Fundy during the fall and winter on many occasions. Tibbo, et al. (1958) record such occurrences in the fall of 1956 and in the fall and winter of 1957-58, Tibbo and Légare (1960) in the fall and winter of 1958-59, Das (1968) in early 1961 and in the fall of 1962, and Tibbo and Lauzier (1970) in the autumn and winter of 1967-68 and 1968-69.

The question then arises whether the upper part of the Bay of Fundy is the major area for overwintering larvae fram the Nova Scotia spawnings and if so, what is the pattern of movement of post larval fish after their first winter. It is assumed by Das (1968) that the movement of these post larvae will be dictated by the non-tidal drift current system set up in the area which, as it develops in the spring, would carry any overwintering larvae out along the New Brunswick coast. It is suggested now that passive drift by larvae, although undoubtedly important during the early stages of development, becomes less important as they develop even relatively modest swimming powers, so that the dispersal of the young of the Nova Scotia stock may not be as widespread as has been supposed. Certainly, "brit" (yearling) herring were also caught during the November cruise well inside the Bay of Fundy and that a return of brit and sardine (two-year-old) herring may be in the direction of the Nova Scotia side of the entrance to the Bay of Fundy cannot be excluded. Iles (1970) produced some evidence to question the commonly held hypothesis that the Nova Scotia spawning provides recruits to the sardine fishery of the New BrunswickMaine coast, suggesting that sardine populations on both sides of the Bay of Fundy are distinct. While neither that evidence, nor the implications of this study are in any way conclusive, they do suggest that more detailed study of the overwintering larval populations of the Bay of Fundy and attempts to complete the life cycle by adequate sampling of later stages are likely to contribute much to what is still the major herring problem in the region - the origin of the sardines of the Bay of Fundy-Gulf of Maine.

If the hypothesis suggested by this study can be substantiated, it will have important implications.

The Nova Scotia fall spawning stock would neither contribute to nor be directly affected by the Gulf of Maine-New Brunswick fisheries for brit (yearling) and sardine fisheries and could be treated as a convenient unit for management purposes. Again, if the Bay of Fundy is a major overwintering area for larval herring, opportunity is created for the study of larval ecology of a discrete population, and the possibility of estimating total abundance at late larval or subsequent stages, if these can be adequately sampled, might allow prediction of subsequent recruitment levels.

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Fig. 1. Herring larval distribution and abundance in the Bay of Fundy Nova Scotia southwest coast area. 7-10 October 1969.


Fig. 2. Herring larval distribution and abundance in the Bay of Fundy Nova Scotia southwest coast area. 28-30 October 1969.


Fig. 3. Herring larval distribution and abundance in the Bay of Fundy Nova Scotia southwest coast area. 14-17 November 1969.


Fig. 4. Frequency distribution of number of larvae per tow for the three cruises.


Fig. 5. Comparison of larval length frequency distributions for the three cruises (see text).
9. Status of the Southwest Newfoundland Herring Stocks, 1965-70 ${ }^{1}$

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## Introduction

Prior to 1965 the Newfoundland herring fishery was associated largely with the demand for herring as bait for the cod fishery and the sporadic demand for pickled herring products as food, especially during and inmediately following World Wars I and II. In 1946 there was a peak landing of 75,000 metric tons, all of which were taken in coastal waters by gillnets and beach seines. Subsequently annual landings decreased to less than 10,000 tons in the early 1960's when the demand for herring as food was very low and most of the herring caught were utilized as bait. Purse-seining was introduced to the south coast of Newfoundland in the autumn of 1964 and landings from that area rapidly increased to 130,000 metric tons in 1968. This report considers the upsurge in herring landings in the light of recent knowledge on the size, distribution, biology, and migratory behaviour of the stocks.

## Mobile fleet landings

In the autumn and winter of 1964-65 large concentrations of herring were found to overwinter in the fjords along the south coast of Newfoundland between Port aux Basques and Bay D'Espoir (Fig. 1, Areas Jl and J2). From a single fishing vessel in 1964 the seiner fleet participating in the winter fishery increased rapidly to 55 seiners during the 1969-70 season. In addition $10-15$ vessels acted as carriers for some of the smaller seiners. Most of the herring are landed at processing plants at Harbour Breton, Burgeo and Isle aux Morts, and seiners account for more than $90 \%$ of the Newfoundland herring yield annually.

The southwest Newfoundland fishery begins in November and continues uninterrupted until April (Table l). In July to October 1969 herring taken in the southwest part of the Gulf of St. Lawrence were landed in Newfoundland for the first time. Annual seiner landings increased from 7000 metric tons in 1965 to nearly 160,000 tons in 1969. The decrease to 148,000 tons in 1970 is largely due to greatly decreased catches in the autumn of that year.

The major winter fishery occurs mostly in the fjords and bays along the south coast in Areas Jl and J2 (Fig. 1, Table 2), which yielded 127,000 tons in 1969 and 119,000 tons in 1970. Up to 1968 the fishery took place largely in Area Jl, but during the winter of 1969 and more so in 1970 the western half of the southwest coast (J2) yielded the largest quantity of herring.
$\overline{1}$ submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res. Doc.71/121.

Seiner landings from catches in other parts of the Newfoundland area have been relatively small and the fisheries based on localized stocks. Along the west coast of Newfoundland (Areas $K$ and M) seiner landings ranged between 3000 and 6000 tons annually with most of the herring taken in November and December in or near Bonne Bay up to 1968 and more recently in Hawke's Bay. The small Fortune Bay stock (Area I), which yielded nearly 12,000 tons to seiners in 1968 but less than 5000 tons in 1969, showed some improvement in the winter of 1970 with a yield of 8000 tons (Table 2). However, the small stock in Areas $G$ and $H$, which yielded over 6000 tons to seiners in the spring of 1968 remained at a very low level of abundance in 1969 and 1970. This stock was greatly affected by phosphorus poisoning in Placentia Bay (H) during the winter and spring of 1969 when a high proportion of the adult population (estimated at more than $80 \%$ ) died.


Fig. 1. Map of the Gulf of St. Lawrence-Newfoundland area for reference to place names and statistical areas mentioned in the text.

Table 1. Monthly distribution of seiner landings (metric tons) in Newfoundland, 1965-70.

| Month | 1965 | 1966 | 1967 | 1968 | 1969 | 1970 |
| :--- | :---: | :---: | :---: | :---: | :---: | ---: |
| Jan. | 395 | 3,285 | 16,807 | 23,379 | 42,028 | 51,691 |
| Feb. | 763 | 4,098 | 11,962 | 22,870 | 33,118 | 30,833 |
| Mar. | 303 | 4,445 | 10,057 | 22,814 | 21,998 | 31,014 |
| Apr. | 313 | 1,864 | 11,848 | 11,294 | 8,645 | 14,055 |
| May | 392 | - | 1,375 | 2,501 | 2,377 | 1,963 |
| Jun. | - | - | - | - | $-6 \overline{1}$ | - |
| Jul. | - | - | - | - | 1,686 | 477 |
| Aug. | - | - | - | - | 3,478 | 332 |
| Sep. | - | - | - | - | 737 | 31 |
| Oct. | - | - | - | 1,057 | 552 | 220 |
| Nov. | 1,534 | 4,013 | 7,247 | 12,631 | 11,071 | 1,873 |
| Dec. | 3,264 | 8,257 | 22,778 | 46,438 | 33,564 | 15,211 |
| Total | 6,964 | 25,962 | 82,074 | 142,984 | 159,254 | 147,700 |

Table 2. Area distribution of seiner and midwater trawler landings (metric tons) in Newfoundland, 1965-70.

| Arȩa | 1965 | 1966 | 1967 | 1968 | 1969 | 1970 |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: |
| GH | 23 | 612 | 1,302 | 6,278 | 220 | 290 |
| I | 19 | - | 4,567 | 11,662 | 4,827 | 7,971 |
| J1 | 3,331 | 15,324 | 57,102 | 82,918 | 42,906 | 30,036 |
| J2 | 472 | 4,054 | 10,408 | 26,450 | 84,020 | 88,612 |
| K | - | 492 | 672 | 1,793 | 240 | 28 |
| M | 3,119 | 5,480 | 5,452 | 3,903 | 2,340 | 2,946 |
| Nfld. | 6,964 | 25,962 | 79,503 | 133,004 | 134,553 | 129,883 |
| 4 T | - | - | 2,571 | 9,633 | 21,503 | 14,741 |
| 4V | - | - | - | 347 | 3,198 | 3,076 |
| T+V | - | - | 2,571 | 9,980 | 24,701 | 17,817 |
| Total | 6,964 | 25,962 | 82,074 | 142,984 | 159,254 | 147,700 |

In recent years the seiners (and occasionally midwater trawlers) landings at Newfoundland ports have enlarged their sphere of fishing activity to the southern Gulf of St. Lawrence ( 4 T ) and northeastern Nova Scotia (4V). Division 4 T catches (Table 2) came mostly from the Magdaien Islands (Bird Rocks) and northern Cape Breton (St. Paul Island), and the $4 V$ catches from Sydney Bight and Chedabucto Bay. In 1969 Newfoundland herring landings from these regions totalled 25,000 tons, taken mostly in November and April, but in 1970 the yield was substantially less due to a poor fishery at Magdalen Islands in the fall.

Catch statistics for the 1971 winter fishery along southwest Newfoundland (Areas Jl and J2) are not yet fully available, but information from plant officials and seiner captains indicate that the herring yield may not exceed one-half of that recorded for the winter of 1970.

## Status of the Southwest Newfoundland herring stock

Investigations began in 1965 to elucidate the size, distribution, biology and migratory behaviour of the herring stocks which support the winter purse seine fishery (Hodder, 1967; Hourston, 1968). These were intensified in 1969. Efforts were continued to improve the collection of information on area of capture through log book records and port interviews. The sampling of seiner landings was carried out at the major ports of landing and several thousand herring were examined annually for information on their biology. During the past 2 years the seasonal migratory behaviour of the stock has been determined by meristic studies and tagging.

There has been a gradual increase in the average size of herring taken along southwestern Newfoundland since the purse seine fishery began in 1965 (Fig. 2). The modal size (total length to the 0.5 cm below) increased from 32 cm in the winter of 1966 to 34 cm in the autumn and winter of 1970-71. Similarly the average length increased by about 2 cm over the 6-year period. More than $95 \%$ of the herring were in the 30-36 cm length range.

The composition of the samples by maturity stages indicates a mixture of spring and autumn spawners, the latter comprising two-thirds or more of the seiner catches. Immature herring were rare in the samples. The relative proportions of spring and autumn spawners in the samples taken for ageing are given in Fig. 3.

Age determinations, based on otolith readings, reveal that the southwest Newfoundland winter fishery since its inception has been largely dependent on a very abundant 1958 year-class of autumn spawners and a smaller 1959 year-class of spring spawners (Fig. 3). Both the length and age frequencies indicate a lack of substantial recruitment to the adult stock.


Fig. 2. Length composition of herring sampled from the southwest Newfoundland fishery, 1965-66 to 1970-71 seasons.


Fig. 3. Relative age composition of spring (S) and autumn (A) spawning herring sampled from the southwest Newfoundland fishery, 196566 to 1969-70 seasons.

The predominance of autumn-spawning herring in the great concentrations which arrive in southwestern Newfoundland in November indicate that autumn spawning must be very extensive in Northwest Atlantic waters. However, no extensive autumn-spawning aress in Newfoundland waters were known prior to the development of the purse-seine fishery. In the autumns of 1968 and 1969 surveys along southern Newfoundland for recently hatched herring larvae produced no evidence of extensive spawning, despite the large numbers of autumn spawners which overwinter in the fjords there.

The distribution of seiner catches at the start of each season indicate that herring arrive from the westward in late November and within a few days are distributed widely in the fjords along the southwest coast. After January the fishery gradually shifts westward and the last catches for the season are usually made off the southwest corner of Newfoundland in April (Fig. 4). For about 3-4 weeks prior to the appearance of herring in southwest Newfoundland waters in late November, there is a fairly intense fishery at Magdalen Islands. Also, about a week after the herring leave Newfoundland in April, a short spring fishery again occurs in an area between St. Paul Island and Magdalen Islands, followed by a substantial summer fishery in the southwestern part of the Gulf of St. Lawrence during June to September.

In 1969 herring samples from seiner catches at Magdalen Islands in November were compared with samples taken from catches in the coastal waters of southwestern Newfoundland in late November and early December (Hodder and Parsons, 1970). In summary the study showed that there was no significant difference between areas for all of the biological characters examined (length, age, maturity condition, vertebral numbers, pectoral, dorsal and anal fin ray numbers, and Anisakis nematodes), although for nearly all of the characters very significant differences were apparent between spring and autumn spawners which were present in the samples from both areas in about the same proportion. The analysis thus supported the hypothesis that the winter fishery along southwest Newfoundland is largely dependent on herring schools which migrate eastward from the southern part of the Gulf of St. Lawrence in the autumn.

Both the eastward movement of herring from the southwestern Gulf of St. Lawrence to southern Newfoundland in the autumn and the westward movement in the spring have recently been confirmed by tag recaptures from more than 80,000 tagged herring which were released in March, May and August 1970 in southwest Newfoundland, at Magdalen Islands and near the Gaspé Peninsula respectively (Hodder and Winters, 1970; Winters, 1971a; Beckett, 1971). The absence of tag recaptures from a substantial purse seine fishery in Chedabucto Bay during the winter of 1971 (J.S. Beckett, 1971) and the recovery of only 2 tags during the 1971 winter fishery along southwest Newfoundland from the liberation of 3400 tagged herring in the northeastern part of the Gulf of St. Lawrence in early December 1970 (G. H. Winters, personal commication) suggest that the southern Gulf of St. Lawrence-southwest


Fig. 4. Monthly distribution of seiner catches along southwestern Newfoundland during the 1969-70 season. (Each full dot represents 1000 tons of herring.)

Newfoundland stock complex of adult herring does not deviate significantly from its seasonal east-west migration route, nor does it appear that the stock, at least when it is in southwest Newfoundland waters, is supplemented by a significant influx of herring from other areas.

Preliminary estimates from tagging data indicate that the stock size of the southwest Newfoundland herring population was about 400,000 tons at the start of the 1969-70 winter fishery and about 250,000 tons at the beginning of the 1970-71 season (Winters, 1971b). The "exploitation rate" was estimated at $4-5 \%$ per month. If a similar rate applies for the surmer fisheries in the southern Gulf of St. Lawrence, the annual rate for the stock as a whole would be about 40-50\%.

## Conclusions

The recent increase in Newfoundland herring landings is due to the development of a substantial autumn and winter purse seine fishery in the fjords along the western part of the south coast. Concurrent with the development of this fishery was a rapid expansion of the surmer fishery in the southern part of the Gulf of St. Lawrence. Consequently, the question arises as to whether or not these fisheries can continue successfully if the same stocks are being fished almost continuously at different times and places along their migratory route.

Recent studies show that herring samples taken during the autumn along southwestern Newfoundland differ in certain biological features from samples taken in other areas of Newfoundland and Labrador and on the northern part of the Scotian Shelf, but are very similar to those taken at Magdalen Islands just prior to the start of the Newfoundland autumn and winter fishery. In the spring the herring schools migrate westward to Magdalen Islands and later in June and July to the Gaspé Peninsula-Chaleur Bay region of the Gulf of St. Lawrence.

About the time that the herring leave the Newfoundland coast in the spring some separation of spring- and autum-spawning herring is evident from samples taken from seiner catches near the end of the season in early April. The short fishery at Magdalen Is lands in late April and early May is based mostly on herring schools heading for shallow water to spawn, but a few catches consist mostly of autumn spawners. However, the latter probably disperse to a large extent and ultimately move westward to the soutnwestern part of the Gulf, where they feed during the summer and spawn in late summer and early autumn.

The winter fishery along southwest Newfoundland is almost entirely on adult herring, that is, herring with gonads well-developed for spawning in the spring and herring which have only begun to recover from autumn spawning, the latter being dominant in the catches. The average size and age of these 2 spawning groups have gradually increased
over the past 5-6 years, and the fisheries have been maintained over the period by the abundance of pre-1960 year-classes ( 1958 year-class of autumn spawners and 1969 year-class of spring spawners). The scarcity of immature herring in the southwest Newfoundland catches suggests that annual recruitment to the stock occurs possibly during the summer in the Gulf of St. Lawrence. Age composition data indicate that recruitment has been relatively poor in recent years.

Between the autumn of 1969 and 1970 the stock size in southwest Newfoundland waters is estimated to have declined by about $40 \%$. The 1971 winter catch of herring in the same area is estimated to be about $50 \%$ of that recorded for 1970. Part of the decline in catch can be attributed to rather severe weather conditions during the past winter, but part is undoubtedly due to poor recruitment of young and to reduced abundance of older fish. Also the estimated exploitation rate of $40-50 \%$ is dangerously close to the level at which the Atlanto-Scandian stock began to collapse according to the report of the Atlanto-Scandian herring working group.

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10. Feeding and migrations of the roundnose grenadier, Macroumus rupestris ${ }^{1}$,
in the Northwest Atlantic and Iceland waters ${ }^{2}$
by S.G. Podrazhanskaya
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This report summarizes the results of a study of some aspects of the feeding and migrations of the roundnose grenadier, Macrourus mpestris, in the Northwest Atlantic and Iceland waters.

The material for the study was collected from 1967 to 1970 in the Iceland, West Greenland, Baffin Island, Labrador and the northern Newfoundland Bank areas.

Papers by Podrazhanskaya (1968, 1969) and Konstantinov and Podrazhanskaya (in press) give detailed analyses of the locality and diurnal rhythm of feeding, changes in the feeding habits with growth, and feeding habits at different depths as well as the food relations of the roundnose grenadier and other deepwater commercial fish species occurring within the same depth range, such as deep-water redfish and Greenland halibut. Therefore, this paper is limited to a general description of feeding and its seasonal dynamics. This will enable us to proceed to the pattern of feeding and spawning migrations of the roundnose grenadier.

The stomach contents of the roundnose grenadier show quite a number of forage species (52) represented mostly by planktonic and nectobenthic forms (Themisto, copepods, Euphausiidae, squid, fish). Only representatives of the order Polychaeta are associated with the bottom; shrimps ( $P$. borealis, genus Pantophilus) also live close to the bottom. Sand, mud and stones are sometimes observed in the stomachs of the roundnose grenadier, which suggests that they may go to the bottom in search of food.

The qualitative composition of forage organisms and the frequency of their occurrence vary considerably within the months of observations. However, the data do not show any consistent pattern, except that the total indices and the degree of fullness of the stomachs increase from September to December (Fig. 1).

The possibility of the feeding of the roundnose grenadier becoming more intensive from September to December is supported by the fatness dynamics (Fig. 1). In the Northwest Atlantic the fat content (percentage weight of liver to weight of whole fish) of the roundnose grenadier increases from August reaching a maximum in December. In the Icelandic area, the fat content follows a downward trend from June and reaches a minimum in August.

Material collected on the fatness of the roundnose grenadier in some areas within the Northwest Atlantic (northern Newfoundland Bank and Baffin Island) has also been analyzed. Fig. 2 shows that in the northern Newfoundland Bank area, the fat content increases with the size of fish. This is typical of all Gadidae.

In the Baffin Island area, the fat content is somewhat stabilized irrespective of the size of fish. In this area the fish seem to accumulate and store

[^8]a certain amount of fat.

It seems appropriate here to consider the question of the migration of the roundnose grenadier in some detail. Investigations conducted by PINRO in the North Atlantic over the past few years (1965-1970) have not revealed the spawning areas of the roundnose grenadier.

Zakharov and Mokanu (1970) suggested that the roundnose grenadier spawned in Icelandic waters. This seems quite likely and the material collected suggests that the eggs and larvae are carried passively by the Irminger Current from Iceland to the waters south of Greenland. From there the western branch of the West Greenland Current takes the young fish to Baffin Island and the waters of the Canadian and then the Labrador Currents take them to the northern Newfoundland Bank where they stay at depths of 600 to 900 m . On reaching a length of 40 to 50 cm , the roundnose grenadier start migrating back to their spawning grounds (Fig. 3).

Unfortunately, it does not seem possible to check and confirm the suggested pattern of migration by tagging experiments because the fish in trawl catches are always severely injured as a result of sharp drop in the pressure.

The suggested pattern of migration is based on the analysis of feeding, fatness, length-weight composition and sex ratio.

The fish, which have accumulated a certain amount of fat at Baffin Island, seem to start their pre-spawning migration to Icelandic waters. It is quite likely that here we have a situation similar to that for the Barents Sea haddock (Sonina, 1969). During the years when haddock feed on benthos, the duration of the feeding period and the time when the fish leave their feeding grounds usually depend on fatness. Haddock leave their feeding grounds having a fat content of not less than $4.5-5.0 \%$. The sooner they accumulate sufficient fat, the earlier they leave the feeding grounds, $i . e$, the cessation of feeding and the migration from the feeding grounds are related to the accumulation of a certain amount of fat. The area where the roundnose grenadier feed before they leave for the spawning grounds appears to be the Baffin Island area. This is supported by the data presented in Table 1.

Table 1. The mean index of the degree of fullness of the stomachs of Macrourus mpestris in different areas of the North Atlantic.

| Area | 1967 | 1969 | 1970 |
| :--- | :---: | :---: | ---: |
| Iceland | 0.8 | 0.8 | 1.3 |
| West Greenland | 0.9 | 1.4 | 1.5 |
| Baffin Island | 1.5 | 1.8 | 2.4 |
| Northern Labrador | 1.5 | 1.8 | 1.7 |
| Northern Newfoundland Bank | 1.3 | - | 0.6 |

The size composition of the roundnose grenadier caught in the areas surveyed is shown in Fig. 4. It shows that the smallest fish occur on the northern Newfoundland Bank where the modal length is $45-47 \mathrm{~cm}$. In the Baffin Island
area, the modal length is $60-62 \mathrm{~cm}$ and at West Greenland it is $78-80 \mathrm{~cm}$. The largest fish occur at Iceland where they are on the average 20 cm longer than in the above areas. These data are in agreement with those obtained earlier (Savvatimsky, 1969; Pechenik and Troyanovsky, 1970) and may be taken as confirmation of the suggested migration pattern if it is assumed that as they grow they move back to their spawning grounds. Many fish species, for example the Labrador cod, are known to perform migrations of this kind (Serebryakov, 1967). The Labrador cod spawn mainly along the slope off northern Labrador, the larvae and young fish in earlier developmental stages drift southward to the northeastern slope of the Grand Newfoundland Bank and as they grow they come back to the northern Newfoundland Bank and Labrador areas.

Roundnose grenadier caught at $145-900 \mathrm{~m}$ in the northern Newfoundland Bank, Northern Labrador, Baffin Island and West Greenland areas in July-August 1970 were immature even at their maximum length of $108-110 \mathrm{~cm}$. At southeast Iceland the population consisted of fish in the pre-spawning and spawning state in May and of spawning and spent fish in August (Table 2). Thus, it is quite likely that the waters of the Canadian and Greenland slopes are the feeding grounds for the roundnose grenadier and the areas from which the grown fish migrate to Iceland, the area where they were born. A similar migration pattern is followed by the Barents Sea herring which spawn mainly outside the Barents Sea along the northwest coast of Norway. From here the larvae and young fish are brought by the North Cape Current and are widely distributed within the Barents Sea (Marty, 1956).

Table 2. Maturity stages of Macrourus mpestris in Iceland waters.

| Month | Sex | No. of fish | No. fish at maturity stages |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | 1 | 2 | 3 | 4 | 5 | 6 | 6-2 |
| May | males | 38 | 3 | 28 | 3 | 2 | 2 | - | - |
|  | females | 17 | - | 14 | 1 | 2 | - | - | - |
| August | males | 54 | - | 39 | 5 | - | - | 7 | 3 |
|  | females | 106 | - | 59 | 11 | 1 | - | 1 | 34 |

It should be noted that the male-female ratio was found to be 2 to 1 in Canadian waters (Fig. 5) and 1 to 2 in Iceland waters.

The material available at present does not permit any definite conclusions but the females seem to mature and to arrive in the spawning area earlier than males.

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Fig. 1. Changes in the intensity of feeding and in the fat content of Macrourus rupestris in different areas of the North Atlantic in summer-winter 1967.

|  |  |  |
| :--- | :--- | :--- |
| Size groups: | 1. | $40-50 \mathrm{~cm}$ |
|  | Iceland |  |
|  | 3. | $51-60 \mathrm{~cm}$ |
|  | 3. | $61-70 \mathrm{~cm}$ |
|  |  | $71-80 \mathrm{~cm}$ |



Fig. 2. Changes with size in the fat content of Macrourus mapestris In different areas of the Northwest Atlantic (July-August 1970).

1. Northern Newfoundland Bank
2. Baffin Island


Fig. 3. Suggested spawning and feeding mbgrations of Macrourus mpestris of North Atlantic.
$\longrightarrow$ spawning migration
$\rightarrow-----\rightarrow$ feeding migration


Fig. 4. Size composition of Macroumu mpestris caught in different areas of the North Atlantic in the summer of 1970.


Fig. 5. Sex ratio of Macrourus mupestris in different areas of the
North Atlantic in the summer of 1970 .

## 11.

Studies of the age and growth of roundnose grenadier (Macrourus rupestris Gunn) ${ }^{1}$ in the North Atlantic, $1967-1970^{2}$
by P.I. Savvatimsky
PINRO, Murmansk

## Introduction

Roundnose grenadier, a valuable commercial fish, is widely distributed in the eastern and western Atlantic Ocean along the continental slopes at depths from 180 to more than $2,000 \mathrm{~m}$ (Leim and Scott, 1966). Other grenadiers are abundant in different areas of the Atlantic and Pacific Oceans (ICNAF, 1968; Konstantinov and Noskov, 1968; Makushok, 1967; Pechenik, 1965; Rass, 1959; Savvatimsky, 1969; Trunov, 1968; Yanulov, 1962). However, the mode of 11fe of roundnose grenadier as well as other species of grenadiers has been investigated very little due to their life in deep-sea areas, and only fragmentary data on their age and growth are available (Kulikova, 1957; Motais, 1960; Yanulov, 1962). In recent years the study of age and growth rate of roundnose grenadier in the North Atlantic has become very important due to the intensification of their fishery. The present paper is dedicated to this problem.

## Material and methods

Attempts to determine the age of grenadiers by otoliths were not very successful. Preparations were made by several methods: non-polished crosssection of the otolith; burnt non-polished cross-section; burnt polished crosssection; cross-cut $0.5-0.6 \mathrm{~mm}$ thick, polished on both sides and set in resin (colophony) on the glass. Of the methods mentioned, best results are obtained using the last two methods; yet they are the most labour-consuming. However, these methods and others do not give sufficient accuracy. An error of about 2 to 4 years can be made because the rings in the central portion of the otolith are faint. The closeness of growth zones and their poor contrast create additional difficulties.

To increase the contrast of the zones, a section 10 microns thick was cut using the microtome. The otolith was first decalcified by means of prolonged exposure in Bouin's solution. Then it was put into alcohols of increasing concentration. The otolith was then sealed with paraffin and stained with haematoxylin. It was impossible to count all the rings because in making the preparation the central portion of the otolith was destroyed. In this case, the rings were clearly marked.

Age studies of grenadier from their otoliths indicate that the fish have a long life because the annual rings are situated close to each other and are numerous.

[^9]The determination of grenadier age by scales when using ordinary methods is hampered by indistinctness of the annual rings; in addition, the scales have minute spines impeding their preparation for examination of the rings.

Staining of grenadier scales with different dyes did not produce more distinct annual rings.

Polarized light was used by Farran (1924), when determining age by scales. After being washed with a weak solution of alkall, the scales were examined by polarized transmitted light. Dark and light rings, that are parallel to the scale edges, were taken as annual rings. These rings were most clearly marked on the anterior spineless portion of the scales.

To confirm the fact that rings seen on the scales of grenadier by polarized light were annual rings, a comparative analysis of age determination of redfish and haddock was made by the regular method and by polarized light.

Farran (1924) who investigated the age of grenadiers of several species using scales and polarized light concluded that it was impossible to determine the exact age of each fish. In our experiments on several scales of the same fish, the number of rings seen by polarized light often proved to be different by 1-2, sometimes 3, rings. Comparatively, a small number of scales which differed from the others in number of rings was not taken into account. Scales were usually taken in the region just above the lateral line between the dorsal fins. Sometimes the scales could only be taken in the undisturbed areas of the body because scales are loosely attached in the scale pockets and during the trawl hauling great numbers of scales are lost. Dry scales become very fragile; therefore, one should protect the packets of collected scales from any physical harm. The edges of the scales are very thin and they can be damaged or broken when washed with a weak solution of alkali.

About $50 \%$ of scales in the collected samples are regenerated and no rings are found in the central portion. The remaining scales usually have the rings clearly marked along the edges, whereas they are most difficult to count in the central portion. All doubtful scales which are regenerated or all with illegible rings were discarded. Careful selection of scales for preparations allows us to determine the age of grenadiers with sufficient accuracy (Fig. 2).

Curves of linear and weight growth of grenadier were calculated according to the formulae $y=a x^{b}$, where $x$ is the age of fishes expressed in years, $y$ is the length expressed in cm or weight expressed in grams, and a and $b$ are the parameters found.

Age samples of grenadier were taken in 1967 and 1969 from bottom trawl catches in different areas of the North Atlantic (Fig. 1). The age of 420 fish was determined by scales (Tables 1,2 ).

## Results of investigations

The length of fish of the same age varied slightly in all the samples (Fig. 2).

The curves of linear growth were calculated according to the formulae: 0.59
$y=14.896 x \quad$ for males; and $y=17.2 x^{0.57}$ for females.

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Table 1. Growth in length (cm) and weight (g) of roundnose grenadier in the eastern North Atlantic in 1967.

Table 2. Growth in length (cm) and weight (g) of roundnose grenadier in the western North Atlantic in 1967 and 1969. (numerator - average length (cm); denominator - average weight (g))


The curves of weight growth were calculated according to the formulae

$$
y=8.77 x^{1.66} \text { for males and } y=13.8 x^{1.6} \text { for females (Fig. 3). }
$$

Females increase their wefght and length somewhat faster than males. The rate of linear and weight growth according to the calculated curves is shown in Fig. 4. Length increments decrease while weight increments increase with age. Fish of commercial size (at an age of $9-14$ years) have a length increment of about $3-4 \mathrm{~cm}$ a year, while weight increases in males by $60-80 \mathrm{~g}$ and in females by $80-100 \mathrm{~g}$. At 19 years of age, males reach a length of 84 cm and weight $1,010 \mathrm{~g}$, while females are 88 cm and $1,250 \mathrm{~g}$.

Despite the fact that samples were collected in different months and years in areas situated a considerable distance apart (Fig. 1, Tables 1, 2), it was established that males have similar length at the same age. This was also true for females.

In the age sample taken on the North Newfoundland Bank on 5-6 November 1969, there were 277 fish from 31 to 88 cm in length and 3 to 19 years of age (Table 2). On conversion of age data into the length composition of catches taken at the same time and in the same area (about 2,562 specimens), it was found out that the bulk of the catches consisted of males 9 to 12 years and females 6 to 10 years (Fig. 5).

Changes in length and sex composition of roundnose grenadier catches taken on North Newfoundland Bank In 1967-1970

In recent years the major fishery for roundnose grenadier has been conducted on a very limited area of the North Newfoundland Bank where the greatest quantity of this fish was caught (Table 3). At present only feeding concentrations of grenadier are utillzed. These are found at shallower depths than the adult fishes. From 1968 to 1969 in this area some changes in the length and sex composition of the catches took place. Fig. 6 shows the length composition of the catches taken in different years at approximately the same depths and during the same period of time (in order to exclude the effect of seasonal migrations of fish on the results of the analysis). The left part of the figure shows that from 1967 to 1969 the modal length of fishes decreased in the catches. In the right part of the figure, where annual deviations from the long-terin mean values are presented, one can see that during these years the relative quantity of large fish in the catches has dropped sharply. When analyzing the data for the whole period of the fishery (from June to December 1967-1970, when 40,690 fish were measured), analogous changes were revealed in the length composition of grenadier (Fig. 7).

The sexual composition of the catches has also changed. From 1967 to 1969 the relative quantity of females dropped from $39.3 \%$ down to $30.8 \%$. Females are larger than males, and apparently because of this they were subjected to fisheries to a greater extent than the males. Nikolsky (1958) believed that under rational exploitation of a stock, especially fishes with a long life cycle, in addition to the age and length composition of the fishes caught one should take into account the sexual composition required for perpetuating the stock, especially when there are differences in lengths of males and females and when a greater quantity of males or females are mainly exterminated by fisheries.

Table 3. USSR yield (in tons) of roundnose grenadier, 1967-1969.

| Year | Subarea |  |  | Total |
| :---: | ---: | ---: | ---: | ---: |
|  | 1 | 2 | - | Yleld |
| 1967 | - | 15,902 | 15,902 |  |
| 1968 | 116 | 2,553 | 24,143 | 26,812 |
| 1969 | 68 | 651 | 11,682 | 12,401 |

On the North Newfoundland Bank the catch per unit of fishing effort has also dropped.

The data presented here on age and growth rate of roundnose grenadier allow us to make the following important conclusion: this fish has a long life cycle and a many-aged population structure, and hence one should regard with caution any intensification of the Roundnose grenadier fishery.

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Fig. 1. Areas and time of collection of age samples of roundnose grenadier in North Atlantic.


Fig. 2. Length of roundnose grenadier of different ages on the North Newfoundland Bank, 5-6 November 1969 (1-males; 2 - females).






Length, cm



Fig. 7. Length composition of roundnose grenadier on the North Newfoundland Bank, June-December 1967-70.

# 12. Observations on selectivity and relative efficiency of salmon drift nets in connection with population studies on salmon at West Greenland ${ }^{1}$ 

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With the object of estimating the age and size distribution of the salmon stock in Greenland waters, investigations on the selectivity of drift nets were carried out in the autumn 1970 on some West Greenland fishing grounds.

The offshore salmon fishery at West Greenland is predominantly a drift net fishery. Salmon long-lines are operated too, but only on a small scale. Before 1969 Danish salmon vessels were exclusively equipped with multifilament nets manufactured from twine of twisted synthetic fibres (polyester). Besides using multifilament nets, two of the Danish vessels participating in the Greenland fishery in 1969 also experimented with monofilament nylon nets (polyamide). In the 1970 salmon fishing season, all the vessels, except one, were equipped with monofilament nets, either exclusively as far as two boats were concerned or as a supplement to the conventional multifilament nets.

The multifilament nets operated in Greenland waters are usually of the type which was introduced in the Baltic salmon drift net fishery during the first half of the sixties, Dimensions were generally 27 m long and $5-6 \mathrm{~m}$ deep; mesh size: 160 mm (stretched mesh); mounting: generally the net proper is mounted with strops to the float line. The net is hauled along the float line by hand. A somewhat modified type was fished from the vessel Polarlaks in 1969, as shall be referred to later on. Strops were inserted not in connection with the float rope, but between the net proper and the footrope which was lightly weighted.

Monofilament nets for salmon fishing are of various makes and qualities as well, with respect to colour, thickness, elasticity of the twine as to kind of knots. Dimensions were generally 34 m long and $5-6 \mathrm{~m}$ deep; mesh size: $130-160 \mathrm{~mm}$; mounting: the net proper is fixed directly to the float rope and connected to the footrope with strops. The gear is hauled along the footrope by a spectal winch.

The present population study is based upon observations on the selectivity of drift nets made on board the Danish commercial fishing vessel Polarlaks during salmon fishing off the coast of West Greenland in the autumn 1970. The report also includes some observations on drift netting from the same vessel the year before. Available data on salmon catches of other vessels have also been used.

When the investigation started in August 1970, Polarlaks was operating 609 monofilament nets distributed as follows:

[^10]| Mesh size | Numbers |
| :---: | :---: |
| 130 mm | 265 |
| 140 mm | 135 |
| 150 mm | 97 |
| 160 mm | 101 |

In addition, 25 nets with 170 -mm meshes were used experimentally during a part of the period of investigation. The nets were shot in chains of about 100 , and within each chain, nets with small and large meshes were tied alternately.

During two periods, 1-7 and $10-17$ September, while fishing in the areas about $69^{\circ} 45^{\prime} \mathrm{N}, 55^{\circ} 15^{\prime} \mathrm{W}$ and about $61^{\circ} 15^{\prime} \mathrm{N}, 49^{\circ} 00^{\prime} \mathrm{W}$ respectively, observations were made on catches by nets of different mesh sizes. The netted salmon were measured (fork length) and the mesh size of the corresponding nets was determined by a mesh gauge. Effort data were obtained by recording the number of nets of the various mesh sizes used and the fishing time of each chain of nets. However, as no correlation was observed between number of salmon retained and the time of fishing, the estimated catch-per-unit-effort values are only expressed as number of salmon retained per net unit.

Table 1 summarizes the observations made during fishing in the two areas concerned. The estimations are based upon measurements of 1,901 salmon. Catch-per-unit-effort data recorded from the fishing area $69^{\circ} 45^{\prime} \mathrm{N}, 55^{\circ} 15^{\prime} \mathrm{W}$ have been used in the calculations of selectivity for both areas, as corresponding data from the fishing area $61^{\circ} 15^{\circ} \mathrm{N}, 49^{\circ} 00^{\prime} \mathrm{W}$ are too scarce and, therefore, not sufficiently reliable. The results are presented graphically in Fig. 1. As the catches in nets of 170 mm were extremely poor, they are excluded from the diagrams. The same was the case with respect to nets of 160 mm in the southernmost area.

Table 1. Number of salmon in $5-\mathrm{cm}$ length-groups retained per 100 monofilament nets in relation to mesh size. Data from commercial catches at $69^{\circ} 45^{\prime} \mathrm{N}, 55^{\circ} 15^{\prime} \mathrm{W}, 1-7$ September, and at $61^{\circ} 15^{\prime} \mathrm{N}$, $49^{\circ} 00^{\prime} \mathrm{W}, 10-17$ September 1970.

| $\begin{gathered} \hline 5-\mathrm{cm} \text { length-groups } \\ \text { (fork length) } \\ (\mathrm{cm}) \\ \hline \end{gathered}$ | $\begin{gathered} 69^{\circ} 45^{\prime} \mathrm{N}, 55^{\circ} 15^{\prime} \mathrm{W} \\ \text { Mesh size }(\mathrm{mm}) \end{gathered}$ |  |  |  |  | $\begin{aligned} & 61^{\circ} 15^{\prime} \mathrm{N}, 49^{\circ} 00^{\prime} \mathrm{W} \\ & \text { Mesh size (mm) } \end{aligned}$ |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 95-100 | 0.1 | - | - | - | - |  |  |  |
| $90-95$ | - | - | - | - | - | - |  |  |
| $85-90$ | 0.1 | 0.3 | 0.4 | 0.6 | - | 0.1 | - | - |
| $80-85$ | 0.4 | 1.4 | 1.1 | 2.5 | - | 0.2 | 0.5 | 0.5 |
| $75-80$ | 1.3 | - | 0.9 | 1.5 | 1.3 | 0.2 | - | - |
| $70-75$ | 4.3 | 2.4 | 2.8 | 3.4 | - | 2.4 | 2.5 | 4.8 |
| 65-70 | 19.7 | 18.6 | 9.6 | 4.5 | 1.3 | 20.4 | 18.2 | 11.5 |
| $60-65$ | 24.4 | 14.9 | 5.8 | 2.1 | 2.5 | 26.6 | 14.9 | 4.3 |
| $55-60$ | 10.2 | 3.6 | 1.3 | 0.7 | - | 10.1 | 5.4 | 0.9 |
| $50-55$ | 1.2 | 0.3 | 0.2 | - | - | 1.6 | - | - |
| 45-50 | 0.1 | - | - | - | - | - | - | - |
| Total number of salmon per 100 nets | 61.7 | 41.5 | 22.0 | 15.3 | 5.1 | 61.7 | 41.5 | 22.0 |
| Number of salmon | 765 | 257 | 103 | 81 | 4 | 492 | 153 | 46 |

For the sake of comparing the relative efficiency of nets fished from Polarlaks with nets of corresponding mesh sizes operated from other vessels, all available information on catch-per-unit effort of a range of mesh sizes is shown in Table 2. The data are listed according to vessels, date and fishing area.

Table 2. Number of salmon retained per 100 monofilament nets in relation to mesh size. Information from four commercial vessels at West Greenland, August-October 1970.

| $\begin{gathered} \text { Vessel } \\ \text { No. } \end{gathered}$ | Date |  | $\begin{aligned} & \text { ICNAF } \\ & \text { Div. } \end{aligned}$ | Number of salmon per 100 nets of Mesh size (mm) |  |  |  | Basis of calculations (No. of nets) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | 130 | 140 | 150 | 160 |  |
| II | 30-31 | August |  | 1B | 76.0 | 57.8 | 18.8 | - | 1,000 |
| I | 1-7 | September | 1A | 61.7 | 41.5 | 22.0 | 15.3 | 2,720 |
| III | 6 | September | 1B | 74.0 | - | 32.0 | 18.0 | 300 |
| III | 5-7 | September | 1B | 64.0 | - | 29.0 | - | 600 |
| II | 1-7 | September | 1A | 94.6 | 72.7 | 24.3 | - | 3,100 |
| II | 11-21 | September | 1E | 96.3 | 86.3 |  | - | 1,300 |
| IV |  | SeptemberOctober | 1E | - | 45.6 | 23.4 | - | 5,820 |

An attempt has been made to estimate the length frequency of the salmon population in Greenland waters. For this purpose a formal model of gill net mesh selection has been used which implies the following assumptions: 1) Mean and standard deviation of the length frequency distribution of the salmon retained is proportional to mesh size. 2) Fishing power of nets is independent of mesh size. The application of the model to the material did not, however, lead to results on which an estimation of a probable length frequency distribution of the salmon population could be based. The reason may be either that the assumptions made are not valid or that the observations were too few.

Considering the remarkable decline in catch efficiency from the smallest to the largest meshes, it cannot be rejected that, on account of certain properties attached to the nets, the fishing power of the various nets may depend on mesh size. One such property may be the elasticity of the monofilament twine. According to tests of tension and breaking strength the twine of the nets in question can be stretched about $20 \%$ before breaking. Consequently, the effective mesh size within certain limits depends on the strain applied. Supposing that the strength of a salmon is proportional to the size of the individual, it might be expected that the number of escaping big salmon entering large meshes will be relatively greater than the corresponding loss of small salmon in nets of small meshes.

Since the study of net selection failed to give any results, the observations only provide rather uncertain information on the age and size composition of the stock of salmon in West Greenland waters. Some conclusions are, however, drawn as shall be mentioned in the following.

In spite of the significantly different selectivity of the nets operated, average length of salmon retained by nets of different mesh sizes varies
relatively little as shown by following means:

| Mesh size: | 130 mm | 140 mm | 150 mm | 160 mm |
| :--- | ---: | ---: | ---: | ---: |
| $69^{\circ} 45^{\prime} \mathrm{N}, 55^{\circ} 15^{\prime} \mathrm{W}$ | 64.8 cm | 66.1 cm | 68.1 cm | 72.2 cm |
| $61^{\circ} 15^{\prime} \mathrm{N}, 49^{\circ} 00^{\prime} \mathrm{W}$ | 63.7 cm | 65.1 cm | 67.2 cm | - |

Unless the properties attached to monofilament nets as mentioned above reduce their fishing power considerably with increasing mesh size, the figures seem to indicate a relatively small proportion of $A .2+$ salmon in the population at West Greenland.

Length distribution of the catches made in the two areas where the majority of the fishery took place during the first half of September, revealed a certain variation of stock composition at West Greenland, as shown in Table 1 and Fig. 1. The proportion of salmon of 75 cm and above - roughly speaking age-group A. $2+$ in nets of different mesh size was as follows:

| Mesh size: | 130 mm | 140 mm | 150 mm |
| :--- | :--- | :--- | ---: |
| $69^{\circ} 45^{\prime} \mathrm{N}, 55^{\circ} 15^{\prime} \mathrm{W}$ | $3.0 \%$ | $4.3 \%$ | $10.8 \%$ |
| $61^{\circ} 15^{\prime} \mathrm{N}, 49^{\circ} 00^{\prime} \mathrm{W}$ | $1.0 \%$ | $1.3 \%$ | $2.2 \%$ |

Differences in condition of salmon taken in the two areas could also be demonstrated. Based on length and weight measurements of a representative sample of 49 and 99 salmon caught at $69^{\circ} 45^{\circ} \mathrm{N}, 55^{\circ} 15^{\prime} \mathrm{W}$ and $61^{\circ} 15^{\prime} \mathrm{N}, 49^{\circ} 00^{\prime} \mathrm{W}$, the condition factor of the two populations was estimated to be 1.02 and 1.07 respectively (fork length and round weight). The difference is statistically significant ( $0.05>P>0.025$ ).

As the ability of gill nets to retain fish of a given size depends on the condition of the fish, differences in length composition of corresponding age-groups of salmon caught in the two areas might be expected. Such differences could not be found, as seen in Fig. 1.

In default of estimates of stock composition and gear selectivity, relative exploitation rates of the individual length- and age-groups cannot be evaluated. It may, however, be pointed out that the highest yield of a lengthgroup per unit of gear was obtained of individuals between 60 cm and 65 cm by nets with the smallest mesh size used, $i . e ., 130$ mm. The mean length of the population of A.lt salmon, therefore, is supposed to be less than 65 cm at the time of investigation, which suggests that the smaller individuals of this agegroup are exploited less than the bigger individuals even by nets of $130-\mathrm{mm}$ meshes. It cannot be excluded, that nets with still smaller meshes might have taken even more salmon per net unit.

Differences with respect to selectivity and catching efficiency of multifilament nets of 160 -mm meshes in relation to monofilament nets of various mesh sizes are shown in Tables 3 and 4. Unfortunately, the material does not allow an exact separation of the catches by monofilament nets with respect to mesh size. Referring to Table 4, monofilament nets are put together in two groups, $130-\mathrm{mm}+140-\mathrm{mm}$ nets and $150-\mathrm{mm}+160-\mathrm{mm}$ nets. In the former group,

Table 3. Number of salmon in $5-\mathrm{cm}$ length-groups retained per 100 nets In relation to type of twine and mesh size of nets. Data from commercial fishing within the area $67^{\circ} 00^{\prime} \mathrm{N}-67^{\circ} 37^{\prime} \mathrm{N}$ and $54^{\circ} 15^{\prime} \mathrm{W}-55^{\circ} 00^{\prime} \mathrm{W}, 18$ October-2 November 1969.

| 5-cm length-groups <br> (fork length) <br> (cm) | Number of <br> multifilament nets <br> 160 mm | Number of <br> monof ilament nets <br> $130 \mathrm{~mm}+140 \mathrm{~mm}$ |
| :--- | :---: | :---: |
| $95-100$ | - | 0.05 |
| $90-95$ | 0.2 | 0.1 |
| $85-90$ | 0.3 | 0.5 |
| $80-85$ | 1.4 | 1.8 |
| $75-80$ | 2.3 | 4.1 |
| $70-75$ | 11.8 | 15.8 |
| $65-70$ | 11.8 | 16.8 |
| $60-65$ | 0.3 | 6.0 |
| $55-60$ | - | 1.0 |
| $50-55$ | 32.7 | 0.05 |
| Total number of salmon |  | 46.3 |
| per 100 nets | 210 | 1,131 |
| Number of salmon |  |  |
| measured |  |  |

Table 4. Number of salmon retained per 100 nets in relation to type of twine of the nets. Information from two commercial vessels at West Greenland, October-November 1969, and August-November 1970.

the two mesh sizes are represented by $65 \%$ and $25 \%$ respectively, while the remaining $10 \%$ consists of nets of 150 mm and 160 mm . The latter group of nets is exclusively represented by $150-m$ and $160-m m$ nets in the ratio $2: 1$. On account of differences in length of the two net types in question, values of catch per 100 multifilament nets are converted in order to make them comparable to catches of monofilament nets.

It appears from the tables that multifilament nets of 160 mm are superior to monofilament nets of, roughly speaking, the same mesh size in catching salmon, but are outnumbered by monofilament nets or 130 mm and 140 mm .

A comparison between the length distribution of catches by 160 -mm multifilament and 130- $+140-m$ monofilament nets (Fig. 2) discloses an amazing uniformity in spite of the differences of mesh size. This probably implies that nets of same mesh size but made of different sorts of twine, monofilament and multifilament, are not only different as to catching efficiency but also with respect to selectivity. Apparently, the two kinds of net do not possess a uniform catching ability; this may be explained in different ways, a.o. (1) Multifilament nets are, no doubt, more visible in the sea than monofilament nets. (2) Multifilament nets - either entirely lacking or only with a light footrope - are not so hard vertically stretched as the heavily weighted monofilament nets. The latter are gill nets in the proper sense of the word, whereas, multifilament nets are retaining salmon more by entangling.


Fig. 2. Number of salmon in 5-cm length-groups retained per 100 nets during fishing within
the area $67^{\circ} 00^{\prime} \mathrm{N}-67^{\circ} 37^{\prime} \mathrm{N}$ and $54^{\circ} 15^{\prime} \mathrm{W}-55^{\circ} 00^{\prime} \mathrm{W}$, 18 October-2 November 1969.
13. Recent scallop recruitment and apparent reduction in cull size by the Canadian fleet on Georges Bank ${ }^{1}$
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## Introduction

Trends in the Georges Bank fishery since the high abundance years of 1961-62 have reflected a continuing decline in scallop stocks. This lower abundance of scallops and consequent increase in landed price has led to changes in fishing practice by certain sections of the Canadian industry and, following a small recruitment to the Northern Edge of the Bank in the fall of 1969, smaller scallops were being exploited in 1970 than during years of good to average abundance.

Population densities of scallops on the Northern Edge were assessed in June 1970 by experimental fishing and bottom photography, and the sizes of landed meats in commercial catches surveyed at three main ports in July-August 1970. These surveys considered together showed that small scallops being landed by the fleet were probably caught within a limited area of new recruits on the Northern Edge.

## Status of the fishery

Combined Canadian and U.S. landings from Georges Bank reached a peak in 1962 at 15.6 million kg of meats ( 130,000 metric tons whole weight) following the abundant recruitment of 1959 (Bourne, 1964) and have since declined by $64.5 \%$ to the 1970 figure of 5.5 million kg ( 46,000 metric tons whole weight) (Table 1). This decline in overall landings has not been so evident from Canadian landed figures, since Canada's share in the overall landings has increased from $36 \%$ to $73 \%$ of the total between 1962 and 1970 as U.S. effort on the grounds has fallen off. Canadian landings continued to go up until 1964, and since then have only declined by $32.6 \%$. However, in order to maintain landings at the present level of 4 million kg of meats ( 34,000 metric tons whole weight), the Canadian fleet has had to extend its operations from the productive Northern Edge and Peak to less productive areas well within the 50-fathom contour, and to deep water below 50 fathoms on the northeastern edge of the Bank. Canadian effort both in days fished per year and in hours dragged per day has increased by $20-40 \%$ since 1962, the peak year for landings (Caddy and Lord, in press).

Because an increasing proportion of the day has been spent dragging, catch per day has underestimated the decline in abundance since the early $1960^{\prime}$ s. Thus, while catch per day dropped by $44 \%$ between 1961 and 1969 , catch per hour (as determined by log book analysis) fell by $80 \%$ over the same period (Fig. 1).

[^11]Table 1. Landings of scallop meats (millions of kg ) from Georges Bank, 1957 to 1970.

|  | 1957 | 1958 | 1959 | 1960 | 1961 | 1962 | 1963 | 1964 | 1965 | 1966 | 1967 | 1968 | 1969 | 1970 |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Total <br> (USA and <br> Canada) | 8.6 | 7.7 | 10.4 | 13.3 | 15.3 | 15.6 | 13.9 | 12.1 | 6.0 | 5.9 | 6.4 | 6.0 | 5.8 | 5.5 |
| Canada | 0.8 | 1.2 | 2.0 | 3.4 | 4.6 | 5.7 | 5.9 | 6.0 | 4.5 | 4.9 | 5.0 | 4.8 | 4.4 | 4.0 |

## Recent recruitment to the fishery

Late in 1969, reports were received of appreciable numbers of young scallops on Georges Bank for the first time since 1959-60. Positions given by fishermen suggested that the area of recent recruits was confined to the Northern Edge of the Bank between $66^{\circ} 35^{\prime}$ and $67^{\circ} 15^{\prime} \mathrm{W}$ long. This area was surveyed in June 1970 and a population consisting predominantly of recent recruits was estimated to occupy approximately 80 square miles ( 274 sq km ) of the Northern Edge of the Bank between $66^{\circ} 45^{\prime}$ and $67^{\circ} 12^{\prime} \mathrm{W}$, in 25-35 fathoms ( $50-70 \mathrm{~m}$ ) (Fig. 2). Bottom photographs within the area showed population densities averaging $0.98 / \mathrm{sq} \mathrm{m}$, corresponding to a population of about 270 million scallops within the area of the concentration.

Calibration of the underwater camera allowed scallops in the photographs to be separated into three size groups: $<50 \mathrm{~mm}, 50-100 \mathrm{~mm}$, and > 100 mm (Table 2).

Table 2. Numbers and sizes of scallops in bottom photographs and dredge tows on the Northern Edge of Georges Bank (June 1970).

| Size group | < 50 mm | $\begin{gathered} 50-100 \mathrm{~mm} \\ \% \end{gathered}$ | $\begin{gathered} >100 \mathrm{~mm} \\ \% \end{gathered}$ | Total No. |
| :---: | :---: | :---: | :---: | :---: |
| Dredge contents | 6.5 | 71.2 | 22.3 | 18,580 |
| Bottom photos | 25.0 | 65.7 | 9.3 | 2,830 |

From bottom photographs, it was estimated that only $10 \%$ of the population exceeded the commercial cull size noted in 1960-62 of 95-100 mon (Bourne, 1964); $25 \%$ of the population were smaller than 50 mm . In comparison, $22.3 \%$ of the dredge catches (using 3 -inch internal diameter commercial rings) were larger than 100 mm , and only $6.5 \%$ smaller than 50 mm . Although some selection for larger scallops was being exerted by the gear, most of the fishable population in this area was smaller than 100 mm , and the largest catches contained the smallest scallops (Fig. 3). Three-ring scallops, with a modal size of 72.5 mm , were the dominant year-class in dredge hauls from within the area of high concentration. A small proportion of 2-ring scallops were also caught, although the actual abundance of this year-class was underestimated as judged from the bottom photos.

Meat sizes landed by the commercial fleet
Information on landed meat sizes was obtained for a single trip by 35 boats, representing $70 \%$ of the 50 Canadian vessels making regular trips to Georges early in 1970.

From each vessel, two $36-1 b(16.3 \mathrm{~kg})$ bags of scallop meats were picked at random from each third of the catch, and a $1-1 \mathrm{~b}$ ( 0.45 kg ) sample of meats weighed from each bag and counted. A 1-lb meat sample from each boat was frozen for later individual weighings of meats.

The mean numbers of meats per pound landed at three principal ports of landing for the offshore fleet were 43,49 , and 26 for ports $A, B$, and $C$ respectively, although counts of up to 79 meats per pound were recorded at port $B$. A highly significant difference in the mean numbers of meats/lb was found between ports $A$ and $C$, and $B$ and $C$ ( $<0.001$ ). The total weight of landings was significantly correlated ( $p<0.01$ ) with the mean number of meats per pound in the catch, confirming research findings that largest catches were made in the area of recent recruits, landings per trip at ports $B, A$ and $C$ diminishing in that sequence (Fig. 4).

Meats per pound were converted to mean weights per sample (gm) and since no data are available for the shell sizes shucked by the fleet, estimates of mean shell lengths per sample were obtained using length-weight data published by Haynes (1966, Table 2) for Georges Bank scallops in the period April to September.

Mean she 11 lengths were plotted as histograms for each port (Fig. 5) and compared with mean shell sizes observed in unculled research tows on the Northern Edge in June 1970. Some culling for size seems to have taken place, as judged by comparison of the ascending side of the distributions from the three ports with that for the unculled research catch. However, since approximately half of the calculated mean shell lengths are below 85 mm for ports $A$ and $B$, cull sizes for these two ports must be smaller than 85 mm , and probably fall between 70 and 85 mm . Half of the mean lengths for port $C$ are below 100 mm , so that cull size for this port has probably remained close to the 95-100 mm cull size recorded for the Canadian fleet in the early $1960^{\prime} \mathrm{s}$.

Examination of $\log$ records from ports $A$ and $B$ for the first half of 1970 revealed a considerable expenditure of effort within the area of recent recruits, while boats from port $C$ appear to have largely avoided this area (Caddy and Sreedharan, 1971).

## Summary

Evidence from port surveys and research catches suggests that following a period of poor recruitment and an $80 \%$ drop in scallop abundance on the Bank, a limited recruitment of 3 -ring scallops to the fishable population occurred on the Northern Edge of Georges Bank in 1969-70. Although the area of the concentration was small (approximately 270 sq km ), the standing stock within it was estimated by bottom photography to be about 270 million scallops in June 1970. The mean length of scallops in the photographs was calculated to fall within the range 65-70 mm. From data in Haynes (1966), this corresponds to an average meat weight of 5 gm , and hence to a standing stock of approximately
1.4 million kg of meats in the area of the concentration. This is approximately one third of the Canadian catch from the Bank in 1969 ( 4.4 million kg meats). This recruitment was being exploited by vessels from two Canadian ports in 1970, and for these two ports cull size had fallen well below that being observed in the early 1960's. Vessels from a third major port were fishing elsewhere on the Bank and landing smaller catches of predominantly large scallops.

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Fig. 1. Decline in two measures of catch/unit effort (Canadian fleet) on Georges Bank since 1961 ( $1-1 \mathrm{~b}$ scallop meats $=0.454 \mathrm{~kg}$ ).

Fig. 2. Area of recent recruitment of scallops to the Northern Edge of Georges Bank discovered


Fig. 3. Relationship between shell length and whole weight ( $1 \mathrm{lb}=$ 0.454 kg ) of scallops per tow made in June 1970 with a $2.4-\mathrm{m}$ wide drag.


Fig. 4. Relationship between mean number of meats/lb in the catch, and weight of meats landed per trip at three Canadian ports by 34 vessels fishing Georges Bank in the period July-August 1970.


Fig. 5. Mean lengths of scallops caught on Georges Bank in the summer of 1970, based on measurements of shell size in the research catch, and conversions of mean numbers of meats per $1 b$ to estimates of mean shell lengths, using data from Haynes (1966).

SECTION H SEALS

# 14. Calculation of production of Harp seals in the Western North Atlantic ${ }^{1}$ 

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## 1. Introduction

Experience has shown that some differences in the biological features of harp seals and fish need to be stressed to fisheries biologists. First, Pagophilus groenlandicus Erxleben is a long-lived mammal in which mortality rates are low and rather constant. Consequently a large number of age-classes of approximately similar strength contribute to annual production, which therefore fluctuates little. The population dynamics of the species are in fact more like those of human beings than fish. Secondly, the yield is taken as newborn or a few weeks old young before virtually any mortality has taken place, for mortality in the first week or so has been measured at 0.01 or less (unpublished data of the author). Therefore, yield to the fishery is a simple fraction of production. Third, the species is aggregated at more or less constant density when the young are fished, regardless of the size of the aggregation or whelping group. (This density is ideally set by an individual distance which exists between adult females with pups, i.e. they are over-dispersed, but the density is further reduced by the inaccessibility to seals of the centres of large ice floes.) Therefore, catch per unit of effort will not change with increasing catch of young until this catch is well above sustainable yield, and such data will not be of much practical value.

Four general methods developed for estimation of production will be discussed here. They are: photographic aerial survey, capture-recapture tagging, study of escapement of year-classes after large and small catches of young, and study of total catch of young when there are grounds, from the first three methods, for believing that the catches of young are a high proportion of production.

Definitions used are as follows: 0-group seals are termed juveniles, approximately 1-5 year old harp seals are sexually immature, and adults are aged some 6 years and up. The Front ice is off the Labrador coast in Subareas 2 and 3, the Gulf of St. Lawrence ice is in Subarea 4.

## 2. Reproductive Rates

The maximum sustainable yield of young has been calculated (Sergeant, MS, 1969) at between 0.24 and 0.38 of production for observed extremes of reproductive rate. The reproductive rate recently measured in the Gulf (Table 1) would favour the lower yield but may not apply to the Front population, where it has not been possible recently to measure the reproductive rate accurately. In 1962 the reproductive rate at the Front was optimal (Sergeant, 1966), as it was in the White Sea under heavy exploitation of both young and adult seals (Yakovenko and Nazarenko, 1967).

[^12]The suboptimal reproductive rate in recent years in the Gulf can be associated with an observed interaction between adults and young immatures at winter feeding grounds (Sergeant, MS, 1969). Segregation occurred and the adults appeared to be using the better feeding areas. Competition for food could affect rates of growth and reproduction. If the same lowered reproductive rate now exists at the Front, as seems likely from small data, then doubt is thrown on the assumption that the optimal yield is obtained from fishing the young animals only. However, it is difficult to reconcile the thesis, developed below, that total population has declined with postulated competition between age-classes. The question has not yet been resolved.

## 3. Aerial Surveys

The main apparent advantage of aerial surveys is that they give the only direct count or estimate, either of production, when applied to young seals or the adult females producing them, or of standing crop when applied to moulting groups of adults and immatures (Dorofeev and Freiman, 1928, and subsequent Soviet authors).

The main disadvantages are high costs of surveys and the need for completeness of each survey. Particularly, surveys of moulting groups must be simultaneous, since the groups may change position daily, and must be repeated to demonstrate patterns of aggregation. 'Ground' control is necessary to measure age and sex composition. Pilots must know how to find seals and at the same time the photography must be competent. No repeat is usually possible in the same year.

Surveys of whelping groups show that many young are invisible and few controls are possible. One control was achieved in Canada in 1967 when the industry decimated a group of seals just previously surveyed and detailed catch figures were available. Twenty thousand whelping adults had been estimated by aerial survey on 10 March. Thirty-five thous and young were taken on 13 and 14 March. Clearly, many adult females were in the water, either because they had not yet pupped, or had left the pup between episodes of suckling. This correction ( 0.57 of adults present in a group) might be applied to all counts made at the same stage of the reproductive cycle, but other variables, e.g. whether the seals are whelping on open pack ice with many leads, or on frozen-together shore ice affect their ability to enter the water. The sex ratio of whelping adults also is not accurately known, since on disturbance the males, present in some numbers ahead of mating time, enter the water more freely than the females attached to pups. Presumably, unwhelped females also enter the water readily. All these corrections might be entered into surveys, given repeated facilities of fixed-wing aircraft and helicopters, but total costs would be high.

Aerial survey was made of whelping groups in the Gulf and on the Front in 1970. The search was highly successful in that all whelping groups of seals were believed located, except that one Gulf group was located by another Government charter plane but not by the photographic plane, and its size was not precisely estimated.

However, the photography was poor. A $70-\mathrm{mm}$ camera, used for
the first time in place of the usual $9 \times 9^{\prime \prime}$ aerographic camera, proved to have film lengths too short for continuous overlapping coverage of the long axis of most seal patches. It was therefore necessary to load the film magazine in mid-survey and the long lines could not be matched afterwards when the mosaics of prints were mounted.

Results are shown in Table 2 with estimated major uncertainties in accuracy. Catches in a fourth Gulf group which was not surveyed were 10,000 seals, bringing the Gulf estimate to at least 75,000 whelping adults or young seals. Previous successful aerial surveys (Sergeant, MS, 1969) gave estimates or indices of 100,000 young seals produced in the Gulf in 1964, and 215,000 produced on the Front in 1960.

The main value of the 1970 survey was that the relative sizes of sets of whelping groups on the Front on the one hand, and in the southern Gulf of St. Lawrence on the other, were compared visually within a short space of time. They were approximately 5 to 3 or 2 to 1 , the Front having the larger set of groups. This estimate is confirmed from the counts (Table 2) and is important in relation to the unsolved problem of mixing of Front and Gulf groups, as follows:

In 1969, lack of ice in the southern Gulf had forced many Gulf animals to whelp on the Front (Section 8). By 1970, the usual numbers of animals had returned to the Gulf. Moreover, the Front has for many years taken a heavier fishery than the Gulf, a trend which has been continued since 1965 when the catch of young in the Gulf was regulated. In spite of this long-continued difference in intensity of catch, the Front whelping groups in 1970 were larger. Therefore, either the Front has not been heavily exploited, which from the evidence presented below is unlikely, or else the population of seals re-sorts itself annually between Gulf and Front.

## 4. Capture-recapture Tagging

This method has been made possible mainly by a latening of the starting dates for hunting young seals from 7 March to 12 March or later. Tagging between 2000 and 3500 young seals has usually been possible from an icebreaker or helicopter. The first figure represents, it is calculated, about $2 \%$ of Gulf production, the second, $2 \%$ of Front production. The interval between tagging and start of the fishery for young seals has varied from no time to up to 2 weeks. The longer the interval, the greater the chance of randomisation of the tagged seals through ice movement. Mortality of young at or immediately after birth, as already stated, is low and not likely much increased in the first few weeks (Popov, 1966). Greatest errors come from a concentration of tagging and recoveries in the same group of seals when other groups are not found or are inaccessible to either tagging or recapture effort. I have determined also that considerable variation occurs in catch returns of tagged seals from ship to ship within one tagged group, apparently because of failure to achieve randomisation of tagging. With such errors, sophisticated treatment seems as unprofitable as for aerial survey, and the best validation has come from three successive experiments in the Gulf, carried out in 1964, 1968 and 1970, which have given comparable estimates (Table 3). One experiment has been performed on the Front. Resulting estimates of Gulf and Front
production, when compared, are in line with the $5 / 3$ ratio between Front and Gulf production observed in 1970, and absolute figures are also broadly comparable when allowance is made for the incompleteness of aerial counts.

## 5. Study of Escapement After Varying Catch

Use of this method started from the observation from a number of age samples (one Danish, several Canadian) collected and analysed in 1953, that after an unusually heavy catch of young seals in 1951, the size of this year-class was consistently reduced. The same result was found after the next unusually heavy catch of young in 1956 and has been observed repeatedly thereafter, particularly following heavy kills of young in 1962 and 1963 (Figs. 1 and 2). The results should be particularly clear since 1961, when closing dates lowered the kill of age-groups older than the young ( 0 -group), making the kill of older age-groups steady at about 35,000 annually as opposed to variable and up to 100,000 prior to 1961. Previously it was possible for a year-class of young, which had received light exploitation and consequently had survived well, to be exploited heavily at a later age, especially as the somewhat segregated 1-year-old moulting animals.

The problem is to quantify the strength of each year-class as a measure of the degree of its survival.

## a. Using all age samples

A first, semi-quantitative method has been to sum the results of all age samples, allowing for various sorts of bias in these age samples which are known from long sampling to be steady, and the biological reasons for which are known. Thus, southward migrant seals lack younger age-classes, the number of immatures progressively increasing with increasing age (Fig. 1). In a real life table, they should of course decrease due to mortality. It is however known, from tagging results and from collection of age samples in West Greenland in winter, that the younger age-classes remain north in the winter up to 2 or 3 months longer than the main groups (Sergeant, 1965). The opposite bias occurs with samples from moulting groups at the icefields in April (Fig. 2, left). One and two year-olds are then over-represented. The reasons may be threefold: concentration and segregation of such immatures in the southern edges of the icefields most easily reached by ships, tameness of the young year-classes, and a movement (demonstrated by tagging) of young year-classes of Gulf seals to the Front at $1-3$ years of age.

Few samples, chosen eclectically, have been "ideal", that is, corresponding to the expected age frequency distribution. Best were a West Greenland sample collected from many points by Danish scientists in 1953, and a unique sample of moulting seals collected in the northern Gulf in 1966. These samples were used to estimate mortality rates. No apology is made for the slight circularity in this argument, since much other unquoted evidence on the expected mortality rates of seals was used in regarding these samples as ideal. An ideal distribution, for example, reasonably gives higher mortality rates to the inexperienced immatures than to the adults. Table 4 shows sources of age samples used and their probable biases. Detailed age frequencies are shown in Appendix Table 1. Table 5 shows the result of combining all samples in
order to estimate the escapement or index of survival of each year-class from 1960 to 1969. Clearly there is more accumulated evidence for earlier year-classes. The index of survival is expressed in four categories: very good rates as 3 , good as 2 , poor as 1 and very poor as 0 . These values are summed and then expressed as a proportion of the maximum possible score (always 3). By this means a year-class which always showed an index of 'very good' would score 1.00, one which always showed an index of 'very poor' would score 0.00 . Results of the analysis are plotted in Figure 3. From the least squares fit, extrapolation down to survival index 0.00 would occur at an estimated catch of 365,000 young seals, which therefore represents an estimate of present production. Survival to index 1.00 would occur at a catch of 115,000 young harp seals. The precise biological significance of this figure is obscure but since it measures "consistently good escapement of a particular year-class in all age samples, as compared with other year-classes," it may represent approximately the present sustainable yield. The ratio $115 / 365=0.32$ whereas the sustainable yield of young seals has been calculated from known mortality and fecundity rates at between 0.24 and 0.38 (mean, 0.31 ). Presumably, below catches of 115,000 natural mortality increases in a density-dependent manner, while above this value, an irreducible minimal mortality of juveniles occurs. This mortality is believed, by extrapolation back from mortality rates of immatures, to be about 0.20 on an annual basis (Sergeant and Fisher, MS, 1960).

## b. Using single sets of age samples

This method is the same as above but confined to one set of age samples at a time, so that variations in selectivity or bias between sets of samples is eliminated. Within one set of samples, bias is seen to be much the same each year (Fig. 2), so that by expressing catch and escapement as proportions between one year and the next, this bias can be eliminated. Thus true quantitative results can be achieved. For proper formulation of the method I am much indebted to Dr. W. E. Ricker.

Consider for example the two sets of samples seen in Fig. 2. One is of seals sampled by net from a shore fishery at or near St. Anthony, northeast Newfoundland, in January to March 1967, 1968, 1969 and 1970. The second set comes from the ship fishery a few miles off this coast and two months later, in April. The animals are then shedding their hair on their annual moult and are shot. In the second set, a bias towards one and two year-old animals is seen as compared with the first set.

Let us consider the shore fishery only. In the first two samples, year-classes 1966 and 1967 show up poorly as one year-old animals but, in the third sample, collected in 1969, year-class 1968 appears in great strength. Now the catch of young seals in 1968 was reduced to half the catch in 1967 or 1966, as the result of a late starting date and temporary reductions in catching effort. The association between low catch and enhanced escapement is seen also in the moulting samples.

We will assume that in successive adjacent years neither stock nor production have changed by an appreciable amount, and that natural mortality during age 0 is constant over this period (i.e. that not even in 1968 was maximal sustainable yield taken).

Let $S$ = stock excluding young (same in both years)
$B=$ number of young just before harvesting (same in both years)
$\mathrm{C}=$ catch (known)
$\mathrm{v}=$ natural mortality rate after kill and before sample taken at age 1 (same in both years)
$K=$ ratio of age 1 seals to $S$ (known for each year).
We wish to estimate B. From the above we can write:
Survivors after catch $=$ B - C
Survivors at age $1=(B-C)(1-v)$
Age 1 divided by stock $=\frac{(B-C)(1-v)}{S}=K$

$$
\frac{K_{1}}{K_{2}}=\frac{B-C_{1}}{B-C_{2}} \quad B=\frac{K_{1} C_{2}-K_{2} C_{1}}{K_{1}-K_{2}}
$$

Table 6 shows the data and Table 7 the calculations using St. Anthony samples on the left side and icefield samples on the right, the mean of 1966 and 1967 catches and escapements compared with 1968, above, and 1968 compared with 1969, below.

For the 1966-1967/1968 comparison, from both areas, results are in good agreement. Using the Front catch figures, production is calculated at close to 200,000 , while using total catch figures, production is close to 300,000 . These figures represent estimates respectively of Front production and total production.

For the 1968/1969 comparison, the Front figure increases and the total figure decreases. This result may be associated with an apparent move to the Front in 1969 of some seals which would normally have whelped in the Gulf (see below, Section 8). With this known abnormal year, it is probably better to use the 1966-1967/1968 comparison as more normal. This is shown graphically in Figure 4, where values of $K$ are plotted against catch $C$. The least squares line is fitted to data for the first three years only. For $K$ equal to 0 , estimates of catch = production are: for the Front - 192,000, for the total production 283,000. Expected further samples will allow refinement of these estimates.

## 6. Use of Catch Statistics

a. Total catch

Table 8 shows recent catches of young, uncontrolled at the Front save by season and partial abstention (1968) but controlled in the Gulf by Canadian regulation and Norwegian abstention since 1965.

Inspection of Front catches of young from 1960 to 1970 shows that in no year have catches exceeded 187,000 in spite of heavy catching effort and uncontrolled (unregulated) catch. This is itself strong evidence that production on the Front does not exceed, and may be less than, 200,000.

Figure 2 and Table 5 showed that, after years of highest catch (e.g. 1963, 1964, 1967) escapement of young was very low. Very low and maintained low escapement after high total catches in 1963, higher than normal in the Gulf, slightly lower than normal at the Front, suggest the importance of mixing between stocks.
b. Whitecoats and beaters

Mr. T. Øritsland (MS, 1967) demonstrated a variant of this method. If all the young are taken as whitecoats, none will escape to be taken at one month of age as "beaters," hence the beater catch will decline. He demonstrated a progressive decline in beater catches by the Norwegian fleet, indicating that a progressively increased proportion of production was being taken.

In order to convert to absolute production, the Canadian (and other) catches of young must be added. Unfortunately Canadian catch statistics did not separate whitecoats and beaters till 1966. In 1966 catches of young at the Front were not maximal $(168,000)$, 1968 was a beater season, and in 1969 the Gulf ice was abnormal so that some whelping females from the Gulf probably moved to the Front. This leaves us 1967, a year of maximal catch of young at the Front $(187,000)$. Indeed in this year beater catches were exceedingly low ( 1700 only) and whitecoat catches 181,000 . (An intermediate category of 'ragged jacket' accounts for the rest.) The combined catch of 187,000 young seals at the Front was therefore very likely close to total production.

## 7. Discussion

If the two populations mix, as now seems very probable, then removal annually of a high proportion of Front production plus a substantial amount of Gulf production will cause the population to decline. In quantitative terms, if production now equals 300,000 and actual removal averages 218,000 (Table 8) while sustainable yield is 125,000, the population as a whole must be declining. Is there any direct evidence that this is so?

First, to revert to Table 8 and the method of Section 5a. Larger maximal catches of young were taken at the Front in the 1950's (up to 248,000 ) than in the 1960 's (up to 188,000 ). This suggests a decline in production since if anything, catching effort has increased, due to experience of skippers and use of more powerful ships. Decline was probably faster in the 1950's since large numbers of adult and immature as well as young seals were then taken, until imposition of a closing date in 1961. Perhaps also the Gulf population, managed since 1965, has contributed part of its production to the overexploited Front. I do not think that the constant mean catches of young (Table 8) are an objection to my thesis, as has been suggested by the sealing industry. On the contrary, I think that they only mean that an increasing proportion of young is being killed.

A second line of evidence comes from a long series of samples, now extending over 20 years, from the net fishery on the North Shore of the Gulf of St. Lawrence at La Tabatiere (Age reading of this series was begun by Dr. H. D. Fisher and continued, with cross-checking of ages, by the author ). In the early 1950's this sample contained a lot of
older immatures or young mature adults, aged 4-6 years. In recent years this proportion has declined (Fig. 1). This can only mean, either that we are underfishing the adults so that they are accumulating, or that we are overfishing the young seals, so that recruitment steadily declines to the population as a whole. All the other evidence cited in this paper favours the second hypothesis. Again, this evidence, so interpreted, favours a high degree of mixing between populations, since it is unlikely that the Gulf, from where these samples come, has itself been overexploited under a controlled fishery.

## 8. Mixing of Stocks

Some evidence has been discussed above. Further evidence is as follows:

In 1969, abnormally restricted pack ice in the southern Gulf interfered with whelping of harp seals. After 32,000 young had been killed, the hunt was called off by Government order. Careful Governmental survey found only 6600 more young. There was no ice elsewhere in the Gulf. Therefore an estimate of production in the Gulf in 1969 was less than 40,000 . This is below the normal production estimated at some 100,000 . The other whelping seals must either have aborted at sea, or looked for and whelped on ice elsewhere. There was no evidence for large numbers of washed-up corpses of neonates, and the second alternative seems logical. The nearest suitable ice was in Hamilton Inlet, Labrador, where a large, compact herd whelped. Here, although the herd was in Canadian territorial limits for the most part so that few Norwegian ships took part in the whitecoat fishery, the total catch of young at the Front was 187,000 young animals, equal to any other maximum (Table 8). Is there any evidence that Gulf animals were involved? I think so, extending the method of Section 5 b above.

If many seals moved to the Front to whelp, escapement of the 1969 year-class will not be much below the expected level after a total catch of 220,000 young (Fig. 3), i.e. fair to poor (index 1 or 2 ) survival. But if they stayed in the Gulf and failed to whelp successfully, then the total Front production was taken, as well as the reduced production in the Gulf, so that escapement would have been very poor (index 0).

In fact, Front samples for 1970 (Tables 5d, e and 6) show that escapement of the 1969 year-class was about one third higher than for 1966 and 1967. If K, the ratio of one year-olds to total animals in age samples (method of Section 5b) is plotted against Front catch for the 4 years of available data from 1967 to 1970 (Fig. 4), in 1970 it falls above the least squares line of fit of the other 3 years, giving an increased value for production at the Front. But if plotted against total catch in 1970, it falls somewhat below the line, giving a lowered value for total production. This suggests to me that most animals which did not whelp in the Gulf whelped on the Front.

During 1969 the ratio of whelping animals was more than 187,000 on the Front to 40,000 in the Gulf, or higher than 4.5 to 1 . In 1970, aerial survey showed the ratio to be nearer 3:2. Therefore, animals had returned to the Gulf. The effect of delayed whelping in the Gulf (and
on the Front too for animals that moved) due to poor ice was apparently reflected in abnormally late whelping in the Gulf in 1970. One large patch was actively whelping between about 8 and 16 March, as compared with normal dates of about 1 to 7 March. (This observation also suggests that the date of whelping in March is fixed by the date of mating in April of the previous year, and not by the date of implantation, which occurs in the arctic about the end of July.)

The simplest hypothesis available at the present time from all the above evidence is that the seals mix freely on the southward migration in January when they feed for some 2 months in ice-free waters before whelping. They then whelp on the nearest available ice, beginning in late February. While this hypothesis does not rule out the possibility of some homing to, for instance, Gulf waters by the same animals yearly, its truth would rule out the possibility that sharp morphological or biochemical differences might be detectable between the populations except for phenotypic influences on the young (e.g. differences in weather which could affect hair quality). A more serious consequence would be that if, as demonstrated above, the Front population is being heavily overfished, and, if the populations mix, this effect will spread to the Gulf population which will also decline. Probably this is happening now. Capture-recapture estimates in the Gulf show a progressive decline (Table 3), and an increase in percentage of recaptured young, and production in the southern Gulf has almost certainly now sunk below 100,000 young.

## 9. Are Any Seals Being Missed?

The possibility of far northern harp seals was explored by flights along the Labrador coast in 1960 (Sergeant and Fisher, MS, 1960). None were found in March north of Hamilton Inlet, the ice being unsuitable due to lack of open water leads until too far out to sea. If such seals exist, they must whelp in April, and may contribute to small numbers of whitecoats seen at the Front late in the season in some years. Catches from these would enter catch figures so that the methods of Section 5 have allowed for them.

More important are a variably sized herd in the northern Gulf, whelping on the rather small and variable amount of ice entering through the Strait of Belle Isle. From catch figures they were known to reach a maximum of perhaps 50,000 in 1965 and 35,000 in 1967. The minimum may be a few hundred seen, for instance, by survey in March 1970. Again such catches enter catch figures, since the seals are taken sometimes by the larger ships and regularly by longliners active in the northern Gulf in April. Thus this production has also been taken into account in the methods of Section 5 .

If the method of Section $6 a$ is applied to total catches it is seen that total catches in the 1960's did not exceed 280,000 (Table 8) less than our postulated 300,000 total production. Big production in the north, occasionally entering the catches, would surely raise catch levels occasionally to a higher figure. Therefore I think that such added northern production is negligible. Higher total catches in the 1950's, up to 350,000 young seals, can be attributed to a higher production at that time.

## 10. Management

The 1971 quota of 200,000 seals of all species assigned to ships is divided 50,000 to the Gulf and 150,000 to the Front. These could all be taken as young harp seals, since the estimated production in each area will allow such catches, and because the fishery for young seals, (beginning about 12 March) comes before the fishery for hood seals on the Front (beginning about 20 March) and the fishery for moulting adult and immature harp seals (beginning about 30 March ). If this quota is fully taken as young harp seals some 50,000 young will be left available in each area, according to the present thesis, to be taken by smaller craft if these are not subject to the 200,000 quota. The numbers of these smaller vessels are known to be increasing rapidly as prices of harp seal furs remain firm.

In this way the proposed 1971 quota, as presently understood, would allow almost total destruction of the 1971 year-class of harp seals. If the population of western harp seals is to remain even at its present, reduced level, the quota must be cut to no more than 125,000 young harp seals taken by all agencies, or else the starting date for sealing must be made so late that the catch of the more highly dispersed "beaters" is no more than this figure. As with the International Whaling Commission a decade ago a major difficulty towards effective regulation is that there exists an excessive and growing catching fleet, the costs of construction of which must be amortized. I have previously suggested (Sergeant, MS, 1966) that the catching fleet should be halved, from some 24 to 12 large vessels, to maintain present sustainable yield and take it efficiently. Now in 1971, this measure would be inadequate. I now suggest that as many sealing ships as possible, most of which constitute a long-distance fleet, be diverted to fishing the stocks of crabeater seals Lobodon carcinophagus and other species in the Antarctic. Their biology, as seen from a pioneering Norwegian voyage, has been described by $\emptyset$ rits land (1970). The sustainable yield of harp seals could then be taken, and the stock possibly allowed to increase, by leaving such ships as are fit only for nearshore waters, together with the increasing fleet of longliners. While such a proposal is outside the scope of formal consideration by a regional organization such as ICNAF, possibly ICNAF should consider an approach to the Antarctic Treaty organization if a diversion of the kind suggested is recommended.

## 11. Summary

Present production of young harp seals in the western North Atlantic is calculated by four independent methods to be about 300,000 and declining. Under normal ice conditions some 200,000 are believed produced on the Front ice in Subareas 2 and 3, and 100,000 in the Gulf of St. Lawrence, Subarea 4. Shifts of population under abnormal ice conditions, and other evidence, suggest intermixing of stocks between the two subareas. The combined sustainable yield at present is estimated to be no more than 125,000 young harp seals. Present annual catches average 218,000 young and effort is increasing. Diversion of the more mobile part of the fleets to sub-Antarctic seal stocks is suggested as an alternative to continued overexploitation.

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Table 1. Reproductive rates of female harp seals collected in the Gulf of St. Lawrence, 1965-1969.

| Age at parturition | Number |  |  |  | Percent pregnant |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | Immature | With regressed corpus luteum | Pregnant | Total |  |
| (yrs) - - - - - |  |  |  |  |  |
| 1 | 25 |  |  | 25 |  |
| 2 | 35 |  |  | 35 |  |
| 3 | 69 |  |  | 69 |  |
| 4 | 77 |  | 2 | 79 | 2.5 |
| 5 | 74 | 1 | 21 | 96 | 21.9 |
| 6 | 35 | 4 | 65 | 104 | 62.5 |
| 7 | 14 | 1 | 81 | 96 | 84.4 |
| 8 | 7 | 5 | 66 | 78 | 84.6 |
| 9 | 3 | 2 | 51 | 56 | 91.1 |
| 10 | 4 |  | 50 | 54 | 92.6 |
| 11 |  | 5 | 43 | 48 | 89.6 |
| 12 | 1 | 2 | 31 | 34 |  |
| 13 | 2 | 4 | 23 | 29 |  |
| 14 | 1 | 1 | 39 | 41 |  |
| 15 |  | 2 | 31 | 33 |  |
| 16 |  | 1 | 23 | 24 |  |
| 17 |  | 4 | 15 | 19 |  |
| 18 | 2 | 1 | 11 | 14 |  |
| 19 | 1 | 1 | 12 | 14 | 88.1 |
| 20 |  | 2 | 4 | 6 |  |
| 21 |  |  | 4 | 4 |  |
| 22 |  |  | 7 | 7 |  |
| 23 |  |  | 2 | 2 |  |
| 24 |  |  | 2 | 2 |  |
| 25 |  | 3 | 1 | 4 |  |
| 26 |  |  | 1 | 1 |  |
| 27 |  |  | 1 | 1 |  |
|  | 350 | 39 | 586 | 975 |  |

Table 2. Aerial photographic survey of whelping seals, 1970.

Table 3. Results of capture-recapture tagging 1964 to 1970.


[^13]| Table 4. Sources of age samples analyzed in Table 5. | Samples arranged in seasonal sequence. |  |  |  |
| :--- | :--- | :--- | :--- | :--- |
| Number | Location | Behaviour of animals | Season | Known biases |
| a. | Labrador | Migrating south | November-December | Few 1-4 year animals |
| b. | Northern Gulf | Migrating south | January | Few 1-4 and >10 year animals |
| c. | Western Gulf | Wintering | January-April | Few >5 year animals |
| d. | Front shore catch | Wintering | January-April | Good |
| e. | Front ship catch | Spring moult | April | Excess 1-2 year animals |
| f. | Arctic | Summering |  | July-October |

Table 5. Index of survival (escapement) of year-classes of harp seals 1960-69, assessed subjectively from the series of age samples identified in Table 4, as very good $=3$, good $=2$, poor $=1$, very poor $=0$. Data in Appendix Table 1. No data shown, where they exist, indicates an expected bias or inadequate data.
$\begin{array}{lllllllllll}\text { Year class } & 1960 & 1961 & 1962 & 1963 & \underline{1964} & \underline{1965} & \underline{1966} & \underline{1967} & \underline{1968} & \underline{1969}\end{array}$
Sample
1970

| e |  |  |  |  |  | 2 | 2 | 0 | 3 | 2 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| d | 3 | 3 | 3 | 1 | 2 | 2 | 2 | 0 | 3 | 2 |
| c |  |  |  |  |  | - | 2 | 3 | 2 | 1 |
| b | 3 | 2 | 2 | 1 | 1 | 3 |  |  |  |  |
| a | 1 | 2 | 1 | 1 | 2 | 3 |  |  |  |  |

1969

| f | 2 | 0 | 1 | 2 | 1 | 3 | 2 | 1 | 3 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| e |  |  |  |  |  |  |  |  | 2 |
| d |  |  |  |  |  |  |  | 1 | 2 |
| c |  |  |  |  |  | - | 2 | 1 | 3 |
| b | 1 | 2 | 2 | 1 | 2 | 3 |  |  |  |
| a | 2 | 2 | 2 | 0 | 1 | 2 | 1 | 1 | 2 |

1968

| e | 3 | 2 | 1 | 1 | 1 | 3 |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| d | 3 | 2 | 1 | 0 | 1 | 1 | 1 | 1 |
| b | 2 | 2 | 2 | 1 | 2 | 3 |  |  |

1967

| f | 2 | 2 | 2 | 1 | 2 | 2 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| e | 3 | 3 | 2 | 1 | 0 |  |  |
| d | 3 | 2 | 1 | 1 | 0 | 1 | 1 |
| b | 2 | 3 | 2 | 1 |  |  |  |
| a | 2 | 2 | 2 | 2 |  |  |  |

1966


1965

| e | 2 | 1 | 1 | 1 |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| b | 3 | 3 |  |  |  |
| a | 2 | 2 |  |  |  |

1964

| b | 3 | 2 | ------ |  |
| :--- | :--- | :--- | :--- | :--- |
| a | 3 | 3 | 1 | 2 |

1963

| e | 3 | 3 | 1 |
| :--- | :--- | :--- | :--- |
| b | 2 | 2 | 1 |
| a | 2 | 3 | 1 |

Table 5. Continued.
$\begin{array}{lllllllllll}\text { Year class } & 1960 & 1961 & 1962 & \underline{1963} & \underline{1964} & \underline{1965} & \underline{1966} & \underline{1967} & \underline{1968} & \underline{1969}\end{array}$
Sample

| 1962 |  |  |
| :---: | :---: | :---: |
| f | 2 | 2 |
| e | 3 | 2 |
| b |  |  |
| a |  |  |

1961
$\begin{array}{ll}\text { e } \\ \text { b } & -----\end{array}$

| Total | 62 | 56 | 32 | 18 | 15 | 28 | 13 | 8 | 20 | 5 |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| Maximal | 72 | 72 | 63 | 51 | 36 | 36 | 24 | 24 | 24 | 9 |
| Ratio $=$ <br> Index | 0.86 | 0.78 | 0.51 | 0.35 | 0.41 | 0.78 | 0.54 | 0.33 | 0.83 | 0.56 |
| Catch <br> $\left(\mathrm{XlO}^{3}\right)$ | 156 | 169 | 207 | 270 | 266 | 183 | 252 | 280 | 156 | 220 |

Table 6. Escapement at one year of age, correlated with catch of young and expressed as percentage of total sample, for samples from (a) St. Anthony and (b) the Front icefields.

| Year <br> class | atch of young (C)$\times 10^{-3}$ |  | Number ¢ percent of juveniles in samples from |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | St. Anthony |  |  | Front icefields |  |  |
|  |  |  | $\underline{1} \mathrm{yr}$ | Total | Ratio(K) | $\underline{19 r}$ | Total | Ratio(K) |
|  | Front | Total |  |  |  |  |  |  |
| 1966 | 180 | 264 | 18 | 315 | . 057 | 77 | 405 | . 190 |
| 1967 | 184 | 276 | 7 | 201 | . 035 | 84 | 576 | . 145 |
| Mean | 182 | 270 |  |  | . 046 |  |  | . 168 |
| 1968 | 98 | 155 | 87 | 205 | . 424 | 62 | 107 | . 579 |
| 1969 | 187 | 220 | 41 | 571 | . 072 | 105 | 431 | . 244 |

Table 7. Calculation of Front and total production comparing mean figures of catch and escapement for 1966-67 with 1968 and 1968 with 1969.

| Catch $\qquad$ <br> Age sample | Front |  | Total |  |
| :---: | :---: | :---: | :---: | :---: |
|  | St. Anthony | Front icefields | St. Anthony | Front icefields |
| Years compared | 1966-67/68 |  |  |  |
| Estimated production | 192,704 | 216,336 | 283,994 | 317,007 |
|  | $(204,520)$ |  | $(300,500)$ |  |
| Years compared | 1968/69 |  |  |  |
| Estimated production | 205,205 | 251,824 | 233,295 | 267,343 |
|  | $(228,514)$ |  | $(250,319)$ |  |

Table 8. Catches of young harp seals $\times 10^{-3}$.

| Year | Gulf of <br> St. Lawrence | Front | Total | 5-yr mean |
| :---: | :---: | :---: | :---: | :---: |
| 1950 | 31 | 195 | 226 |  |
| 1951 | 90 | 229 | 319 |  |
| 1952 | 63 | 135 | 198 | 223 |
| 1953 | 32 | 166 | 198 | 228 |
| 1954 | 74 | 101 | 175 | 233 |
| 1955 | 94 | 158 | 252 | 226 |
| 1956 | 93 | 248 | 341 | 215 |
| 1957 | 74 | 91 | 165 | 228 |
| 1958 | 90 | 51 | 141 | 208 |
| 1959 | 62 | 177 | 239 | 174 |
| 1960 | 85 | 71 | 156 | 182 |
| 1961 | 41 | 128 | 169 | 208 |
| 1962 | 89 | 118 | 207 | 214 |
| 1963 | 110 | 160 | 270 | 219 |
| 1964 | 84 | 182 | 266 | 236 |
| 1965 | 90 | 93 | 183 | 250 |
| 1966 | 84 | 180 | 264 | 227 |
| 1967 | 92 | 184 | 276 | 218 |
| 1968 | 57 | 98 | 155 |  |
| 1969 | 33 | 187 | 220 |  |
| (1970 preliminary | 57 | 150 | 207) |  |

Appendix Table 1. Age samples.
(a) Labrador (November of previous year to January of current year).

| Age (yrs) | 1970 | 1969 | $\underline{1968}$ | 1967 | 1966 | 1965 | 1964 | 1963 | $\underline{1962}$ | 1961 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| nearly |  |  |  |  |  |  |  |  |  |  |
| 1 | 21 | 5 |  | 6 | 1 | 4 | 43 | 26 | 6 |  |
| 2 | 60 | 4 |  | 11 | 2 | 10 | 42 | 83 | 15 |  |
| 3 | 67 | 29 |  | 17 | 3 | 24 | 50 | 54 | 16 |  |
| 4 | 63 | 18 |  | 24 | 5 | 34 | 32 | 33 | 30 |  |
| 5 | 68 | 11 |  | 30 | 7 | 24 | 32 | 26 | 21 |  |
| 6 | 44 | 6 |  | 25 | 5 | 18 | 42 | 21 | 7 |  |
| 7 | 24 | 11 |  | 27 | 4 | 15 | 29 | 8 | 14 |  |
| 8 | 22 | 10 |  | 14 |  | 16 | 35 | 10 | 16 |  |
| 9 | 25 | 11 |  | 10 | 4 | 4 | 23 | 12 | 13 |  |
| 10 | 24 | 2 |  | 9 | 2 | 5 | 27 | 15 | 16 |  |
| 11 | 13 | 3 |  | 11 | 1 | 5 | 18 | 14 | 10 |  |
| 12 | 8 | 6 |  | 5 |  | 6 | 21 | 10 | 3 |  |
| 13 | 8 | 1 |  | 6 |  | 7 | 20 | 9 | 11 |  |
| 14 | 5 | 1 |  | 7 |  | 7 | 21 | 14 | 8 |  |
| 15 | 10 |  |  | 7 |  | 10 | 7 | 11 | 23 |  |
| 16 | 2 | 1 |  | 6 | 1 | 6 | 11 | 9 | 10 |  |
| 17 | 9 |  |  | 3 | 2 | 5 | 12 | 11 | 4 |  |
| 18 | 6 | 1 |  | 4 |  | 4 | 6 | 4 | 4 |  |
| 19 | 6 |  |  | 6 | 2 | 4 | 14 | 12 | 6 |  |
| 20 | 6 | 1 |  | 4 |  | 1 | 6 | 10 | 12 |  |
| 21 | 1 | 1 |  | 4 |  | 2 | 6 | 6 | 2 |  |
| 22 | 1 |  |  |  | 1 | 1 | 3 | 8 | 5 |  |
| 23 | 3 |  |  | 1 |  |  | 5 | 10 | 3 |  |
| 24 | 1 |  |  | 2 |  | 2 | 2 | 6 | 2 |  |
| 25 |  |  |  | 1 | 2 | 2 | 6 | 2 | 3 |  |
| 26 | - |  |  |  |  | 1 | 3 | 2 |  |  |
| 27 |  |  |  |  |  | 1 |  | 1 | 3 |  |
| 28 |  |  |  | 1 |  |  | 1 | 2 | 2 |  |
| 29 |  |  |  |  |  |  |  | 3 |  |  |
| 30 |  |  |  | 2 |  |  |  | 4 |  |  |
| 30+ |  |  |  |  |  |  | 2 |  | 3 |  |
| N | 497 | 122 |  | 243 | 42 | 218 | 519 | 439 | 266 |  |

Appendix Table 1 (Continued). Age samples.
(b) Northern Gulf (January).

| Age (yrs) | 1970 | $\underline{1969}$ | 1968 | 1967 | 1966 | $\underline{1965}$ | 1964 | 1963 | $\underline{1962}$ | 1961 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| nearly |  |  |  |  |  |  |  |  |  |  |
| 1 | 5 | 9 | 11 | 12 | 10 | 4 | 10 | 18 | 20 | 16 |
| 2 | 12 | 20 | 48 | 42 | 29 | 17 | 31 | 52 | 50 | 65 |
| 3 | 32 | 49 | 68 | 46 | 42 | 45 | 74 | 86 | 74 | 100 |
| 4 | 55 | 69 | 56 | 37 | 44 | 72 | 64 | 90 | 95 | 126 |
| 5 | 77 | 71 | 39 | 58 | 42 | 71 | 65 | 80 | 95 | 39 |
| 6 | 38 | 49 | 54 | 72 | 50 | 66 | 40 | 56 | 33 | 52 |
| 7 | 35 | 60 | 41 | 64 | 33 | 39 | 27 | 32 | 46 | 43 |
| 8 | 64 | 59 | 39 | 43 | 19 | 32 | 11 | 36 | 42 | 27 |
| 9 | 66 | 43 | 17 | 31 | 14 | 18 | 10 | 22 | 29 | 28 |
| 10 | 82 | 40 | 18 | 27 | 8 | 9 | 6 | 25 | 18 | 24 |
| 11 | 61 | 38 | 14 | 14 | 10 | 10 | 11 | 24 | 14 | 15 |
| 12 | 42 | 22 | 14 | 15 | 5 | 12 | 5 | 20 | 17 | 20 |
| 13 | 37 | 20 | 7 | 10 | 7 | 16 | 6 | 11 | 11 | 16 |
| 14 | 30 | 16 | 9 | 16 | 7 | 8 | 3 | 17 | 13 | 7 |
| 15 | 32 | 24 | 10 | 12 | 7 | 8 | 4 | 22 | 21 | 12 |
| 16 | 27 | 18 | 6 | 4 | 7 | 6 | 2 | 15 | 7 | 5 |
| 17 | 30 | 11 | 9 | 9 | 3 | 5 | 4 | 13 | 9 | 3 |
| 18 | 21 | 11 | 4 | 7 | 1 | 2 | 1 | 9 | 11 | 2 |
| 19 | 17 | 5 | 5 | 4 | 1 | 6 |  | 4 | 1 |  |
| 20 | 16 | 4 | 2 | 5 | 2 | 3 | 2 | 10 | 4 | 7 |
| 21 | 10 | 5 | 1 | 3 | 1 | 3 |  | 5 | 1 |  |
| 22 | 9 | 2 | 3 | 4 | 1 | 1 | 2 | 5 | 6 |  |
| 23 | 4 |  |  |  | 1 | 2 |  | 4 | 1 | 2 |
| 24 | 5 | 1 |  | 2 |  | 1 |  | 2 | 1 | 1 |
| 25 | 6 |  | 1 | 3 |  | 1 | 1 | 1 | 2 | 4 |
| 26 | 3 |  | 1 |  | 1 | 1 |  | 3 |  | 1 |
| 27 | 3 |  | 2 |  |  |  |  | 2 |  |  |
| 28 | 4 |  |  |  | 2 |  |  | 3 |  | 2 |
| 29 |  |  |  |  |  |  |  |  |  |  |
| 30 | 2 |  |  |  |  | 1 | 1 | 5 |  | 2 |
| N | 824 | 646 | 479 | 540 | 347 | 459 | 380 | 672 | 621 | 619 |

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Appendix Table 1 (Continued). Age samples.

| Age (yrs) | (c) Western Gulf <br> January-April |  | (d) Front shore catch January-April |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1970 | $\underline{1969}$ | 1970 | 1969 | 1968 | 1967 |
| 1 | 31 | 37 | 41 | 87 | 7 | 17 |
| 2 | 69 | 14 | 41 | 13 | 8 | 14 |
| 3 | 89 | 21 | 15 | 13 | 8 | 1 |
| 4 | 45 | 10 | 25 | 13 | 8 | 9 |
| 5 | 40 | 11 | 23 | 11 | 5 | 11 |
| 6 | 16 | 2 | 19 | 10 | 9 | 21 |
| 7 | 8 | 5 | 14 | 5 | 12 | 26 |
| 8 | 12 | 4 | 43 | 7 | 15 | 21 |
| 9 | 8 | 2 | 49 | 1 | 16 | 26 |
| 10 | 8 | 11 | 39 | 6 | 15 | 16 |
| 11 | 4 | 6 | 24 | 4 | 10 | 21 |
| 12 | 4 | 2 | 28 | 4 | 13 | 14 |
| 13 | 1 | 5 | 26 | 5 | 9 | 16 |
| 14 |  | 2 | 19 | 3 | 7 | 15 |
| 15 | 2 | 1 | 29 | 3 | 9 | 13 |
| 16 | 1 | 1 | 30 | 2 | 11 | 17 |
| 17 |  | 5 | 20 | 6 | 10 | 12 |
| 18 | 1 |  | 12 | 2 | 7 | 9 |
| 19 | 1 | 2 | 18 | 3 | 5 | 8 |
| 20 | 3 |  | 22 | 2 | 3 | 11 |
| 21 | 1 |  | 10 | 2 | 4 | 3 |
| 22 |  | 1 | 5 | 1 | 4 | 2 |
| 23 |  |  | 7 | 2 | 2 | 3 |
| 24 |  | 1 | 5 |  | 1 | 5 |
| 25 |  |  | 3 |  |  | 1 |
| 26 |  |  | 1 |  |  | 1 |
| 27 |  |  |  |  | 1 |  |
| 28 |  |  |  |  |  | 1 |
| 29 |  |  |  |  |  |  |
| 30 |  |  | 1 |  | 1 |  |
| 30+ |  |  |  |  | 1 | 1 |
| N | 359 | 144 | 571 | 205 | 201 | 315 |

Appendix Table 1 (Continued). Age samples.
(e) Ships' catch of moulters. All or mostly from Front except 1966 when from Gulf.

| Age (yrs) | 1970 | 1969 | $\underline{1968}$ | 1967 | $\underline{1966}$ | 1965 | 1964 | $\underline{1963}$ | 1962 | 1961 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 105 | 62 | 84 | 176 | 120 | 32 |  | 111 | 201 | 122 |
| 2 | 98 | 2 | 77 | 61 | 100 | 13 |  | 121 | 245 | 34 |
| 3 | 11 | 6 | 48 | 18 | 39 | 13 |  | 106 | 61 | 25 |
| 4 | 17 | 1 | 20 | 18 | 40 | 15 |  | 55 | 56 | 30 |
| 5 | 18 | 2 | 17 | 29 | 45 | 23 |  | 46 | 39 | 11 |
| 6 | 12 | 4 | 23 | 46 | 45 | 22 |  | 61 | 17 | 13 |
| 7 | 15 | 1 | 27 | 40 | 44 | 7 |  | 67 | 15 | 15 |
| 8 | 8 | 6 | 39 | 30 | 31 | 2 |  | 66 | 16 | 9 |
| 9 | 13 | 2 | 23 | 23 | 15 | 2 |  | 63 | 17 | 8 |
| 10 | 18 |  | 26 | 16 | 17 | 5 |  | 72 | 6 | 17 |
| 11 | 13 | 3 | 19 | 23 | 23 | 1 |  | 70 | 8 | 10 |
| 12 | 20 | 1 | 18 | 18 | 17 | 7 |  | 51 | 13 | 3 |
| 13 | 12 | 2 | 16 | 14 | 15 |  |  | 57 | 6 | 4 |
| 14 | 6 | 2 | 24 | 19 | 17 | 5 |  | 56 | 9 | 6 |
| 15 | 11 | 2 | 20 | 20 | 14 | 4 |  | 42 | 12 | 1 |
| 16 | 13 | 2 | 20 | 14 | 14 | 2 |  | 54 | 4 | 5 |
| 17 | 7 |  | 11 | 18 | 10 | 1 |  | 36 | 12 | 4 |
| 18 | 7 |  | 17 | 20 | 17 | 3 |  | 24 | 5 | 2 |
| 19 | 7 |  | 16 | 21 | 10 | 3 |  | 18 | 4 | 3 |
| 20 | 6 |  | 11 | 15 | 11 | 1 |  | 18 | 9 | 2 |
| 21 | 4 | 1 | 6 | 8 | 8 | 2 |  | 17 | 1 |  |
| 22 | 5 | 2 | 5 | 8 | 3 | 1 |  | 12 | 4 | 2 |
| 23 | 3 | 2 | 4 | 8 | 7 |  |  | 9 |  | 1 |
| 24 | 3 | 1 | 2 | 4 | 4 |  |  | 8 | 1 |  |
| 25 |  |  | 1 | 4 | 2 |  |  | 4 | 1 | 1 |
| 26 | 1 | 1 | 1 | 1 | 4 | 3 |  | 4 | 1 | 1 |
| 27 |  |  | 2 | 3 | 1 |  |  | 1 | 1 |  |

Appendix Table 1 (Continued). Age samples.
(e) Continued.

| Age (yrs) | 1970 | 1969 | $\underline{1968}$ | 1967 | 1966 | 1965 | 1964 | 1963 | 1962 | 1961 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 28 | 2 |  |  |  | 1 |  |  | 1 | 1 |  |
| 29 |  | 1 | 1 |  |  |  |  |  |  |  |
| 30 |  |  |  | 4 |  |  |  |  |  |  |
| 30+ |  |  |  |  | 3 |  |  |  |  |  |
| N | 431 | 107 | 578 | 679 | 677 | 167 |  | 1247 | 767 | 330 |

No. of ships
$\begin{array}{lllll}2 & 1 & 4 & 4^{\mathrm{X}} & 1\end{array}$
1*
1+
*USSR (Khuzin, 1963)
$x_{\text {including }} 1$ Norwegian
${ }^{+}$USSR (from Popov and Timoshenko, 1965)

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Appendix Table 1 (Continued). Age samples.
(f) Arctic, July-September.

| Age (yrs) | Baffin Is land |  | West Greenland |
| :---: | :---: | :---: | :---: |
|  | 1969 | $\underline{1967}$ | 1962 |
| 0 | 9 | 5 | 275 |
| 1 | 23 | 11 | 37 |
| 2 | 11 | 25 | 19 |
| 3 | 19 | 13 | 3 |
| 4 | 23 | 9 | 1 |
| 5 | 12 | 17 | 3 |
| 6 | 17 | 20 | 2 |
| 7 | 9 | 14 | 3 |
| 8 | 9 | 10 | 1 |
| 9 | 19 | 14 | 2 |
| 10 | 14 | 9 | 1 |
| 11 | 11 | 14 |  |
| 12 | 9 | 8 |  |
| 13 | 8 | 10 | 2 |
| 14 | 17 | 12 |  |
| 15 | 5 | 7 | 9 |
| 16 | 6 | 7 |  |
| 17 | 5 | 6 |  |
| 18 | 5 | 2 |  |
| 19 | 10 | 5 |  |
| 20 | 3 | 4 |  |
| 21 | 1 | 3 |  |
| 22 | 3 | 2 |  |
| 23 | 6 | 1 |  |
| 24 | 2 | 2 |  |
| 25 | 1 |  |  |
| 26 |  |  |  |
| 27 | 1 |  |  |
| 28 |  |  |  |
| 29 |  |  |  |
| 30 |  |  |  |
| 30+ |  |  |  |
| N | 264 | 232 | 358 |



Fig. 1. Age samples of southward migrant seals from the same site, La Tabatiere, in 1953 and 1970.


Fig. 2. Age samples from St. Anthony, Newfoundland, and the Front icefields, 1967-1970.


Fig. 3. Index of survival plotted against total catch for yearclasses 1960 to 1969. See text.


Fig. 4. Index $K$, the ratio of young to total seals in four annual age samples (1967-1970) from St. Anthony, Newfoundland, plotted against total catch of young of the same year-class, and catch of young at the Front, to show the aberrant year, 1969. See text.
15. The status of Norwegian studies of harp seals at Newfoundland ${ }^{1}$
by Torger Øritsland
Institute of Marine Research
Bergen, Norway

## Introduction

Recent Norwegian seal research at Newfoundland was started in 1964. Field work in that season was concentrated on collection of samples for blood protein studies and studies of breeding biology. In 1965 physiological studies were performed, and in 1967, 1969, 1970 and 1971 material and data were collected for general biological studies of both harp and hooded seals. No field work was done in 1966 or 1968, but a good sample for age analysis of harp seals was collected by sealers in 1968. Provisional results of these studies were reported to the ICNAF Panel A (Seals) In 1968 and 1969 (Naevdal, MS 1968; $\emptyset$ ritsland, MS 1969a and $b$ ).

Some effort has also been made to revise and analyze Norwegian sealing statistics. Provisional analyses were reported to the ICNAF Seal Panel in 1966 and 1967 ( $\emptyset$ ritsland, MS 1966 and MS 1967b).

The purpose of this report is to sumarize results at hand from Norwegian studies of harp seals at Newfoundland from 1964 to 1971.

## Distribution of seals

Observed and reported ice edges and patches of seals have been plotted in 1965, 1967, 1969, 1970 and 1971 as shown in Figs. 1-5. In these years one, two or more separate harp seal breeding lairs have formed in the area from Hamilton Inlet southwards to Roundhill Island before the middle of March. During lactation these lairs drift south with the pack-ice, and may end up in the Strait of Belle Isle or Notre Dame Bay before the last moulted pups have taken to sea or been killed in late March or early April. Patches of juvenile seals, bedlamers, form in the area from Funk Island to Gray Islands off Notre Dame Bay or further north in late March. As bedlamers gradually mix with adult males in moulting patches in early April, the animals move north, and may be found of $f$ Hamilton Inlet or further north in late April.

## Populations

Results from blood protein studies (Naevdal, MS 1968 and 1969) failed to reveal any significant difference between the breeding herds of harp seals in the North Atlantic.

A total of 127 harp seal pups have been tagged on the Front off Newfound-land-Labrador in 1969, 1970 and 1971. Out of these a total of 14 were recaptured in the hunting area before the seasons ended. Two animals have been recaptured off West Greenland in the autumn of the year they were tagged: one near Jakobshavn in August 1969 and one near Ritenbank in October 1970. Finally, two animals were recaptured on the Front during the hunting season 1971, one year after they were tagged. So far Norwegian taggings have not yielded definite

[^14]information on the distinctness of harp seal populations, but none of the 441 harp seal pups which have been tagged near Jan Mayen through the years, has been recaptured west of Cape Farvel.

Some 450 harp seal skulls which have been collected at Newfoundland (147), in the Jan Mayen area (about 100) and in the Barents Sea, are deposited in the Zoological Museum in Trondheim. No results are yet available from this collection.

Age composition of catches
Age-group frequencies in samples collected in 1964, 1967 and 1968 are given in Table 1. Hrifortunately, the small 1964 sample contains 33 specimens of breeding femajes, and therefore it is not representative for moulting animals. Howeyer, the other two samples are not "contaminated" in this way. Data from samples of moulting animals collected in 1969 and 1970 are presented in Table 2. An additional sample of 559 moulting harps collected in 1971 has not been processed yet.

Table 1, Age-group frequencies of moulting harp seals in Front area, Newfoundland, from samples collected in 1964, 1967 and 1968.

| $\begin{aligned} & \text { Age- } \\ & \text { group } \end{aligned}$ | 1964 |  | 1967 |  | 1968 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | No. | \% | No. | \% | No. | \% |
| 1 | 3 | 2.4 | 55 | 19.6 | 135 | 19.2 |
| 2 | 6 | 4.7 | 31 | 11.0 | 74 | 10.5 |
| 3 | 8 | 6.3 | 5 | 1.8 | 42 | 6.0 |
| 4 | 11 | 8.7 | 6 | 2.1 | 27 | 3.8 |
| 5 | 7 | 5.5 | 19 | 6.8 | 30 | 4.3 |
| 6 | 12 | 9.4 | 22 | 7.8 | 20 | 2.8 |
| 7 | 6 | 4.7 | 19 | 6.8 | 46 | 6.6 |
| 8 | 5 | 3.9 | 13 | 4.6 | 38 | 5.4 |
| 9 | 5 | 3.9 | 5 | 1.8 | 31 | 4.4 |
| 10 | 8 | 6.3 | 8 | 2.8 | 28 | 4.0 |
| 11 | 4 | 3.1 | 13 | 4.6 | 22 | 3.1 |
| 12 | 4 | 3.1 | 6 | 2.1 | 12 | 1.7 |
| 13 | 3 | 2.4 | 5 | 1.8 | 15 | 2.1 |
| 14 | 3 | 2.4 | 10 | 3.6 | 18 | 2.6 |
| 15 | 5 | 3.9 | 10 | 3.6 | 14 | 1.9 |
| 16 | 4 | 3.1 | 6 | 2.1 | 12 | 1.7 |
| 17 | 5 | 3.9 | 8 | 2.8 | 15 | 2.1 |
| 18 | 8 | 6.3 | 9 | 3.2 | 26 | 3.7 |
| 19 | 4 | 3.1 | 9 | 3.2 | 21 | 3.0 |
| 20 | 0 | 0.0 | 4 | 1.4 | 16 | 2.3 |
| 21 | 4 | 3.1 | 4 | 1.4 | 10 | 1.4 |
| 22 | 2 | 1.6 | 1 | 0.4 | 17 | 2.4 |
| 23 | 2 | 1.6 | 3 | 1.1 | 8 | 1.1 |
| 24 | 3 | 2.4 | 2 | 0.7 |  | 1.0 |
| 25 | 1 | 0.8 | 2 | 0.7 | 3 | 0.4 |
| 26 | 0 | 0.0 | 1 | 0.4 | 5 | 0.7 |
| 27+ | 4 | 3.1 | 5 | 1.8 | 10 | 1.4 |
| Sum <br> Undet. | 127 | 99.7 | 281 | 100.0 | $\begin{array}{r} 702 \\ 35 \end{array}$ | $\left(\begin{array}{c} 99.6 \\ 4.7 \% \\ \text { of total } \end{array}\right)$ |

Table 2. Age-group frequencies of moulting harp seals in Front area, Newfoundland, from samples collected in 1969 and 1970.

| Agegroup | 1969 |  | 1970 |  |
| :---: | :---: | :---: | :---: | :---: |
|  | No. | \% | No. | \% |
| 1 | 371 | 40.6 | 132 | 25.6 |
| 2 | 42 | 4.6 | 109 | 21.1 |
| 3 | 34 | 3.7 | 33 | 6.4 |
| 4 | 30 | 3.3 | 30 | 5.8 |
| 5 | 39 | 4.3 | 29 | 5.6 |
| 6 | 28 | 3.1 | 15 | 2.9 |
| 7 | 36 | 3.9 | 18 | 3.5 |
| 8 | 44 | 4.8 | 26 | 5.0 |
| 9 | 35 | 3.8 | 15 | 2.9 |
| 10 | 29 | 3.2 | 22 | 4.3 |
| 11 | 23 | 2.5 | 13 | 2.5 |
| 12 | 16 | 1.8 | 7 | 1.4 |
| 13 | 19 | 2.1 | 6 | 1.2 |
| 14 | 21 | 2.3 | 11 | 2.1 |
| 15 | 21 | 2.3 | 8 | 1.6 |
| 16 | 12 | 1.3 | 11 | 2.1 |
| 17 | 25 | 2.7 | 3 | 0.6 |
| 18 | 16 | 1.8 | 6 | 1.2 |
| 19 | 19 | 2.1 | 6 | 1.2 |
| 20 | 16 | 1.8 | 4 | 0.8 |
| 21 | 10 | 1.1 | 5 | 1.0 |
| 22 | 5 | 0.5 | 2 | 0.4 |
| 23 | 7 | 0.8 | 2 | 0.4 |
| 24 | 3 | 0.3 | 0 | 0.0 |
| 25 | 4 | 0.4 | 1 | 0.2 |
| 26 | 4 | 0.4 | 0 | 0.0 |
| 27+ | 6 | 0.7 | 2 | 0.4 |
| Sum | 915 | 100.2 | 516 | 100.2 |
| Undet. | 106 | $\text { (10.4 of } \begin{aligned} & \text { total) } \end{aligned}$ | 26 | $\left(4.8 \begin{array}{l} \text { of } \\ \text { total } \end{array}\right)$ |

It is still an open question, however, whether the age-composition of catches of moulting harp seals does represent the age-composition in the stock. Impressions from field work in all hunting areas of the North Atlantic and the information on distribution given in Figs. 1-5 suggest that adult males gather in patches near the lairs of breeding females shortly after the formation of the breeding lairs. Somewhat later the juvenile seals of both sexes congregate, and after the pairing season the patches of juveniles and the adult males gradually mix to form moulting patches (in early April on the Front). Adult females foin these moulters in increasing numbers during April.

Information from counts of pelts in 1967, 1969, 1970 and 1971 on the change in age- and sex-composition through April in patches of moulting harps is available, and a summary of all counts is presented in Table 3 and in Fig. 6B. Independent data from sampling for age analysis during April in 1967, 1969 and 1970 are presented in Table 4 and Fig. 6A. The data available up to

Table 3. Moulting harp seals in Front area, Newfoundland. Catch composition from counts of pelts by age-group (bedlamers and saddlers), sex and date in 1967, 1969, 1970 and 1971.

| $\begin{aligned} & \text { Date in } \\ & \text { April } \end{aligned}$ | $\begin{aligned} & \text { Sample } \\ & \text { size } \end{aligned}$ | Juveniles |  |  |  | Adults |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | $\mathrm{o}^{\text {a }}$ |  | 9 |  | ${ }^{\prime}$ |  | ¢ |  |
|  |  | No. | \% | No. | \% | No. | \% | No. | \% |
| 1967 |  |  |  |  |  |  |  |  |  |
| 12-14 | 237 | 30 | 12.7 | 25 | 10.5 | 163 | 68.8 | 19 | 8.0 |
| 23 | 119 | 8 | 6.7 | 9 | 7.5 | 74 | 62.2 | 28 | 23.5 |
| Subtotal | 356 | 38 | 10.7 | 34 | 9.6 | 237 | 66.6 | 47 | 13.2 |
| 1969 |  |  |  |  |  |  |  |  |  |
| 02-05 | 391 | 97 | 24.8 | 87 | 22.3 | 205 | 52.4 | 2 | 0.5 |
| 12-13 | 343 | 110 | 32.1 | 104 | 30.3 | 125 | 36.4 | 4 | 1.2 |
| 16-20 | 720 | 188 | 26.1 | 193 | 26.8 | 261 | 36.3 | 78 | 10.4 |
| 22 | 132 | 39 | 29.5 | 44 | 33.3 | 35 | 26.5 | 14 | 10.6 |
| Subtotal | 1,586 | 434 | 27.4 | 428 | 27.0 | 626 | 39.5 | 98 | 6.2 |
| 1970 |  |  |  |  |  |  |  |  |  |
| 19 | 76 | 32 | 42.1 | 29 | 38.2 | 13 | 17.1 | 2 | 2.6 |
| 28-29 | 280 | 75 | 26.7 | 80 | 28.6 | 79 | 28.2 | 46 | 16.4 |
| Subtotal | 356 | 107 | 30.1 | 109 | 30.6 | 92 | 25.8 | 48 | 13.5 |
| 1971 |  |  |  |  |  |  |  |  |  |
| 10 | 258 | 80 | 31.0 | 68 | 26.4 | 71 | 27.5 | 39 | 15.1 |
| 13 | 395 | 102 | 25.8 | 91 | 23.0 | 146 | 37.0 | 56 | 14.2 |
| Subtotal | 653 | 182 | 27.9 | 159 | 24.3 | 217 | 33.2 | 95 | 14.5 |
| Summary, $1967+1969+1970+1971$ |  |  |  |  |  |  |  |  |  |
| 01-05 | 391 | 97 | 24.8 | 87 | 22.3 | 205 | 52.4 | 2 | 0.5 |
| 06-10 | 258 | 80 | 31.0 | 68 | 26.4 | 71 | 27.5 | 39 | 15.1 |
| 11-15 | 975 | 242 | 24.8 | 220 | 22.6 | 434 | 44.5 | 79 | 8.1 |
| 16-20 | 796 | 220 | 27.6 | 222 | 27.9 | 274 | 34.4 | 80 | 10.1 |
| 21-25 | 251 | 47 | 18.7 | 53 | 21.1 | 109 | 43.4 | 42 | 16.7 |
| 26-30 | 280 | 75 | 26.7 | 80 | 28.6 | 79 | 28.2 | 46 | 16.4 |
| Total | 2,951 | 761 | 25.8 | 730 | 24.7 | 1,172 | 39.7 | 288 | 9.8 |

now suggest that the proportion of juveniles of both sexes remains fairly constant through April, whereas the proportion of adult males decreases. The important point, however, is that adult females do not join these moulting patches in appreciable numbers before the second half of April. The data suggest that adult females constitute no more than $15-20 \%$ of the catches in late April. Corresponding data from the Jan Mayen area and the Barents Sea indicate similar changes through April in these areas ( $\varnothing$ ritsland, unpub1ished).

Further evidence derived from subsamples for age analysis is presented in Table 5 and Fig. 7. These data indicate a decreasing trend in the mean age of males and a definite increase of the mean age of females through April. The mean age and sex ratio of an age sample collected in the moulting season therefore will depend upon the date of sampling. To be comparable, samples from different years should be collected at comparable dates. Characteristics of the present samples, partly derived from Table 4, are given below:

Table 4. Moulting harp seals in Front area, Newfoundland. Catch composition from age samples by age-group (1-6 years and 7 years + ), sex and date in 1967, 1969 and 1970. Because of inadequate dating, the 1967 subsamples marked with asterisks are omitted from the summary; they are, however, included in the total.

| Date in April | $\begin{gathered} \text { Sample } \\ \text { size } \end{gathered}$ | Juveniles |  |  |  | Adults |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | ${ }^{7}$ |  | 아 |  | $\mathrm{F}^{7}$ |  | + |  |
|  |  | No. | \% | No. | \% | No. | \% | No. | \% |
|  |  |  |  |  |  |  |  |  |  |
| 02-09* | 180 | 65 | 36.1 | 43 | 23.9 | 65 | 36.1 | 7 | 3.9 |
| 13-15 | 84 | 14 | (16.7) | 10 | (11.9) | 50 | (59.5) | 10 | (11.9) |
| 23-26* | 17 | 4 | - | 2 | ) | 7 | (59.5) | 4 | (11.9) |
| Subtotal | 281 | 83 | 29.5 | 55 | 19.6 | 122 | 43.4 | 21 | 7.5 |
| 1969 |  |  |  |  |  |  |  |  |  |
| 02-05 | 309 | 100 | 32.4 | 96 | 31.1 | 110 | 35.6 | 3 | 1.0 |
| 06-07 | 208 | 84 | 40.4 | 46 | 22.1 | 72 | 34.6 | 6 | 2.9 |
| 13 | 114 | 21 | (18.4) | 15 | (13.2) | 68 | (59.6) | 10 | (8.8) |
| 18-20 | 257 | 88 | 34.2 | 74 | 28.8 | 63 | 24.5 | 32 | 12.5 |
| 22 | 27 | 12 | - | 8 | - | 4 | - | 3 | - |
| Subtotal | 915 | 305 | 33.3 | 239 | 26.1 | 317 | 34.6 | 54 | 5.9 |
| 1970 |  |  |  |  |  |  |  |  |  |
| 19 | 48 | 15 | - | 19 | - | 14 | - | 0 | - |
| 28-29 | 260 | 90 | 34.6 | 61 | 23.5 | 71 | 27.3 | 38 | 14.6 |
| Subtotal | 308 | 105 | 34.1 | 80 | 26.0 | 85 | 27.6 | 38 | 12.3 |
| Summary, $1967+1969+1970$ |  |  |  |  |  |  |  |  |  |
| 01-05 | 309 | 100 | 32.4 | 96 | 31.1 | 110 | 35.6 | 3 | 1.0 |
| 06-10 | 208 | 84 | 40.4 | 46 | 22.1 | 72 | 34.6 | 6 | 2.9 |
| 11-15 | 198 | 35 | 17.7 | 25 | 12.6 | 118 | 59.6 | 20 | 10.1 |
| 16-20 | 305 | 103 | 33.8 | 93 | 30.5 | 77 | 25.2 | 32 | 10.5 |
| 21-25 | 27 | 12 | - | 8 | - | 4 | - | 3 | . |
| 26-30 | 260 | 90 | 34.6 | 61 | 23.5 | 71 | 27.3 | 38 | 14.6 |
| Total | 1,504 | 493 | 32.8 | 374 | 24.9 | 524 | 34.8 | 113 | 7.5 |


| Year | No. | Percentage by date in April |  |  | Mean age | \% $0^{\prime \prime}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | 1-10 | 11-20 | 21-30 |  |  |
| 1964 | 127 | 15.7 | 7.9 | 37.0 | 11.0 | 22.0 |
| 1967 | 281 | 64.1 | 29.9 | 6.0 | 8.5 | 72.9 |
| 1968 | 702 | (5) | (90) | (5) | 8.7 | - |
| 1969 | 915 | 56.5 | 40.5 | 3.0 | 6.7 | 67.9 |
| 1970 | 516 | 0.0 | 9.3 | 50.4 | 5.6 | 61.7 |

It is evident that no sample is strictly comparable to any of the others. It is assumed, however, that the 1967 and 1969 samples which both have a load in early April, can be compared. Tentatively, it may also be assumed that the 1968 sample, definitely loaded in mid-April, can be compared to the sample of 1970 which is loaded in late April. If this is accepted, it follows that the indicated decreasing trend of the mean age may be real.

Table 5. Moulting harp seals in Front area, Newfoundland. Mean age from age samples, by sex and date.

| $\begin{array}{\|c} \text { Date in } \\ \text { Apri1 } \end{array}$ | $0^{7}$ |  | 앙 |  |
| :---: | :---: | :---: | :---: | :---: |
|  | No. | $\begin{array}{r} \hline \text { Mean } \\ \text { age } \\ \hline \end{array}$ | No. | Mean age |
| 1967 |  |  |  |  |
| 02-09 | 130 | 8.4 | 50 | 3.1 |
| 13-15 | 64 | (12.3) | 20 | (8.2) |
| 22-26 | 11 | (12.3) | 6 | (8.2) |
| Sub total | 205 | 9.8 | 76 | 5.1 |
| 1969 |  |  |  |  |
| 02-05 | 210 | 8.0 | 99 | 1.7 |
| '06-07 | 156 | 8.2 | 52 | 3.1 |
| 13 | 89 | (10.4) | 25 | (4.8) |
| 18-20 | 151 | 7.0 | 106 | 5.1 |
| 22 | 16 | - | 11 | 5.1 |
| Subtotal | 622 | 8.0 | 293 | 3.6 |
| 1970 |  |  |  |  |
| 19 | 29 | - | 19 | - |
| 28-29 | 161 | 7.0 | 99 | 6.1 |
| Subtotal | 190 | 6.9 | 118 | 5.4 |

## Sex ratio

Counts of pups and juvenile seals were performed in 1967, 1969, 1970 and 1971. These data are shown in Table 6. A possible change with date in the sex ratio of pups is suggested by subsamples. The total ratios for pups and fuveniles are very close to each other, both showing that males are slightly more numerous than females (51.0\%).

Table 6. Sex ratio of harp seals in Front area, Newfoundland, 1967, 1969, 1970 and 1971.

| Dates | Males |  | Females |  | Sum |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | No. | \% | No. | \% |  |
| Pups |  |  |  |  |  |
| 11-23 March 1967 | 241 | 48.2 | 258 | 51.7 | 499 |
| 25 March-10 April 1969 | 234 | 51.2 | 223 | 48.8 | 457 |
| 26 March-17 April 1970 | 534 | 51.3 | 507 | 48.7 | 1,041 |
| 23 March-06 April 1971 | 105 | 55.9 | 83 | 44.1 | 188 |
| Subtotal | 1,114 | 51.0 | 1,071 | 49.0 | 2,185 |
| Juveniles |  |  |  |  |  |
| 12-23 Apri1 1967 | 38 | 52.8 | 34 | 47.2 | 72 |
| 02-22 April 1969 | 434 | 50.3 | 428 | 49.7 | 862 |
| 19 - 29 April 1970 | 107 | 49.5 | 109 | 50.5 | 216 |
| $10-13$ April 1971 | 182 | 53.4 | 159 | 46.6 | 341 |
| Subtotal | 761 | 51.0 | 730 | 49.0 | 1,491 |
| Total | 1,875 | 51.0 | 1,801 | 49.0 | 3,676 |

## Sexual maturity of females

Data on age at first ovulation from known-age breeding and moulting female harp seals were collected in 1964 and 1967. An additional sample of 173 breeding females was collected in 1971, but these specimens have not been processed yet. The data from analyses of ovaries collected in 1964 and 1967 are shown in Tables 7 and 8. This information indicates that female harp seals mature in age-groups from 4 to 12 or 13 years. The age of 50 percent maturity was found to be about 6.5 for breeding and about 5.5 for moulting females. However, the small sample from moulting seals is hardly representative of the maturing age-groups (see discussion of age composition above) and can only give supporting evidence on age at maturity, whereas samples collected in the breeding lairs apparently are representative of the mature females. When age at first ovulation can be established as suggested in Table 7, samples of breeding females therefore may yield direct evidence on age at maturity in the stock. The actual age distribution of breeding females supports the data from ovary analyses (Fig. 8). The time-lag between maturity and breeding (Parturition) reflects the one-year sexual cycle and the somewhat lower fertility of newly-matured females.

Table 7. Harp seals in Front area, Newfoundland. Sexual maturity of breeding females collected 12-20 March 1964 and 11-23 March 1967.

| Age at last ovulation | No. | Age at 'first' ovulation |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 |
| 4 | 0 | - | - | - | - | - | - | - | - | - | - | - |
| 5 | 2 | 1 | 1 | - | - | - | - | - | - | - | - | - |
| 6 | 11 | 2 | 3 | 6 | - | - | - | - | - | - | - | - |
| 7 | 7 | $\frac{1}{1}$ | $\underline{1}$ | 4 | 1 | - | - | - | - | - | - | - |
| 8 | 8 | - | $\underline{-}$ | 4 | 3 | 1 | - | - | - | - | _ | - |
| 9 | 2 | - | - | - | $\underline{1}$ | 0 | 1 | - | - | - | - | - |
| 10 | 6 | - | - | 1 | 1 | $\underline{3}$ | 1 | 0 | - | - | - | - |
| 11 | 3 | - | - | - | - | - | $\underline{2}$ | 1 | 0 | - | - | - |
| 12 | 8 | - | - | - | 2 | 3 | 0 | 2 | 1 | 0 | - | - |
| 13 | 9 | - | - | - | - | - | 1 | $\frac{2}{2}$ | 4 | 2 | 0 | - |
| 14 | 7 | - | - | - | - | - | 1 | 1 | $\frac{1}{2}$ | $\underline{0}$ | 2 | 1 |
|  | 63 |  |  |  |  |  |  |  |  |  |  |  |
| Sum | (48) | 3 | 5 | 14 | 5 | 4 | 4 | 3 | 5 | 2 | 2 | 1 |
| \% |  | 6.3 | 10.4 | 29.2 | 10.4 | 8.3 | 8.3 | 6.3 | 10.4 | 4.2 | 4.2 | 2.1 |
| Acc. \% |  | 6.3 | 16.7 | 45.9 | 56.3 | 64.4 | 72.9 | 79.2 | 89.6 | 93.8 | 98.0 | 100.1 |

Table 8. Harp seals in Front area, Newfoundland. Sexual maturity of moulting females collected 3-30 April 1964 and 14-26 April 1967.

| Age | No. | Age at 'first' ovulation |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 |
| 4 | 10 | 0 | - | - | - | - | - | - | - | - | - | - |
| 5 | 5 | 0 | 1 | - | - | - | - | - | - | - | - | - |
| 6 | 2 | 1 | 1 | 0 | - | - | - | - | _ | - | _ | - |
| 7 | 5 | $\underline{0}$ | 3 | 1 | 1 | - | - | - | - | - | - |  |
| 8 | 2 | - | 1 | 0 | 1 | 0 | - | - | - | - | - | - |
| 9 | 2 | - | 1 | 0 | 0 | 1 | 0 | - | - | - | - | - |
| 10 | 4 | - | - | $\overline{3}$ | 1 | 0 | 0 | 0 | - | - | - | - |
| 11 | 3 | - | - | 2 | 0 | $\underline{0}$ | 1 | 0 | 0 | - | - | - |
| 12 | 1 | - | - | - | - | - | 0 | 0 | 1 | 0 | - | - |
| 13 | 1 | - | - | - | 0 | 0 | $\overline{0}$ | 1 | 0 | 0 | 0 | - |
| 14 | 1 | - | - | - | - | - | - | $\frac{1}{1}$ | $\underline{0}$ | 0 | 0 | 0 |
|  | 36 |  |  |  |  |  |  |  |  |  |  |  |
| Sum | (15) | 1 | 6 | 1 | 3 | 1 | 1 | 1 | 1 | 0 | 0 | 0 |
| \% |  | 6.7 | 40.0 | 6.7 | 20.0 | 6.7 | 6.7 | 6.7 | 6.7 | 0.0 | 0.0 | 0.0 |
| Acc. \% |  | 6.7 | 46.7 | 53.4 | 73.4 | 80.1 | 86.8 | 93.5 | 100.2 | 100.2 | 100.2 | 100.2 |

## Fertility

Date on the rate of births or pregnancies are not available. Therefore, data from ovaries of breeding and moulting mature seals sampled in 1964 and 1967 were analyzed in order to get some information on fertility. Presuming that mature females ovulate once every year, that ovulations alternate regularly between the two ovaries and that successful matings result in coxpora lutea which are retained as visible corpora albicantia for at least three years after their formation, frequencies of corpora misaing in the regular sequence were recorded for the three most recent breeding seasons. If these missing corpora indicate missed pregnancies, the counts suggest an overall fertility of about $92 \%$ among mature harp seals. An increase from about $90 \%$ in age-groups 6-12 to about $93 \%$ in older age-groups is suggested. There is also some evidence for an increase in overall fertility of about $1 \%$ from 1964 to 1967. The counts of corpora are shown in Table 9.

## Pup catches and survival of year-classes

Tentative indexes of survival of year-classes based on relative frequencies of age-groups in samples from the Front, in relation to corresponding frequencies derived from data in a life table (Table 10) constructed by Sergeant (MS 1960) are given in Table 11. They are based on all available Norwegian age samples, and calculated by dividing relative frequencies in age samples by corresponding relative frequencies in column "Young excl." in Table 10. Weighted means have been calculated for each year-class, using factors roughly corresponding to the size of each age sample (1964:1, 1967:2, 1968:4, 1969:5 and 1970:3).

These indexes are offered as an attempt to quantify earlier rather subjective guesstimates of year-class survival for harp seals. Bearing in mind

Table 9. Fertility of sexually mature female harp seals in Front area, Newfoundland. Counts of corpora lutea and albicantia, and estimates of missing corpora for the three most recent breeding seasons (ovulations), presuming that ovulations alternate regularly between ovaries.

| $\begin{aligned} & \text { Age } \\ & \text { groups } \end{aligned}$ | $\begin{gathered} \text { Year } \\ \text { of } \\ \text { coll. } \end{gathered}$ | Breeding females |  |  | Moulting females |  |  | Total |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | No. count. | $\begin{gathered} \text { No. } \\ \text { miss. } \end{gathered}$ | $\begin{gathered} \% \\ \text { miss. } \\ \hline \end{gathered}$ | No. count. | $\begin{gathered} \text { No. } \\ \text { miss. } \end{gathered}$ | $\begin{gathered} \% \\ \text { miss. } \end{gathered}$ | No. count. | $\begin{aligned} & \text { No. } \\ & \text { miss. } \end{aligned}$ | $\begin{gathered} \% \\ \text { miss. } \end{gathered}$ |
| 6-12 | $\begin{aligned} & 1964 \\ & 1967 \end{aligned}$ | $\begin{array}{r} 8 \\ 68 \end{array}$ | $\begin{aligned} & 1 \\ & 8 \end{aligned}$ |  | $\begin{aligned} & 36 \\ & 17 \end{aligned}$ | 3 1 |  | 124 | 13 | 9.5 |
| 13-18 | 1964 | $\begin{aligned} & 35 \\ & 91 \end{aligned}$ | $\begin{aligned} & 4 \\ & 4 \end{aligned}$ |  | $\begin{aligned} & 23 \\ & 16 \end{aligned}$ | $\begin{aligned} & 1 \\ & 2 \end{aligned}$ |  | 165 | 11 | 6.3 |
| 19 up | $\begin{aligned} & 1964 \\ & 1967 \end{aligned}$ | $\begin{aligned} & 38 \\ & 80 \end{aligned}$ | $\begin{aligned} & 2 \\ & 7 \end{aligned}$ |  | $\begin{aligned} & 8 \\ & 9 \end{aligned}$ | $\begin{aligned} & 1 \\ & 0 \end{aligned}$ |  | 135 | 10 | 6.9 |
| Unknown | 1964 | 23 | 3 |  | 9 | 2 |  | 32 | 5 |  |
| A11 | $\begin{aligned} & 1964 \\ & 1967 \end{aligned}$ | $\begin{aligned} & 104 \\ & 239 \end{aligned}$ | 10 19 | $\begin{aligned} & 8.8 \\ & 7.4 \end{aligned}$ | $\begin{aligned} & 76 \\ & 37 \end{aligned}$ | 7 3 | $\begin{aligned} & 8.4 \\ & 7.5 \end{aligned}$ | 456 | 39 | 7.9 |

Table 10. Relative age-group frequencies of harp seals of the Northwest Atlantic derived from life table by Sergeant (MS 1960).

| Age | Percent |  |
| :---: | :---: | :---: |
|  | Young incl. | Young excl. |
| 0 | 18.6 | - |
| 1 | 11.4 | 14.0 |
| 2 | 9.0 | 11.0 |
| 3 | 7.4 | 9.1 |
| 4 | 6.3 | 7.7 |
| 5 | 5.4 | 6.6 |
| 6 | 4.9 | 6.0 |
| 7 | 4.4 | 5.4 |
| 8 | 4.0 | 4.9 |
| 9 | 3.6 | 4.4 |
| 10 | 3.2 | 3.9 |
| 11 | 2.9 | 3.6 |
| 12 | 2.6 | 3.2 |
| 13 | 2.3 | 2.8 |
| 14 | 2.1 | 2.6 |
| 15 | 1.9 | 2.3 |
| 16 | 1.7 | 2.1 |
| 17 | 1.5 | 1.8 |
| 18 | 1.2 | 1.5 |
| 19 | 1.0 | 1.3 |
| 20 | 0.9 | 1.1 |
| 21 | 0.8 | 0.9 |
| 22 | 0.6 | 0.7 |
| 23 | 0.5 | 0.6 |
| 24 | 0.4 | 0.5 |
| 25 | 0.4 | 0.5 |
| 26 | 0.3 | 0.4 |
| 27+ | 0.8 | 0.9 |
| Sum | 100.1 | 99.8 |

Table 11. Harp seal pup catches at Newfoundland and survival of corresponding year-classes expressed by relative frequency in age samples from the Front, Newfoundland, divided by relative frequency of age-groups in life table (Sergeant, MS 1960). Parentheses indicate estimated catches and size of age samples.

| Year- <br> class | Catch <br> of pups <br> Front <br> $10^{-3}$ | Total catch of pups$10^{-3}$ | "Survival Index" from age samples |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | $\begin{gathered} 1 \\ 1964 \\ (127) \end{gathered}$ | $\begin{gathered} 2 \\ 1967 \\ (281) \end{gathered}$ | $\begin{gathered} 4 \\ 1968 \\ (702) \end{gathered}$ | $\begin{gathered} 5 \\ 1969 \\ (915) \end{gathered}$ | $\begin{gathered} 3 \\ 1970 \\ (516) \end{gathered}$ | Weighted mean |
| 1945 | - | - | 2.38 | 0.57 | 1.83 | 0.60 | 0.40 | 1.00 |
| 46 | - | 73 | 4.20 | 1.56 | 3.43 | 1.33 | 0.00 | 1.85 |
| 47 | - | 102 | 2.17 | 1.27 | 1.55 | 0.71 | 0.67 | 1.10 |
| 48 | - | 137 | 1.48 | 2.46 | 2.09 | 1.22 | 0.57 | 1.50 |
| 49 | (169) | 227 | 1.70 | 2.13 | 2.31 | 1.62 | 1.11 | 1.78 |
| 1950 | (174) | 226 | 0.92 | 1.56 | 2.47 | 1.62 | 0.73 | 1.61 |
| 51 | (215) | 319 | 0.86 | 1.00 | 1.17 | 1.20 | 0.92 | 1.09 |
| 52 | (126) | 198 | 0.97 | 1.57 | 0.81 | 1.50 | 0.80 | 1.15 |
| 53 | (161) | 198 | 0.86 | 1.38 | 0.83 | 0.62 | 0.33 | 0.74 |
| 54 | (109) | 175 | 1.62 | 0.64 | 1.00 | 1.00 | 1.00 | 0.99 |
| 1955 | (159) | 252 | 0.89 | 0.66 | 0.75 | 0.88 | 0.70 | 0.78 |
| 56 | (245) | 341 | 0.80 | 1.28 | 0.53 | 0.75 | 0.81 | 0.78 |
| 57 | (71) | 165 | 0.87 | 0.72 | 0.86 | 0.56 | 0.43 | 0.66 |
| 58 | (56) | 141 | 1.57 | 0.41 | 1.03 | 0.69 | 0.44 | 0.75 |
| 59 | (181) | 239 | 0.89 | 0.94 | 1.00 | 0.82 | 0.69 | 0.86 |
| 1960 | (81) | 171 | 1.13 | 1.26 | 1.10 | 0.86 | 1.10 | 1.04 |
| 61 | 126 | 179 | 0.69 | 1.30 | 1.22 | 0.98 | 0.66 | 1.00 |
| 62 | (114) | 214 | 0.43 | 1.03 | 0.47 | 0.72 | 1.02 | 0.74 |
| 63 | (159) | 278 | 0.17 | 0.27 | 0.65 | 0.52 | 0.65 | 0.52 |
| 64 | (168) | 273 | - | 0.20 | 0.49 | 0.65 | 0.48 | 0.50 |
| 1965 | (90) | 190 | - | 1.00 | 0.66 | 0.43 | 0.85 | 0.67 |
| 66 | (167) | 257 | - | 1.40 | 0.95 | 0.41 | 0.75 | 0.78 |
| 67 | 185 | 280 | - | - | 1.37 | 0.42 | 0.70 | 0.81 |
| 68 | 97 | 156 | - | - | - | 2.90 | 1.92 | 2.53 |
| 69 | 186 | (238) | - | - | - | - | 1.83 | (1.83) |

that age samples hardly are representative of the stock, it is worth attention that the "survival index" is rather consistently high or low through several or all age samples for some year-classes.

Revised sealing statistics for the ICNAF area up to 1968 have been published by ICNAF Headquarters (1970) and statistics for 1969 are available in manuscript (ICNAF, MS 1970). Provisional data on Norwegian sealing in 1970 also have been reported ( $\emptyset$ ritsland, MS 1970). The published Canadian statistics (ICNAF, 1970) are somewhat lacking in detail up to 1965, but Sergeant (MS 1967) has given some estimates of division between subareas. Csnadian data for 1970 are not available. Some information on Danish (Greenland) and Soviet catches in the ICNAF area are included in the published statistics (ICNAF, 1970), but occasional French and U.S. catches have not been reported.

By and large, the available statistics seem to give a fairly good coverage for the years from 1947 to 1969, although somewhat lacking in detail before 1965.

In Table 11 the "survival indexes" are compared to total pup catches in the Front area and in the ICNAF area. No definite correlation has been discovered between survival index and estimated pup catch on the Front. Weighted index means in relation to total pup catches are plotted in Fig. 9. Plots for the years 1946-1959 show no obvious trend, but plotted values for 1960-1968 suggest that total pup catches higher than 170-180 thousand may have had an adverse effect on the survival of year-classes in these years.

Catch in relation to hunting activity
An attempt to calculate catch per unit of hunting activity for Norwegian sealing at Newfoundland is presented in Fig. 10, showing total Norwegian catches of harp seals (all age-groups) on the Front per unit of hunting activity for the years 1946-1970. The estimates are based on the formula

C/ $\mathrm{n}_{\mathrm{T}}^{\mathrm{P}}$
where $C$ is catch in number of animals,
$n$ is number of ships, and
$\frac{P}{T}$ is an average efficiency index:brake horse power per gross ton
The data suggest that the availability of harp seals may have been reduced by some 60 percent during the years from 1955 to 1970 . Catches per unit of activity are plotted against units of activity in Fig. 11.

Accumulated catch of year-classes
Estimated total catches by 1970 of some year-classes which were selected because pup catches in these years were exceptionally high, are shown in Table 12. To the actual catches of pups have been added the estimated catches represented by each year-class in each year. These estimates are based on actual catches of one-year-old or older seals, and "normal" relative age-group frequencies under the heading "Young excl." in Table 10.

Although the estimated catches of one-year-old or older animals should be recalculated from actual age-group frequencies in age samples, the accumulated catches may serve as a useful reminder. According to these estimates even the year-class produced in 1967 may have yielded more than 295 thousand seals by the end of the season 1970. For this year-class, however, the accumulated catch may be corrected, using actual age-group frequencies in Norwegian samples from 1968, 1969 and 1970:

| Year | Percent in <br> sample | Catch of one <br> year or older | Catch of 1967 <br> year-class |
| :--- | :---: | :---: | :---: |
|  | - | - | 280,442 |
| 1967 | 19.2 | 36,238 | 6,958 |
| 1969 | 4.6 | 55,472 | 2,552 |
| 1970 | 6.4 | $(33,403)$ | $-(2,138)$ |
|  |  | Accumulated catch | 292,090 |

From this it may be assumed that pup production in 1967 was at least 292 thousand.

Table 12. Total catches of selected year-classes of harp seals in the Northwest Atlantic.

| Year | Year-class |  |  |  |
| :---: | ---: | ---: | ---: | ---: |
|  | 1951 | 1956 | 1963 | 1967 |
| 1951 | 318,626 | - | - | - |
| 52 | 15,266 | - | - | - |
| 53 | 8,240 | - | - | - |
| 54 | 8,134 | - | - | - |
| 1955 | 6,243 | - | - | - |
| 56 | 3,169 | 341,397 | - | - |
| 57 | 4,803 | 11,206 | - | - |
| 58 | 8,467 | 17,247 | - | - |
| 59 | 3,984 | 7,398 | - | - |
| 1960 | 5,413 | 9,473 | - | - |
| 61 | 797 | 1,349 | -105 | 6,841 |
| 62 | 2,358 | 3,980 | 278,350 | - |
| 63 | 2,171 | 3,799 | 10,855 | - |
| 64 | 1,396 | 2,363 | 5,907 | - |
| 1965 | 1,703 | 2,888 | 6,738 | - |
| 66 | 1,221 | 2,093 | 4,476 | 280,442 |
| 67 | 652 | 1,160 | 2,392 | 5,073 |
| 68 | 834 | 1,556 | 3,334 | 6,113 |
| 69 | 650 | 1,300 | 2,700 | 4,550 |
| 1970 |  |  |  |  |
| Accumulated | 398,232 | 414,050 | 314,752 | 296,178 |
| catch |  |  |  | - |

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Fig. 1.


Fig. 2.


Fig. 3.


Fig. 4.


Fig. 5.


Fig. 6. Age and sex composition of catches of moulting Harp seals in Front area, Newfoundland, through April.
A) From age samples (Tab. 4), 0 1967, +1969 , o 1970.
B) From counts of pelts (Tab. 3).


Fig. 7. Mean age of males and females in catches of moulting Harp seals in Front area, Newfoundland, through April (Tab. 5). - 1967, + 1969, o 1970.




16. Distribution and abundance of yellowtail flounder populations on Georges Bank and off southern New England according to trawl survey data
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## Introduction

In 1970 the absolute abundance of yellowtail flounder populations on Georges Bank and in southern New England was calculated by Brown (1970). He described the abundance of the 1958-1961 year-classes at the age of 2 years and older estimated by Gulland's model (1963) and also gave its evaluation for 1968 on the basis of the catch data. This paper calculates the abundance of two yellowtall stocks in 1967-1970 from trawl survey data.

## Materials and methods

This study is based upon material obtained during the joint USSR-US trawl surveys conducted in the autumn of 1967-1970. Charts of distribution and relative abundance were constructed as follows: the data on a trawl survey were plotted on plane-tables and then interpolated, using zones of species density of $1-10$ specimens, 11-25 specimens, $26-100$ specimens, $101-200$ specimens, 201 and more specimens per 30 -minute haul and treating each separately. The areas inhabited by the species with a certain density were determined with the help of a planimeter.

Mean individual weights by sample sizes for the populations were calculated by the formula:

$$
P=\frac{\sum n 1 P 1}{N}
$$

```
where \(\quad P=\) mean individual weights by sample sizes,
nl \(=\) number of specimens of a certain size,
Pl = mean individual weight by fish sizes,
    \(N=\) total number of specimens in the size frequency.
```

Since the populations from Georges Bank and southern New England come in contact in nature, calculation of the absolute abundance is made along the $70^{\circ}$ meridian which is used as a conventional boundary between the above populations (Fig. 1). We have used the following formula:

No. $=\frac{1}{K} \sum \frac{S_{i} \times n_{1}}{S}$
where No. $\ddagger$ the abundance of the population in numbers, $S_{1}=$ the area of a certain density, $S^{1}=$ the tow area ( 0.016 square miles), $n_{1}=$ average abundance within the layer of a certain density, $k^{1}=$ general catchability coefficient by Edwards (1968) which is 0.39.

[^15]According to Scott (1954) there are three populations of yellowtail flounder in the area off New England: (1) the population inhabiting the Cape Cod area, (2) the Georges Bank population, and (3) the population off southern New England. Later, tagging experiments and morphometric studies by Lux (1963) confirmed this suggestion. The present paper deals with only two populations: one from southern New England and the other from Georges Bank.

The distribution of yellowtail flounder on Georges Bank is given in Figs. 3, 4, and 5. These figures show that no substantial changes in habitat were observed during the period of 1968-1970. However, some change of areas of different density took place within the habitat, e.g. In 1968 concentrations of yellowtail flounder with a density of $101-200$ specimens per $30-\mathrm{min}$ haul were observed over an area of 0.46 thousand square miles, the area decreased to 0.08 thousand square miles before 1970. The tendency to decrease is also observed over the areas with a density of $26-100$ specimens; respectively, the areas with the least density of yellowtail flounder tended to increase (Table 1).

Table 1. Areas (in thousands of square miles) occupled by various densities of yellowtail flounder in the southern New England and Georges Bank populations, 1967-1970.

| Year | Areas of fish densities |  |  |  |  | Total area |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1-10 | 11-25 | 26-100 | 101-200 | 201-max |  |
|  | Southern New England |  |  |  |  |  |
| 1967 | 3.70 | 3.05 | 4.24 | 2.53 | 0.08 | 13.60 |
| 1968 | 3.91 | 2.24 | 4.96 | 2.41 | 0.51 | 14.03 |
| 1969 | 5.96 | 3.56 | 2.52 | 1.83 | 1.15 | 15.02 |
| 1970 | 5.18 | 2.36 | 2.98 | 0.95 | 0.49 | 11.96 |
| Average | 4.96 | 2.80 | 3.65 | 1.93 | 0.56 | 13.63 |
|  | Georges Bank |  |  |  |  |  |
| 1968 | 3.79 | 2.49 | 3.61 | 0.46 | - | 10.35 |
| 1969 | 5.53 | 2.93 | 3.19 | 0.16 | - | 11.81 |
| 1970 | 7.07 | 2.07 | 2.09 | 0.08 | - | 11.31 |
| Average | 5.46 | 2.50 | 2.96 | 0.23 | - | 11.15 |

In southern New England (Figs. 2, 3, 4, and 5) an increase of the habitat areas due to the southerly extension of yellowtall flounder was noted up to 1970, e.g. In 1967 this species occurred over an area of 13.60 thousand square miles, while in 1968 it occupied 14.03 thousand square miles, and in 1969 15.02 thousand square miles (Table 1). In 1970 the habitat areas decreased abruptly and constituted only 11.96 thousand square miles. It should be noted that up to 1970, the areas with frequency of occurrence of 201 specimens and more increased substantially. In 1967 they occupied 0.08 thousand square miles, in 1968-0.51 thousand square miles, in 1969-1.15 thousand square miles, while in 1970 these areas decreased somewhat and were only 0.49 thousand square miles $(0.6 \%, 4.3 \%, 7.6 \%$ and $4.0 \%$ of the total habitat area, respectively).

## Size composition

The results of mass measurements of yellowtail flounder are given in figs. 6 and 7. These figures show that the areas under study undergo some changes in size composition of the catches. A decrease in mean length is observed on Georges Bank. The population is characterized by an increase of the young size-groups and a decrease of the older ones. The reverse is recorded in southern New England where the mean length of yellowtall flounder increases: the proportion of the young size-groups decreases and that of the older ones increases from year to year. The percentage recruitment of $12-28-\mathrm{cm}$ specimens decreases in both populations. This can be explained by a lower abundance of 1967, 1968 and 1969 year-classes as pointed out earlier by Brown (1970). The decrease in mean size of yellowtail flounder from Georges Bank results in a decrease in mean weight. Mean weight of yellowtail flounder in southern New England increases (Table 2).

Table 2. Mean weights (g) of yellowtail flounder populations in southern New England and Georges Bank, by sample sizes based on the trawl surveys, 1967-70.

| Year | Southern New England | Georges Bank |
| :---: | :---: | :---: |
| 1967 | 190 | - |
| 1968 | 211 | 304 |
| 1969 | 224 | 287 |
| 1970 | 253 | 291 |

## Studies of the absolute abundance of the populations

The absolute abundance of the two populations was evaluated on the basis of the frequency of occurrence and the general catchability coefficient. In 1968-1970 the average absolute abundance on Georges Bank was $54.5 \times 10^{6}$ specimens, but the abundance decreases from year to year (Table 3). In 1968 it was $66.8 \times 10^{6}$ specimens, but before 1970 it decreased to $40.2 \times 10^{6}$ specimens. In the period 1967-1970 the average absolute abundance in southern New England was $140.8 \times 10^{6}$ specimens. Highest abundance was observed in 1968 ( $167.6 \times 10^{6}$ specimens). That year also had the highest catches per unit of effort (Graham, 1969). Later, the abundance began to decrease and by 1970, was $100.0 \times 10^{6}$ specimens (Table 3). After estimating mean individual weights of yellowtail flounder by sample sizes, and its absolute abundance, we calculated the biomass of the two populations (Table 4). Biotmass in autumn 1968 was 55.7 thousand tons, in 1969-52.1 thousand tons, and in $1970-37.0$ thousand tons. Thus, in 1971 the hiomass of these two populations will be lower than in the previous years.

## Conclusion

The following conclusion can be drawn from these investigations.
The habitat of the Georges Bank yellowtail flounder population showed no changes during the period of investigation, but showed some tendency toward a decrease in areas in which the maximum fish density was observed within the habitat.

Table 3. Absolute abundance (in pieces $x 10^{6}$ ) of the southern New England and Georges Bank populations by densities.

| Year | Numbers of fish with densities |  |  |  |  | Total area |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1-10 | 11-25 | 26-100 | 101-200 | 201-max |  |
|  | Southern New England |  |  |  |  |  |
| 1967 | 3.7 | 10.1 | 47.8 | 69.8 | 4.4 | 135.8 |
| 1968 | 3.9 | 7.4 | 57.3 | 66.3 | 32.7 | 167.6 |
| 1969 | 6.0 | 11.7 | 29.0 | 50.5 | 62.9 | 160.1 |
| 1970 | 5.2 | 7.8 | 34.4 | 26.1 | 26.5 | 100.0 |
| Average | 4.7 | 9.2 | 42.1 | 53.2 | 31.6 | 140.8 |
|  |  |  | George |  |  |  |
| 1968 | 3.8 | 8.2 | 42.1 | 12.7 | - | 66.8 |
| 1969 | 5.6 | 9.7 | 36.8 | 4.3 | - | 56.4 |
| 1970 | 7.1 | 6.9 | 24.1 | 2.1 | - | 40.2 |
| Average | 5.5 | 8.3 | 34.3 | 6.4 | - | 54.5 |

Table 4. Biomass (in thousand tons) of the southern New England and Georges Bank yellowtail flounder populations, 1967-70.

| Year | Southern New England | Georges Bank |
| :---: | :---: | :---: |
| 1967 | 25.8 | - |
| 1968 | 35.4 | 20.3 |
| 1969 | 35.9 | 16.2 |
| 1970 | 25.3 | 11.7 |
| Average | 30.6 | 16.1 |

The boundaries of the southern New England population expanded to 1970, the habitat areas increased (by 1.5 thousand square miles) from 13.51 thousand square miles in 1967 to 15.01 thousand square miles in 1969; in 1970 the area decreased sharply to 11.95 thousand square miles.

Mean length of yellowtail flounder from Georges Bank decreased and in southern New England tended to increase.

The absolute abundance of the Georges Bank population in 1968 was 66.8 x $10^{6}$ specimens, in $1969-56.3 \times 10^{6}$ specimens, and in $1970-40.2 \times 10^{6}$ specimens. It decreased in 1970 by $26.6 \times 10^{6}$ specimens ( $40 \%$ ), as compared with 1968. Average biomass was about 16.1 thousand tons.

In southern New England the maximum abundance was recorded in 1968 $167.6 \times 10^{6}$ specimens, the minimum abundance in 1970 ( $100.0 \times 10^{6}$ specimens).

The average biomass during the period in studies was 30.6 thousand tons.

## Discussion

Our results differ slightly from the abundance data given by Brown (1970) who estimated abundance at $105,297 \times 10^{3}$ specimens based on catch data, and the yield at 32.6 thousand tons with an $F=0.8$, or 31.2 thousand tons with an $F=1.15$. Richter and Vinogradov (1969) determined the absolute abundance of the Georges Bank population at 11.148 thousand tons from 1968 trawl survey data.

The data on the absolute abundance and biomass of the populations from Georges Bank and southern New England obtained by using the general catchability coefficient of 0.39 proposed by Edwards, reflecting the state of the populations and the tendency of the abundance and biomass to decrease, are doubtful as absolute; the biomass is practically equal to the catches. The general catchability coefficient is, evidently, somewhat overestimated and, hence, the values of the absolute abundance and biomass should be considered as relative.

## Summary

1. The abundance of the Georges Bank population is considerably lower than that of the population in southern New England.
2. In recent years, these two populations have tended to decrease in abundance.

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Fig. 5. Distribution and abundance of the Georges Bank and southern New England yellowtail flounder populations in 1970.


Fig. 6. Length frequency of Georges Bank yellowtail flounder for 1968-70.


Fig. 7. Length frequency of southern New England yellowtail flounder for 1967-70.

## 17. Prediction of yellowtail flounder population size from pre-recruit catches

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## Introduction

Research vessel survey cruises (Grosslein, 1969) conducted each autumn since 1963 provide an estimate of relative abundance of pre-recruit yellowtail flounder. These values may be used to predict population size in following years assuming certain rates of mortality. The relationship between this prediction and actual catch can be utilized to establish a catch quota one year in advance that would provide for the desired fishing rate. This procedure has been used to estimate a 1972 quota for yellowtail flounder in the southern New England area - west of $69^{\circ}$ and south of $41^{\circ} 15^{\prime}$ within Subarea 5.

## Methods

It was possible to identify the fish of age-group I with reasonable accuracy in the length frequencies of the autum (October-November) Albatross IV survey cruises. These are the youngest fish which are consistently caught in sufficient quantity to provide an estimate of abundance with reasonable precision. The stratified mean numbers of yellowtail flounder in age-group I caught per tow in the autumn survey cruises was estimated for the southern New England strata (Fig. 1). These estimates (Table 1) were taken to represent the relative abundance of age-group II at the beginning of the year following that in which the cruise was made.

Table 1. Number of age I yellowtail flounder caught per tow in autumn survey cruises in the southern New England strata (see Fig. 1).

| Year of <br> cruise | 1963 | 1964 | 1965 | 1966 | 1967 | 1968 | 1969 | 1970 |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Number <br> per tow | 16.3 | 18.5 | 11.7 | 34.4 | 19.9 | 9.0 | 7.0 | 8.3 |

At present yellowtail flounder first enter the US commercial fishery in significant numbers at age II. Fish are almost completely recruited by age III. Most of the comercial catch is represented by age-groups II through $V$ (Brown and Hennemuth, 1971).

$$
\begin{aligned}
\text { Let } n_{i, j} & =\text { relative abundance of age-group } j \text { in year } i, \\
\text { where } \quad j & =0,3 ; i=1,2, \ldots \ldots
\end{aligned}
$$

For our data, $n_{1,0}$ represents the relative abundance of age-group II at the ${ }^{1}$ Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/115.
beginning of 1964, the initial observation. The $n_{1,0}$ are observed, and the $n_{i, j}, j=1,3$ are calculated.

The relative abundance of the available total population at the beginning of any year is
$N_{i}=n_{i, 0}+e^{-Z_{1}}\left(n_{1-1,0}+n_{1-2,0} e^{-Z_{2}}+n_{i-3,0} e^{-2 Z_{2}}\right)$.
$\mathrm{Z}_{1}=$ total mortality during the first year in the fishery (age-group II).
$\mathrm{Z}_{2}=$ total mortality during the second, third or fourth years in the fishery (age-groups III-V).

The numbers caught,
$C(N)_{i}=n_{i, 0}\left(1-e^{-Z}\right) E_{1}+\left(1-e^{-Z_{2}}\right) E_{2} e^{-Z_{1}}\left(n_{i-1,0}+n_{i-2,0} e^{-Z_{2}}+n_{i-3,0} e^{-2 Z_{2}}\right)$. (2)
$E=F / Z$, the percent of total mortality due to fishing.
The same indices in terms of weight, $W_{i}$, and $C(W)$, may be obtained by multiplying the $n_{i, j}$ in (1) by the mean weight per fish, adding the instantaneous growth rate, $G, 1, j$ in the exponential of the catch equation (2), viz. $e^{-Z+G}$, and expressing $E$ as $\frac{F}{Z-G}$. The mean weight per fish at age was calculated from the growth equation estimated by Lux and Nichy (1969) and the length weight equation estimated by Lux (1969).

Results
The survey population index, $N_{i}$, and the survey catch index, $C(N)$, are tabulated in Table 2 and plotted in Fig. 2 (top line). The average fishing rate, $\Sigma C(N)_{i} / \Sigma N_{i}$, was 40 percent.

Table 2. Estimated population and catch indices for two fishing intensities, and the observed catch in numbers of fish.

| Year | Survey Population ${ }^{1}$ Index N | Survey <br> Catch ${ }^{1}$ <br> Index <br> C(N) | $\begin{gathered} \text { Observed } \\ \text { Catch } \\ \text { Nos. } \times \quad 10^{-6} \end{gathered}$ |
| :---: | :---: | :---: | :---: |
| 1967 | 45.8 | 14.6 | 77 |
| 1968 | 43.9 | 18.6 | 78 |
| 1969 | 28.0 | 13.0 | 87 |
| 1970 | 17.9 | 7.9 | 57 |
| 1971 | 15.2 | 5.8 | - |

${ }^{1} Z_{1}=0.5 ; \quad Z_{2}=1.2 ; \quad M=0.2$
The mortality rates given in Table 2 were derived from an analysis of the

US commercial fishery statistics for the years 1960-1968 (Brown and Hennemuth, 1971). This study also indicated that a reduction in fishing mortality (F) of at least $20 \%$ would be required to maximize sustainable yield. A $20 \%$ reduction in F corresponds to an average fishing rate, as defined above, of 35\% (bottom line in Fig. 2).

The observed commercial catch in numbers of fish plotted against the survey catch index (Fig. 3) contains too few points to verify the accuracy of precision of the relationships. In comparison to Fig. 2 it does appear that the catches during 1969 and 1970 did produce higher fishing intensities than in 1967-1968, assuming that the survey indices are valid.

To be useful in determining the quota for 1972, the relation between the catch and the population index in terms of weight and also the age II index for 1972 are required. The latter is, of course, not available until autumn 1971, and we have used instead the average of the age II index for 1969-1971, which equals 8.1.

The indication, in Fig. 3, of an increased fishing rate in 1969 and 1970 requires an upward adjustment of the fishing mortality rate. The required increases were taken to be proportional to the increased fishing intensity, and the adjusted rates are listed in Table 3, along with the estimated population indices for each age-group. The resulting survey population weight index, $W_{1}$, and the observed catches are plotted in Fig. 4.

Table 3. Weight indices of yellowtail flounder abundance by age-group.

| Year-class | Age |  |  |  |
| :--- | ---: | :---: | :---: | :---: |
|  | 2 | 3 | 4 | 5 |
| 1962 | 25.9 | 32.0 | 15.3 | 6.1 |
| 1963 | 29.4 | 36.3 | 17.3 | 6.8 |
| 1964 | 18.6 | 23.0 | 10.7 | 4.1 |
| 1965 | 54.7 | 67.7 | 32.1 | 7.4 |
| 1966 | 31.6 | 39.2 | 15.3 | 4.8 |
| 1967 | 14.3 | 15.9 | 6.1 | 2.7 |
| 1968 | 11.1 | 12.3 | 5.6 | 2.0 |
| 1969 | 13.2 | 16.2 | 7.6 | 2.7 |
| $1970^{1}$ | 12.9 | - | - | - |

T Average of 1967-69 year-classes
$\mathrm{M}=.2$
Age 2, $F=.3,1964-68,1971$
Age 2, $F=.4,1969,1970$
Age 3, $F=1.0,1964-68,1971-74$
Age 3, $F=1.2,1969,1970$
The points for 1967 and 1968 determine a line of yield corresponding to a fishing rate believed to be moderately higher than that corresponding to sustained yield under the conditions of high average recruitment which occurred during the period 1960-1968. The line through the 1969 and 1970 points indicates over-fishing. The point for 1971, assuming the entire catch quota of 13,000 metric tons is taken from the southern New England stock, falls between the two lines. Even if 2,000 metric tons of the quota is taken from the Cape

Fig. 1. Southern New England survey strata.


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Fig. 3. Actual catch in numbers versus catch index.


Fig. 4. Population weight index versus catch in weight.
18. Comparisons of abundance indices from research vessel surveys and commercial statistics for cod and haddock in ICNAF Subarea $4^{15}$
by

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## Introduction

On recommending that an ICNAF Working Group on Coordinated Groundfish Surveys be established, the Standing Committee on Research and Statistics emphasized that determination of accuracy of abundance indices derived from research vessel surveys should be one of its first tasks. Obviously, future survey planning decisions should be based on a realistic assessment of the ability of surveys to provide the type of information required.

To this end, a number of comparisons of survey and commercial abundance indices for cod and haddock in ICNAF Subarea 4 are brought together here. Complete analysis has not been possible on these data because of time limitations and this report should be considered preliminary.
Div. 4X cod

The U.S. National Marine Fisheries Service has surveyed most of ICNAF Div. 4X at least once a year since 1965. These surveys by the research vessel $A l b a t r o s s ~ I V$, using a stratified random sampling design, extended as far as the mouth of the Bay of Fundy and included LaHave and Roseway Banks. Stratified mean catch of cod in pounds per tow, calculated after log transformation of the data, is used here as an index of cod abundance (Table 1). Pre-recruits are included in this index but they make a very smali contribution to the survey index on a total weight basis because of their relative unavailability to the survey trawl. The strata considered (numbers 31-35, 41, 42 - see Lux and Grosslein, MS 1970) cover most of Div. 4X but exclude the inner Bay of Fundy, inshore southwestern Nova Scotia, and depths greater than 100 fathoms. These indices are, therefore, estimates of abundance of the offshore cod stock in Div. 4X.

Halliday (MS 1971) has taken the catch per hour of Canadian side otter trawlers of 151-500 gross tons which caught predominantly cod in "unit areas" $\mathrm{N}, \mathrm{O}, \mathrm{P}$ of $\mathrm{Div}$.4 X as being the best available commercial index of abundance of the offshore cod stock (Table 1). This area includes deep waters of the Fundian Channel and eastern Gulf of Maine while excluding the region between northern Brown's Bank and the Bay of Fundy. Thus, the areas from which survey and commercial data come do not coincide exactly. However, both include the

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major areas of cod concentration on Brown's and LaHave Banks.

Table 1. ICNAF Div. 4X cod abundance indices from USA research vessel Albatross IV and Canadian commercial otter trawlers.

| Year | Quarter | Season | $\begin{aligned} & \text { Survey Index } \\ & \log _{\mathrm{e}}(1 \mathrm{~b} / \text { tow }) \end{aligned}$ | Commercial index metric tons/hour |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | Quarterly | Annual |
| 1965 | 1 | Winter | - | . 307 |  |
|  | 2 | Spring | - | . 355 |  |
|  | 3 | Summer | 2.66 | . 376 |  |
|  | 4 | Fall | 3.25 | . 333 |  |
|  |  |  |  |  | . 343 |
| 1966 | 1 | Winter | - | . 362 |  |
|  | 2 | Spring | 3.28 | . 433 |  |
|  | 3 | Summer | - | . 263 |  |
|  | 4 | Fall | 2.71 | . 225 |  |
|  |  |  |  |  | . 321 |
| 1967 | 1 | Winter | - | . 362 |  |
|  | 2 | Spring | - | . 381 |  |
|  | 3 | Summer | - | . 273 |  |
|  | 4 | Fall | 2.16 | . 265 |  |
|  |  |  |  |  | . 320 |
| 1968 | 1 | Winter | - | . 332 |  |
|  | 2 | Spring | 2.47 | . 317 |  |
|  | 3 | Summer |  | . 270 |  |
|  | 4 | Fall | 1.86 | . 240 |  |
|  |  |  |  |  | . 290 |
| 1969 | 1 | Winter | - | . 302 |  |
|  | 2 | Spring | 1.86 | . 307 |  |
|  | 3 | Summer | 1.88 | . 192 |  |
|  | 4 | Fall | 1.74 | . 137 |  |
|  |  |  |  |  | . 235 |

Expressing the abundance estimates as percentages of the mean values facilitates comparison of trends (Fig. 1). Both survey and commercial indices show a pronounced downward trend in cod abundance during the period 1965-69. A measure of the similarity in these trends is provided by the fact that the correlation coefficient ( $r$ ) of survey indices with their corresponding commercial quarterly indices is quite high ( $\mathrm{r}=.74$ ), there being only a probability of about . 015 (with 8 degrees of freedom) of as high a value occurring by chance. While this is encouraging, more precise comparisons require that we take into account seasonal availability factors and related differences in the accuracy of abumdance indices for different seasons.

Obviously, there are large seasonal variations in availability of cod to
the commercial fishery with the maximum usually occurring in the winter and spring quarters (lst and 2nd) during the pre-spawning and spawning aggregations (Table 1, Fig. 1). The survey data at hand are insufficient to determine whether availability of cod to the survey trawl changes with the season, but it is reasonable to assume that it does. In general, however, differences in fishing practices between commercial and survey operations will result in larger seasonal availability changes in the commercial index because commercial vessels concentrate on fish aggregations, and thus, the relative differences between availability factors for the two types of index may be quite different in different seasons. Furthermore, in the case of the commercial fishery, the precision of the first and second quarter indices probably is greater than that for the third and fourth quarters because the volume of the fishery is much greater in the first two quarters.

For these reasons we should expect a higher correlation between survey and commercial indices taken one season at a time (or averages of season comblnations) than if all seasons are included in the correlation; and furthermore, on the average, we might expect higher correlations of survey indices with winter or spring commercial indices than with summer or fall commercial indices. In fact, the data for 4 X cod are consistent with such expectations.

The correlation of fall survey indices with fourth quarter comercial indices was higher ( $r=.77, P=.14$ ) than the correlation based on all seasons represented by survey indices, and it was higher still for fall survey indices and annual commercial indices ( $r=.83, P=.08$ ) even though these correlations each had only 3 degrees of freedom. Fall survey indices and the following winter (lst quarter) commercial indices have a still higher correlation of .91 ( $\mathrm{P}=.09, \mathrm{df}=2$ ).

Thus, in spite of the fact that there are so few comparisons available and that the survey indices are on a log scale (thereby not strictly comparable with the commercial indices on the linear scale), there is reasonably good agreement between the two data series for 4 X cod.
Div. 4X haddock

Survey and commercial data on Div. 4X haddock abundance are available in an identical form to that described above for Div. 4X cod (Table 2). Both survey and commercial indices indicate that abundance was lower in the latter part of 1968 and in 1969 than in earlier years but the trend was less prominent than for cod (Fig. 2). In fact, abundance held steady or even increased in 1966 and 1967 due to recruitment of the strong 1963 year-class. However, abundance began dropping rapidly in 1968. The correlation between all haddock survey indices and their corresponding commercial quarterly indices is only 0.32 ( $P=.38, \mathrm{df}=8$ ), which is not surprising in view of the non-ifnear trend over the entire period. However, fall indices for the two data sets give $r=$ $0.76(\mathrm{P}=.14, \mathrm{df}=3)$; and fali survey vs. annual commercial indices give $r=0.70(P=.19)$. Fall survey indices and the following winter (lst quarter) commercial indices have a correlation of 0.98 ( $\mathrm{P}=.02$ ). Thus, as for Div. 4X cod, the agreement between the two data series is encouraging even though the trend for haddock is not a simple linear one.

The inclusion of pre-recruits in the survey indices for haddock, does inflate the 1965 indices somewhat because large numbers of the 1963 year-class were present in 1965 survey catches. However, removal of haddock less than

40 cm from the survey indices (thus approximating commercial sizes) made only a minor change in the comparison. For example, the correlation between fall surveys and following first quarter comercial indices was reduced from .98 to . 91 .

Table 2. ICNAF Div. 4X haddock abundance indices from USA research vessel Albatross IV and Canadian commercial otter trawlers.

| Year | Quarter | Season | Survey Index $\log _{e}$ ( $1 \mathrm{~b} /$ tow) | Commercial index metric tons/hour |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | Quarterly | Annual |
| 1965 | 1 | Winter | - | . 306 |  |
|  | 2 | Spring | - | . 400 |  |
|  | 3 | Summer | 2.78 | . 250 |  |
|  | 4 | Fall | 3.61 | . 237 |  |
|  |  |  |  |  | . 298 |
| 1966 | 1 | Winter | - | . 391 |  |
|  | 2 | Spring | 3.72 | . 390 |  |
|  | 3 | Summer | - | . 279 |  |
|  | 4 | Fall | 3.22 | . 228 |  |
|  |  |  |  |  | . 322 |
| 1967 | 1 | Winter | - | . 370 |  |
|  | 2 | Spring | - | . 397 |  |
|  | 3 | Summer | - | . 236 |  |
|  | 4 | Fall | 3.87 | . 212 |  |
|  |  |  |  |  | . 304 |
| 1968 | 1 | Winter | - | . 423 |  |
|  | 2 | Spring | 3.13 | . 296 |  |
|  | 3 | Surmmer | - | . 248 |  |
|  | 4 | Fall | 2.93 | . 148 |  |
|  |  |  |  |  | . 279 |
| 1969 | 1 | Winter | - | . 352 |  |
|  | 2 | Spring | 2.53 | . 286 |  |
|  |  | Summer | 2.96 | . 215 |  |
|  | 4 | Fall | 2.68 | . 162 |  |
|  |  |  |  |  | . 254 |

Div. 4 T cod

The Groundfish Investigation of the St. Andrews Biological Station of the Fisheries Research Board of Canada has undertaken annual fall (SeptemberOctober) cod surveys in the southern Gulf of St. Lawrence (ICNAF Div. 4T) since 1957. On these surveys, 13 fixed stations were normally occupied, two halfhour otter-trawl tows being made at each station, the station locations being originally chosen to coincide with popular areas for comercial fishing. Those surveys undertaken in 1957-65 have been analyzed to provide abundance

Indices for the "summer" concentrations of the Div. 4T-4VN cod stock (Paloheimo and Kohler, 1968). In this case, survey catches, which were taken using otter trawls with small mesh codend liners, were adjusted by using selection ogives for the mesh sizes which were currently used by the commercial fleet. Thus, survey indices are expressed as pounds per tow of commercial-size cod (Table 3). Catches per trip by Gloucester-class otter trawlers ( $26-50$ gross tons) in August to October inclusive provide suitable commercial abundance indices for comparison (Table 3). Variations in the two series are almost identical (Fig. 3). The largest discrepancy lies in the 1957 values. However, the survey in that year was undertaken by a different vessel from those in following years, and the 1957 abundance index was adjusted on the basis of results from comparative fishing experiments between the two vessels. The discrepancy between survey and commercial values in 1957 may reflect errors in this adjustment. Even including the 1957 value the correlation coefficient between the two series is 0.90 which is highly significant ( $P<.001$, $\mathrm{df}=7$ ).

Table 3. ICNAF Div. 4 T cod abundance indices from Canadian research vessel surveys and Canadian commercial otter trawlers.

| Year | Survey index <br> lb/tow | Commercial index <br> 1000 lb/trip |
| :--- | :---: | :---: |
| 1957 | 393 | 14.3 |
| 1958 | 227 | 11.5 |
| 1959 | 250 | 12.8 |
| 1960 | 119 | 7.8 |
| 1961 | 136 | 8.6 |
| 1962 | 178 | 11.5 |
| 1963 | 159 | 9.6 |
| 1964 | 140 | 8.6 |
| 1965 | 111 | 6.2 |

Div. 4W haddock

Summer survey cruises in ICNAF Div. 4 W in 1958-60 by the St. Andrews Biological Station to investigate the distribution of haddock have been used to obtain abundance indices of haddock at ages 1-3 (Halliday, MS 1970). Abundance indices at ages 1-3 were found to be significantly correlated with abundance at age 4 estimated from the comercial fishery for the 1954-59 yearclasses ( $r=0.83, P=.04$, $\mathrm{df}=4$ ), when both data sets were transformed to $\log _{e}$ scale (Fig. 4).

Discussion
The extremely close agreement between research and commercial abundance indices in the 4 T cod study indicates that under certain circumstances very precise abundance indices can be obtained with modest sampling intensity by a research vessel. The key circumstances in this case probably are that since the survey was designed specifically for measuring cod abundance, stations were selected in locations known to be good cod fishing areas, and then exactly
the same stations were fished each year. An obvious requirement for this approach to be successful is that the density of fish in the selected habitats should remain in direct proportion to the total stock abundance from year to year. While this condition seems to have held in the case of $4 \mathrm{~T} \operatorname{cod}$, it would be risky to assume that constant proportionality holds generally, particularly when large stock changes occur.

In the case of the U.S. surveys in 4 X which employed stratified random sampling, the objective was to sample all trawlable habitats with probability proportional to their area, and to estimate abundance of all spectes. The random sampling design avoids the type of blas referred to above, but it does so at the expense of statistical precision. That is, sampling errors are almost certain to be larger than in cases where specific habitats are selected and then sampled each year. It is particularly encouraging then that the correspondence between commercial and survey indices in the 4 X studies appeared to be reasonably good. Admittedly, we would be on firmer ground with a few more years' data.

The 4 W haddock surveys represent a compromise between the 4 T and 4 X survey approaches, with sampling directed specifically toward haddock, but not confined to the same stations each year. Precision of the 4 W haddock pre-recrult indices would appear adequate for assessment purposes.

In conclusion, the degree of correspondence between abundance indices from research vessel surveys and commercial statistics in Subarea 4 provides very encouraging evidence that surveys can produce accurate measures of groundfish abundance. If sampling is directed toward a single stock, then the relatively modest sampling intensity represented by the surveys in 4 T and 4 W appears sufficient to provide a level of accuracy required for certain assessment needs. If sampling is directed toward estimation of the total biomass of groundfish, then it would appear that the sampling intensity of the U.S. surveys in 4 X is adequate to measure major stock changes. However, substantial increases in sampling intensity may be necessary in order to achieve the accuracy required for assessment of less drastic changes which nevertheless are significant from the standpoint of management.

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Fig. 1.


Fig. 2.



19. Accuracy of abundance indices for cod from St. Pierre Bank (ICNAF Div. 3Ps) based on Canada (Newfoundland) research vessel surveys in terms of comparisons with commercial abundance indices ${ }^{1}$<br>by A.T. Pinhorn<br>Fisheries Research Board of Canada Biological Station, St. John's, Newfoundland

## Introduction

One of the chief objectives of research vessel surveys is to estimate the absolute or relative abundance of year-classes of fish several years prior to their recruitment to the commercial fishery. The surveys conducted by the St. John's Biological Station have been no exception and the survival of yearclasses of cod and haddock has been followed for many years, although no actual quantitative predictions have been made. The purpose of the present paper is to evaluate the survey data from St. Pierre Bank (ICNAF Div. 3Ps) and to correlate abundance estimates of year-classes of cod at the pre-recruit stage with abundance estimates of these same year-classes after recruitment to the commercial fishery.

## Materials and methods

The sampling design traditionally used on groundfish cruises has been a standard line system. Lines and stations selected under this system are described by Pinhorn (1971) in a document presented to this meeting. Figure 1 shows the lines and stations fished on St. Plerre Bank during 1957-70, the period considered in the present paper.

Temperature conditions in this area vary with season and even from year to year within the same season (Templeman and Hodder, 1965). Also, primarily as a response to these changing temperature conditions, cod are known to migrate between depth zones from season to season (Templeman and May, 1965) and even to migrate to the Newfoundland coast in this area in the May-June period (Templeman, 1962). In view of these facts and since the survey cruises in 1957-65 were conducted during May-June but in 1967-70 were conducted during March-May (Fig. 2), the length compositions of the catches from each cruise were combined by depth zones of $0-50$ fathoms, $51-100$ fathoms and 101-150 fathoms. These were then converted to number per hour fished for each depth zone and the results are shown in Fig. 2. It became obvious that very few cod were caught deeper than 100 fathoms and this depth zone is omitted from Fig. 2. In fact, movements of cod occurred within depths less than 100 fathoms. It is further obvious that, except for occasional years of peculiar hydrographic conditions (e.g. 1958), the young cod of $2-3$ years of age ( $<37 \mathrm{~cm}$ ) remained for the most part in the 51-100-fathom depth zone during the entire period from March to June, whereas the older cod were found in the 51-100-fathom depth zone during the early part of this period, but moved onto the top of the bank in depths less than 50 fathoms during the later part of this March-June period (Fig. 2).

To evaluate the similarity of abundance indices calculated from research vessel surveys in successive years, indices of abundance of 2 - and 3-year-old cod of each year-class were compared with indices of abundance of 4- and 5-yearold cod of the same year-class in later cruises. In view of the facts outlined

[^16]above, three separate methods of computing these indices were used as follows:

1. Numbers of cod caught per hour fished for each year-class at each age were computed by applying an age-length key to numbers per hour length compositions as shown in Fig. 2 from sets made in 51-100 fathoms only for each cruise.
2. Numbers of cod caught per hour fished for each year-class at each age were computed for 2- and 3-year-old cod by applying an agelength key to number per hour length compositions from sets made in 51-100 fathoms only and for 4 - and 5 -year-old cod from sets made in 0-50 fathoms for the 1957-65 late spring to early summer cruises and from sets made in 51-100 fathoms for 1967-70 late winter to early spring cruises.
3. Numbers of cod caught per hour fished for each year-class at each age were computed by weighting the number per hour length compositions in each depth zone by the area of the depth zone as shown in Fig. 1. These areas are as follows:

| $0-50$ | fathoms | $-5,749$ | square miles |  |
| :---: | :---: | :---: | :---: | :---: |
| $51-100$ | $"$ | $-2,776$ | $"$ | $"$ |
| $101-150$ | $"$ | - | 869 | $"$ |

In the above computations, in years where no survey data were available (1961 and 1966) and for year-classes at the beginning and end of the period, numbers caught per hour for the missing ages were calculated by applying the average survival rates of the remaining year-classes between successive ages to the known number per hour at the age immediately before or after the missing ages.

In comparing indices of abundance of 2- and 3-year-old cod from the research cruises with indices of abundance of 4 - and 5 -year-old cod from the comercial fishery, the indices for 2- and 3-year-old cod calculated by Methods 1-3 above were used. The indices of abundance for 4- and 5-year-old cod from the commercial fishery were calculated in the following manner: Nominal catches and hours fished by Canada (Newfoundland) side trawlers of 151-500 tons were tabulated for the spring fishery for the $1959-68$ period. Average weight caught per hour fished was then calculated for each year and average number caught per hour fished calculated from this figure by using average weight values as determined from the Sampling Yearbooks for the respective years. Per mille length frequencies of catches of Canada (Newfoundland) side trawlers of 151-500 tons, as well as length frequencies of other countries which appeared similar to Canada (Newfoundland) frequencies, where the latter were lacking or few in number, were averaged and the resulting per mille length frequency adjusted to number per hour caught by these Canada (Newfoundland) side trawlers in each year. Research age-length keys were applied to these length frequencles for each year and indices of abundance calculated for 4and 5-year-old cod of each year-class. For 1961 and 1966, in which no agelength keys were available, keys from the adjacent years were averaged and applled to the length frequency in the particular year. Numbers of measurements used for each country are given below:

| Year |  | Country | No. of Measurements |
| :---: | :---: | :---: | :---: |
| 1959 | Canada | (Newfoundland) | 2,047 |
| 1960 | 1 | " | 520 |
|  | United | Kingdom | 2,013 |
| 1961 | Canada | (Newfoundland) | 834 |
|  | Portug |  | 500 |
| 1962 | United | Kingdom | 1,890 |
| 1963 |  | - | - |
| 1964 | Canada | (Newfoundland) | 267 |
| 1965 | " | " | 999 |
| 1966 | " | " | 2,428 |
| 1967 | " | " | 600 |
| 1968 | " | " | 945 |

## Results

Logarithmic transformations best described the relationships between abundance indices of 2- to 3-year-old cod and 4- to 5-year-old cod in successive research surveys. Correlation coefficients ( $r$ ) were generally larger than with the arithmetic straight line fits and were significantly different from zero in all cases, except ages 2-3 versus age 5 calculated by Method 3 (Table 1 and Fig. 3). Only two r-values were significant using arithmetic straight lines, between ages $2-3$ and age 5 and ages $2-3$ and ages $4-5$ as calculated by Method 1.

Table 1. Correlation coefficients and tests of significance between indices of abundance of 2- and 3-year-old cod and 4- and 5-year-old cod from Canada (Newfoundland) research surveys of St. Pierre Bank (ICNAF Div. 3Ps), 1957-70. Indices of abundance are computed by three different methods which are described in text.

| Method | Ages compared | Arithmetic straight line |  |  | Log-log straight line |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | r | df | t | r | df | t |
| 1 | 2-3 vs 4 | 0.52 | 10 | 1.91 | 0.64 | 10 | $2.64{ }^{1}$ |
|  | 2-3 vs 5 | 0.83 | 10 | $4.73{ }^{2}$ | 0.74 | 10 | $3.45{ }^{2}$ |
|  | 2-3 vs 4-5 | 0.72 | 10 | $3.20{ }^{2}$ | 0.72 | 10 | $3.26{ }^{2}$ |
| 2 | 2-3 vs 4 | 0.51 | 10 | 1.89 | 0.76 | 10 | $3.66{ }^{2}$ |
|  | 2-3 vs 5 | 0.56 | 10 | 2.13 | 0.59 | 10 | $2.31{ }^{1}$ |
|  | 2-3 vs 4-5 | 0.57 | 10 | 2.21 | 0.85 | 10 | $5.07{ }^{2}$ |
| 3 | 2-3 vs 4 | 0.40 | 10 | 1.40 | 0.59 | 10 | $2.32{ }^{1}$ |
|  | 2-3 vs 5 | 0.42 . | 10 | 1.45 | 0.50 | 10 | 1.84 |
|  | 2-3 vs 4-5 | 0.45 | 10 | 1.60 | 0.59 | 10 | $2.31{ }^{1}$ |
| ${ }^{1}$ si | icant at th | leve1 level |  |  |  |  |  |

On the other hand, arithmetic straight line fits best described the relationships between ages $2-3$ in the research surveys and ages $4-5$ in the commercial catches. Correlation coefficients were generally larger than with
the logarithmic transformation and were significant in all cases, except between ages 2-3 and age 5 calculated by Method 1 (Table 2 and Fig. 4). However, all r-values, except two, were still significant with the logarithmic transformation.

Table 2. Correlation coefficients and tests of significance between indices of abundance of 2- and 3-year-old cod from Canada (Newfoundland) survey catches and 4- and 5-year-old cod from Canada (Newfoundland) commercial catches of 151-500-ton side trawlers, St. Pierre Bank (ICNAF Div. 3Ps), 1959-68.

| Method | Ages compared | Arithmetic straight line |  |  | Log-log straight line |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | r | df | t |  | $\mathrm{df}$ | t |
| 1 | 2-3 vs 4 | 0.83 | 8 | $4.15{ }^{2}$ | 0.79 | 8 | $3.59{ }^{2}$ |
|  | 2-3 vs 5 | 0.54 | 8 | 1.79 | 0.32 | 8 | 0.94 |
|  | 2-3 vs 4-5 | 0.88 | 8 | $5.29{ }^{2}$ | 0.67 | 8 | $2.57^{1}$ |
| 3 | 2~3 vs 4 | 0.82 | 8 | 3.992 | 0.82 | 8 | $4.04^{2}$ |
|  | 2-3 vs 5 | 0.63 | 8 | 2.291 | 0.50 | 8 | 1.62 |
|  | 2-3 vs 4-5 | 0.88 | 8 | $5.33{ }^{2}$ | $\cdot 0.76$ | 8 | $3.29^{1}$ |
| 1 significant at the $5 \%$ level2 significant at the $1 \%$ level |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |

Except for the 1958 year-class, survival of the $1954-65$ year-classes as determined from research vessel surveys fluctuated only moderately with the weakest year-classes being about $75 \%$ less than the long-term average and the strongest year-classes being about equal to the average (Fig. 5A). The 1958 year-class was represented in the 1960 research vessel catches in significant numbers but was only caught in small numbers in later research vessel cruises and in commercial catches. The catch of this year-class as 2 -year-olds was not considered representative of its abundance and consequently, this yearclass is omitted from the above correlations. The 1966 and 1968 year-classes appear to be considerably stronger than average, while the 1967 year-class was slightly below average.

## Discussion

It appears from the results presented in this paper that indices of abundance as determined from survey cruises to St. Plerre Bank are accurate enough, or at least consistent enough from year to year, to allow for prediction of relative levels of recruitment to the commercial fishery, even with the present standard line system as used by the St. John's Biological Station. However, attempts at estimating absolute abundance levels of individual yearclasses would be more difficult with this system, since it does not allow for random coverage of the entire habitat of the species and hence, any attempt at stratifying the area and using the stations fished on each line as random sets in the various strata would result in some strata having no sets at all and others with very few.

In considering the problem of stratifying the area according to the system of sampling described by Grosslein (1968), if only the pre-recruits of
age-groups $0-3$ are being considered, then it appears that the problem may be simplified since in most years these age-groups are confined largely to the depth zone 51-100 fathoms and a stratification system within this zone may be sufficient. However, if older age-groups are considered, then the fact that these age-groups migrate to the top of the Bank and more important to the Newfoundland coast must be considered. Any survey then undertaken would have to be in winter or early spring before this migration takes place.

Comparison of year-class strengths with those given by Bulatova (1970) indicates some differences (Fig. 5C). Deviations from the long-term mean of the 1960-66 year-classes were similar in both sets of data for the 1960-63 year-classes, ranging from $25-75 \%$ below this mean. Also, the deviations for the 1965 year-class was reasonably similar in both cases, being equal to the mean for Canada (Newfoundland) and $25 \%$ above for Bulatova (1970). However, the strength of the 1964 year-class, while being $10 \%$ below the mean from Canada (Newfoundland) data, was $125 \%$ above the mean from Bulatova (1970). Also, the 1966 year-class, while being $200 \%$ above the mean from Canada (Newfoundland) data was only $60 \%$ above from Bulatova (1970) data. Bulatova (1970) also states that the 1967 year-class is probably not better than average, while the 1968 year-class was especially strong. This agrees favourably with the Canada (Newfoundland) data (Fig. 5A), from which the 1967 year-class was judged to be about $25 \%$ less than the long-term 1954-68 mean and the 1968 year-class about $150 \%$ greater than the mean.

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Fig. 1. Chart of St. Pierre Bank showing depth zones used in calculation of indices of abundance of cod and lines and stations fished during the period considered. Solid lines marked A, B and C delineate sections used for determination of depth zone areas.


Fig. 2. Number caught per 1-hour trawling in each 3-cm length-group in the two major depth zones from Canada (Newfoundland) research vessel surveys, St. Plerre Bank (ICNAF Div. 3Ps). Areas shaded and hatched to the left of vertical lines indicate cod less than 4 years old.


Fig. 3. Relationship between average indices of abundance of 2- to 3 -year-old cod and 4- to 5 -year-old cod of each year-class in successive research vessel surveys. See text for explanation of Methods l-3 of calculating indices.


Fig. 4. Relationship between average indices of abundance of 2- to 3. year-ild coí from reseaich veasel surveys and 4- to 5 -yearold cod from the commercial fishery by Canada (Newfoundland) side trawlers of 151-500 tons for each year-class.


Fig. 5. Percent deviations by year-class of abundance indices of cod from long-term average of: (A) 1954-68 year-classes from research data alone; (B) 1954-64 year-classes from research data and commercial data; (C) 1960-66 year-classes from Canada (Newfoundland) research data and data of Bulatova (1970).
20. Some observations on accuracy of abundance indices derived from research vessel surveys ${ }^{1}$

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## Introduction

In order to help evaluate the cost-benefit ratio of surveys it is necessary to have some idea of the magnitude of change in stock size that is considered significant, as well as the magnitude of change we are able to detect and with what probability. Clearly one of the most important questions is whether surveys can measure changes in abundance with sufficient accuracy to permit meaningful assessment of the short-term affects of fishing. However, it is important to remember that we are also concerned with long tern changes involving not just a few priority species but the entire groundfish community. In general a lower level of accuracy probably would suffice for monitoring long term changes than in the case of assessment on a year-to-year basis. My principal aim here is to provide some information on what accuracy is possible with catch-per-haul statistics from research vessel surveys.

When considering accuracy of estimates, we must distinguish between statistical precision or sampling error (variance) and bias. That is, an estimate may be very precise in terms of a small variance but have a large bias, and therefore not be very accurate. In our problem we are concerned not only with precision but also with the possible biases in the survey abundance index (catch per standard haul) as a relative measure of absolute abundance. That is, we may consider our index unbiased if there is a constant proportionality (catchability coefficient) between the relative abundance index and the true absolute abundance of the stock. The next step is to estimate this coefficient so that we can estimate actual total numbers in the population.

Evidence to be presented later suggests that the assumption of constant proportionality is not unreasonable for certain species and observed stock changes in the case of joint US-USSR surveys. Consequently the following data on precision of abundance indices from these surveys probably reflects the general order of accuracy obtainable in measures of change in absolute stock size. Admittedly we will be on firmer ground when we can estimate variability of catchability coefficients, by utilizing direct (camera, acoustic) measures of abundance in conjunction with trawling.

Statistical characteristics of trawl catch data
As is well known trawl catches are highly variable even within relatively restricted areas because fish are not uniformly distributed; and random trawl hauls result in a frequency distribution of catches which is highly skewed. A major consequence of this skewness is that the variance is generally much larger than the mean resulting in very imprecise (although unbiased in the statistical sense) estimates of the mean, and even less reliable estimates of the variance

[^17]itself, except with very large sample sizes. That is, the standard error associated with the variance is particularly susceptible to departures from normality, and without a rellable estimate of the variance of course it is not possible to calcualte meaningful confidence limits about the mean.

A standard approach to this general problem is to stratify the population to be sampled into high and low density units or strata, and then sample randomly within individual strata within each of which skewness is then reduced. Control of varlabillty in this manner is one of the primary advantages to be gained from the technique of stratified random sampling. However in the case of trawl catches, considerable skewness remains even after stratification. For example the varlability of varlance estimates for haddock trawl catches on U.S. surveys, reflects the fact that catches within individual strata are still highly skewed (Table 1). Sampling strata used in the surveys discussed here are shown in Figures 1 and 2.

Another well known approach is to try to find a transformation which normalizes the frequency distribution of variables. We have found that on the average, stratum variances of trawl catches are approximately proportional to the square of the mean, i.e. the standard deviation is proportional to the mean. This is true for haddock (Fig. 3) and for many other species as well. This relation indicates that a log transformation is appropriate, and such a transformation tends to normalize the data and stabilize the variance (i.e. make means and variances independent). Also the log transformation converts multiplicative effects into linear additive effects. In terms of our problem of estimating proportional changes in abundance, this means that linear changes on a log scale represent estimates of multiple or factor changes on the original scale. That is, the anti-log of the difference between two 10 g means represents the proportionality constant relating means on the linear scale. The estimates of proportional change on the original scale are believed to be essentially unbiased in the statistical sense, but it should be noted that the re-transformed mean is a biased estimate of the true mean on the linear scale (an unbiased estimate is theoretically possible).

## Calculation of stratified mean and variance

The basic index of abundance dealt with here is the stratified mean catch per standard haul, calculated by weighting each stratum mean according to the proportional size (area) of the stratum relative to all strata in the set. The variance of a stratified mean is similarly derived by weighting each stratum variance in proportion to the stratum area and inversely according to the number of hauls in the stratum.
Table 1. Bank. A $\grave{b}$ batross IV surveys.

|  | STRATUM 16 |  |  |  | STRATUM 19 |  |  |  | Stratum 20 |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| CRUISE | No. hauls | Mean | Variance | Std. deviation | No. hauls | Mean | Variance | Std. deviation | No. hauls | Mean | Variance | Std. deviation |
| 63-05 | 7 | 41 | 2,740 | 52 | 4 | 126 | 22,442 | 150 | 3 | 7 | 52 | 7 |
| 63-07 | 7 | 101 | 4,330 | 66 | 4 | 291 | 66,992 | 259 | 4 | 115 | 33,379 | 183 |
| 64-01 | 10 | 41 | 857 | 29 | 7 | 147 | 37,875 | 194 | 5 | 37 | 1,322 | 36 |
| 64-210 | 8 | 300 | 338,823 | 582 | 5 | 364 | 209,248 | 457 | 5 | 356 | 70,072 | 264 |
| 64-13 | 7 | 148 | 31,926 | 179 | 6 | 168 | 26,652 | 163 | 5 | 335 | 155,074 | 394 |
| 65-2 | 6 | 73 | 6,309 | 80 | 6 | 392 | 243,932 | 494 | 5 | 21 | 338 | 18 |
| 65-510 | 8 | 405 | 682,555 | 826 | 6 | 800 | 2,019,784 | 1421 | 5 | 618 | 188,942 | 435 |
| 65-14 | 7 | 78 | 3,266 | 57 | 5 | 171 | 14,377 | 120 | 5 | 332 | 160,830 | 401 |
| 66-601 | 7 | 73 | 17,357 | 132 | 6 | 49 | 6,058 | 78 | 5 | 43 | 1,243 | 35 |
| 66-614 | 7 | 62 | 1,423 | 38 | 6 | 54 | 15,495 | 124 | 5 | 126 | 11,584 | 108 |
| 67-721 | 8 | 14 | 564 | 24 | 9 | 52 | 4,096 | 64 | 6 | 37 | 4,140 | 65 |
| 68-803 | 9 | 49 | 5,533 | 74 | 8 | 42 | 1,189 | 34 | 6 | 13 | 351 | 19 |
| 68-817 | 8 | 19 | 2,850 | 53 | 9 | 0 | - | - | 6 | 25 | 3,574 | 60 |
| 69-902 | 14 | 71 | 26,570 | 163 | 8 | 45 | 1,831 | 43 | 6 | 3 | 41 | 6 |
| 69-908 | 10 | 7 | 185 | 14 | 9 | 6 | 124 | 11 | 6 | 23 | 2,610 | 51 |
| 69-911 | 12 | 4 | 117 | 11 | 9 | 7 | 413 | 20 | 6 | 16 | 1,137 | 34 |
| 70-703 | 10 | 130 | 120,926 | 348 | 8 | 11 | 409 | 20 | 5 | 5 | 1,76 | 9 |

Computational formulae are:

$$
\begin{aligned}
& \overline{\mathrm{y}}_{\boldsymbol{s t}}=\frac{1}{\mathrm{~N}} \sum_{\mathrm{h}} \mathrm{~N}_{\mathrm{h}} \overline{\mathrm{y}}_{\mathrm{h}} \\
& \mathrm{v}\left(\overline{\mathrm{y}}_{\mathrm{st}}\right)=\frac{1}{\mathrm{~N}^{2}} \sum_{\mathrm{h}} \frac{\mathrm{~N}_{\mathrm{h}^{2} \mathrm{~S}_{h^{2}}}^{\mathrm{n}_{\mathrm{h}}},}{}
\end{aligned}
$$

where $\bar{y}_{\text {st }}$ and $V\left(\bar{y}_{\text {st }}\right)$ are the stratified mean catch per haul and its variance respectively, of some set of strata, and
$N_{h} \quad=$ area of the $h^{\text {th }}$ stratum
$\mathrm{N} \quad={ }_{h}^{\Sigma} N_{h}=$ total area of all strata in the set
$\vec{y}_{h} \quad=$ mean catch per haul in the $h^{\text {th }}$ stratum
$n_{h}=$ number of standard hauls in $h^{\text {th }}$ stratum
$S_{h^{2}}=$ variance of catches in the $h^{\text {th }}$ stratum

## Examples of precision on 1inear scale

It is of interest to look at some examples of sampling errors of stratified means on a non-transformed scale before proceeding on to the log scale. Recall that in the examples of haddock data for individual strata, the standard deviation was on the average about equal to the mean (Fig. 3, Table 1). That is, coefficients of variation (ratio of standard deviation to the mean) were on the order of 100 percent with 5-7 hauls per sample. In the case of stratified means for haddock on Georges Bank (representing about 60 hauls in strata 13-25 combined) the average $C V$ is only about 25 percent (Table 2). Similar values were obtained for cod.

In spite of the observed variability in estimates of individual stratum variances, we note that the CV's of the stratified means are reasonably consistent from year to year suggesting that the estimates of $V\left(\bar{y}_{s t}\right)$ may be approximately correct. Essentially we have computed a weighted mean of variances from 13 strata, and since most of these strata appear to have about the same variance this would account for the consistency among estimates of $v\left(\bar{y}_{s t}\right)$.

Stratified means for yellowtail on Georges Bank show CV's similar to those for cod and haddock (Table 3). Also shown in Table 3 are stratified means for the three principal strata for yellowtail, representing about half of the total area of the strata set, 13-25. The CV's are only slightly greater on average for this subset of strata than for the entire set, although there were less than half as many hauls in the subset. Very little information on yellowtail was gained by sampling outside these three principal strata.

Table 2. Stratified mean catch per haul (pounds, linear) of cod and haddock on Georges Bank, (strata 13-25), and estimates of precision. Albatross IV fall surveys.

|  |  | COD |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Year | Mean | Variance | S.D. | S.D. $/$ Mean | Mean <br> 2 S.D. |
| 1963 | 24.18 | 43.35 | 6.58 | .27 | $11.0-37.3$ |
| 1964 | 15.74 | 20.89 | 4.57 | .29 | $6.6-24.9$ |
| 1965 | 15.90 | 26.04 | 5.10 | .32 | $5.7-26.1$ |
| 1966 | 11.10 | 5.87 | 2.42 | .22 | $6.3-15.9$ |
| 1967 | 18.43 | 17.85 | 4.22 | .23 | $10.0-26.9$ |
| 1968 | 11.66 | 8.54 | 2.92 | .25 | $5.8-17.5$ |
| 1969 | 10.91 | 4.79 | 2.19 | .20 | $6.5-15.3$ |
|  |  |  |  |  |  |
|  |  |  |  | HADDOCK |  |
| 1963 | 112.83 | 590.75 | 24.30 | .22 | $64.2-161.4$ |
| 1964 | 165.68 | 1032.11 | 32.13 | .19 | $101.4-229.9$ |
| 1965 | 123.66 | 411.58 | 20.29 | .16 | $83.1-164.2$ |
| 1966 | 47.22 | 99.39 | 9.97 | .21 | $27.3-67.2$ |
| 1967 | 44.05 | 103.86 | 10.19 | .23 | $23.7-64.4$ |
| 1968 | 20.53 | 52.18 | 7.22 | .35 | $6.1-35.0$ |
| 1969 | 12.70 | 16.62 | 4.08 | .32 | $4.5-20.9$ |

Table 3. Stratified mean catch per haul (pounds, linear) of yellowtail on Georges Bank, and estimates of precision. Albatross IV fall surveys.

| STRATA 13-25 (15,300 sq miles) |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Year | Mean | Variance | S.D. | S.D./Mean | $\begin{aligned} & \text { Mean } \pm \\ & 2 \text { S.D. } \end{aligned}$ | No. Hauls |
| 1963 | 18.00 | 11.56 | 3.40 | . 19 | 11.2-24.8 | 57 |
| 1964 | 18.58 | 53.27 | 7.30 | . 39 | 4.0-33.2 | 63 |
| - 1965 | 12.36 | 15.73 | 3.97 | . 32 | 4.4-20.3 | 66 |
| 1966 | 5.38 | 3.07 | 1.75 | . 32 | 2.1-8.6 | 67 |
| 1967 | 9.71 | 6.91 | 2.63 | . 27 | 4.4-15.0 | 65 |
| 1968 | 14.73 | 11.33 | 3.37 | . 23 | 8.0-21.5 | 62 |
| 1969 | 12.02 | 9.73 | 3.12 | . 26 | 5.8-18.3 | 66 |
| 1970 | 6.37 | 3.49 | 1.87 | . 29 | 2.6-10.1 | 70 |
|  |  |  |  |  |  |  |
| Strata 13, 16, 19 (7,800 sq miles) |  |  |  |  |  |  |
| 1963 | 23.10 | 33.19 | 5.76 | . 25 | 11.6-34.6 | 16 |
| 1964 | 32.10 | 194.97 | 13.96 | . 43 | 4.2-60.0 | 18 |
| 1965 | 18.48 | 56.99 | 7.55 | . 41 | 3.4-33.6 | 19 |
| 1966 | 8.71 | 11.35 | 3.37 | . 39 | 2.0-15.4 | 19 |
| 1967 | 16.58 | 25.96 | 5.10 | . 31 | 6.4-26.8 | 25 |
| 1968 | 24.50 | 40.78 | 6.38 | . 26 | 11.7-37.3 | 25 |
| 1969 | 21.44 | 36.96 | 6.08 | . 28 | 9.3-33.6 | 30 |
| 1970 | 10.69 | 12.44 | 3.53 | . 33 | 3.6-17.8 | 24 |

Table 4. Stratified mean catch per haul ( $1 \mathrm{~b}, \log _{\mathrm{e}}$ scale) and measures of precision for selected species. Albatross IV fall surveys, Strata 13-25.

| YELLOWTAIL |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Year | Mean | Variance | S.D. | $\begin{aligned} & \text { S.D. / } \\ & \text { Mean } \end{aligned}$ | 2 S.D. | Mean $\pm$ $2 \text { S.D. }$ | Factor diff. |
| 1963 | 1.97 | . 026805 | . 1637 | . 08 | . 33 | 1.64-2.30 | 1.9 |
| 1964 | 1.41 | . 037142 | . 1927 | . 14 | . 38 | 1.03-1.79 | 2.1 |
| 1965 | 1.32 | . 029119 | . 1706 | . 13 | . 34 | .98-1.66 | 2.0 |
| 1966 | 0.96 | . 025860 | . 1608 | . 17 | . 32 | .64-1.28 | 1.9 |
| 1967 | 1.32 | . 027724 | . 1665 | . 13 | . 33 | .99-1.65 | 1.9 |
| 1968 | 1.40 | . 038260 | . 1956 | . 14 | . 39 | 1.01-1.79 | 2.2 |
| 1969 | 1.35 | . 025200 | . 1587 | . 12 | . 32 | 1.03-1.67 | 1.9 |
| 1970 | 0.96 | . 0204 | . 1428 | . 15 | . 28 | .68-1.24 | 1.8 |
| HADDOCK |  |  |  |  |  |  |  |
| 1963 | 3.34 | . 052176 | . 2284 | . 07 | . 46 | 2.88-3.80 | 2.5 |
| 1964 | 3.86 | . 080315 | . 2834 | . 07 | . 57 | 3.29-4.43 | 3.1 |
| 1965 | 4.02 | . 042355 | . 2058 | . 05 | . 41 | 3.61-4.43 | 2.3 |
| 1966. | 2.43 | . 044512 | . 2110 | . 09 | . 42 | 2.01-2.85 | 2.3 |
| 1967 | 2.45 | . 052075 | . 2282 | . 09 | . 46 | 1.99-2.91 | 2.5 |
| 1968 | 1.15 | . 029587 | . 1720 | . 15 | . 34 | 0.81-1.49 | 2.0 |
| 1969 | 1.10 | . 021536 | . 1467 | . 13 | . 29 | 0.81-1. 39 | 1.8 |
| 1970 | 1.35 | . 0345 | . 1857 | . 14 | . 37 | 0.98-1.72 | 2.1 |
| COD |  |  |  |  |  |  |  |
| 1963 | 1.75 | . 084829 | . 2912 | . 17 | . 58 | 1.17-2.33 | 3.2 |
| 1964 | 1.29 | . 056270 | . 2372 | . 18 | . 47 | 0.82-1.76 | 2.6 |
| 1965 | 1.32 | . 041737 | . 2043 | . 15 | . 41 | 0.91-1.73 | 2.2 |
| 1966 | 1.20 | . 040673 | . 2017 | . 17 | . 40 | 0.80-1.60 | 2.2 |
| 1967 | 1.74 | . 047301 | . 2175 | . 12 | . 44 | 1.30-2.18 | 2.4 |
| 1968 | 1.04 | . 031888 | . 1786 | . 17 | . 36 | 0.68-1.40 | 2.1 |
| 1969 | 1.32 | . 025381 | . 1593 | . 12 | . 32 | 1.00-1.64 | 1.9 |
| 1970 | 1.35 | . 0332 | . 1822 | . 13 | . 36 | 0.99-1.71 | 2.1 |

Examples of precision on $\log$ scale
On the log scale the variances are nearly stabilized and the $C V^{\prime} s$ of stratified means are on the order of $10-15$ percent for the same species and strata (Table 4). However note that now we are interested in the absolute rather than relative size of the standard deviation. For haddock $\pm 2$ S.D.'s $( \pm .40)$ corresponds to $\pm 50$ percent on the linear scale. Thus there is no great improvement in the size of difference (proportional change on linear scale) we are able to detect as compared with the non-transformed scale, but we have more efficient estimates of those differences over the range of abundance levels, and the estimated confidence limits more closely approximate true 95 percent confidence intervals. Results of stratified estimates for cod and haddock off western Nova Scotia are comparable to those on Georges Bank (Table 5).

Table 5. Stratified mean catch per haul (loge pounds) and variance estimates for cod and haddock off western Nova Scotia. Albatross IV autumn surveys in strata 31-35, 41, 42.

| HADDOCK |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Year | Mean | Variance | S.D. | S.D. / <br> mean | 2 S.D. | 95\% CI | Factor difference |
| 1965 | 3.61 | . 1918 | . 4379 | . 12 | . 88 | 2.73-4.49 | 5.8 |
| 1966 | 3.22 | . 1321 | . 3634 | . 11 | . 73 | 2.50-3.94 | 4.3 |
| 1967 | 3.87 | . 1073 | . 3276 | . 08 | . 66 | 3.21-4.53 | 3.7 |
| 1968 | 2.93 | . 0598 | . 2445 | . 08 | . 49 | 2.45-3.41 | 2.7 |
| 1969 | 2.68 | . 0593 | . 2435 | . 09 | . 49 | 2.20-3.16 | 2.7 |
| 1970 | 2.82 | . 0352 | . 1876 | . 07 | . 38 | 2.44-3.20 | 2.1 |
| COD |  |  |  |  |  |  |  |
| 1965 | 3.25 | . 1492 | .3863 | . 12 | . 77 | 2.47-4.03 | 4.7 |
| 1966 | 2.71 | . 1608 | . 4010 | . 15 | . 80 | 1.91-3.51 | 5.0 |
| 1967 | 2.16 | . 1051 | . 3242 | . 15 | . 65 | 1.52-2.80 | 3.7 |
| 1968 | 1.86 | . 0949 | . 3080 | . 16 | . 62 | 1.24-2.48 | 3.5 |
| 1969 | 1.74 | . 0887 | . 2978 | . 17 | . 60 | 1.14-2.34 | 3.3 |
| 1970 | 1.77 | . 0500 | . 2236 | . 13 | . 45 | 1.32-2. 22 | 2.5 |

The most significant feature of these data is that they indicate the present survey cannot detect with high probability, proportional changes in abundance which are less than a factor of about 2. That is, the $\log _{e}$ difference between the lower and upper limits of the 95 percent $C$. I. is about 0.7 corresponding to a factor difference of 2 on the linear scale; and to be very sure that two means are significantly different there must be no overlap in the 95 percent confidence intervals.

Sample size vs. precision
Some first approximations have been made of the relation between precision of stratified means and sample size (total number of hauls). The calculations are based on the general formula for estimating required sample size in stratified random sampling:

$$
n=\frac{\sum_{h h_{h}{ }^{2} S_{h}^{2}}^{h}}{V+\frac{1}{N} \sum_{h} W_{h} S_{h}^{2}}
$$

and in terms of this problem,
$W_{h}$ and $S_{h}$ are as defined earlier,
$w_{h}=\frac{n_{h}}{n}$, the observed relative sampling effort in the
$h^{\text {th }}$ stratum (the ratio of the number of hauls in
the $h^{\text {th }}$ stratum to the total number of hauls,
$n$, in all strata of the specified set)
$V=$ desired variance of the stratified mean
$N=$ total number of possible hauls in the area represented by strata in the set.

Since the number of hauls in our survey is very small relative to the total number possible (strata $13-25$ cover roughly 15,000 square miles and each standard haul covers approximately . 01 square mile), the second term in the denominator is extremely small compared with the first.

Thus,

$$
\mathrm{n}=\frac{1}{V} \sum_{\mathrm{h}} \frac{\mathrm{~W}_{h}^{2} \mathrm{~S}_{h}^{2}}{\mathrm{~W}_{\mathrm{h}}}
$$

Using the above formula and average values of $S_{h}{ }^{2}$ for haddock, and $w_{h}$ based on eight Albatross IV fall surveys, estimates were made of the sample sizes required to achieve various levels of precision. For example, if we wanted to be able to detect proportional changes in abundances of $\pm 20$ percent with high probability, this would require an interval of $\pm 2 \mathrm{~S} . \mathrm{D}^{\prime} \mathrm{s}= \pm .18$ on the natural log scale, and thus S.D. $=.09$ and $V=.0081$. Substituting this value of $V$ in the above formula, $n=338$ hauls. Results of calculations for levels of precision between $10-100$ percent of the stratifled mean for haddock are given in Table 6. The same computations for yellowtail in strata 13, 16 , and 19 (representing about half of Georges Bank) are also shown in Table 6 .

Table 6. First approximations to sample sizes (total number hauls) required for specified precision of stratified mean abundance indices ( $\log _{\mathrm{e}}$ catch/haul in pounds) from Albatross IV surveys on Georges Bank. ${ }^{1}$

| LEVEL OF PRECISION |  | Total number hauls required, (approximately proportional allocation) |  |
| :---: | :---: | :---: | :---: |
| Percentage | 2 standard |  |  |
| change <br> linear scale | deviations, <br> log scale | $\frac{\text { Haddock }}{\left(\frac{\text { strata }}{13-25)}\right.}$ | $\frac{\text { Yellowta11 }}{(\text { strata } 13,16,19)}$ |
| $\pm 10 \%$ | $\pm .10$ | >500 | >500 |
| $\pm 20 \%$ | $\pm .18$ | 338 | 253 |
| $\pm 30 \%$ | $\pm .26$ | 164 | 120 |
| $\pm 50 \%$ | $\pm .40$ | 70 | 51 |
| $\pm 100 \%$ | $\pm .69$ | 23 | 17 |

1 An empirical measure of the improvement in precision with increase In samyle size was obtained on the 1971 spring groundfish survey by pooling results of two cruises on Georges Bank, one in March and one in May. The pooled data shown below represent an increase in numbers of hauls of about 50 percent over the standard sampling rate, and resulted in reductions in standard deviations of about the magnitude predicted by the analysis based on the $1963-70$ series of cruises shown above.
--Spring 1971 groundfish survey.

|  | Cruise 1 <br> No. <br> 2SD's Hauls |  | Cruise  <br> 2 SD's |  | $\begin{aligned} & \text { Cruises } \\ & 2 \text { SD's } \\ & \hline \end{aligned}$ | $\begin{gathered} 1 \& 2 \\ \text { No. } \\ \text { Hauls } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $\begin{gathered} \text { Haddock } \\ \text { (Strata 13-25) } \end{gathered}$ | . 38 | 71 | . 53 | 37 | . 32 | 108 |
| $\begin{gathered} \text { Yellowtail } \\ \text { (Strata 13, 16, 19) } \end{gathered}$ | . 52 | 30 | . 51 | 17 | . 37 | 47 |

These data suggest that the cost of detecting with high probability changes of stock size as small as $\pm 10$ percent would be extremely high. It is even doubtful that we could justify the cost of measuring changes within $\pm 20$ percent; to get to this level it would appear that for haddock we would need to make nearly 5 times as many hauls as in the current survey which employs about 65 hauls in strata $13-25$ and achieves a precision of roughly $\pm 50$ percent (Table 6). In' sampling for yellowtail it would appear that we would need almost as many hauls for strata 13,16 and 19 alone, in order to obtain comparable levels of precision.

These results should be considered as first approximations since we have not fully investigated all of the characteristics of these data. For example it is possible that some improvement could be achieved with a modified log transformation which would further improve normalization of the data. Also it is possible that we could make a significant gain in precision by additional stratification according to time of day, for those species exhibiting strong
diurnal variations in availability. Additional stratification would cost something however, either in terms of fewer degrees of freedom for estimating stratum means and variances, or additional time at sea, or both. Thus there is no guarantee that additional stratification would achieve a net gain in information per unit cost. Further it is possible that the region could be more effectively stratified, for example by utilizing additional information on bottom sediments relative to groundfish distribution. However this too could only result in slight gains so long as we are interested in many species distributed over a wide area.

I think the most promising approach lies in controling or at least monitoring the haul-to-haul performance of the trawl; for example we do not have a precise measure of groundspeed, nor do we know what variations occur in wingspread, headrope height or bottom contact. Even direction of tow relative to bottom currents may be important for some species.

Even after all such improvements are incorporated however, it seems clear to me that there cannot be any drastic change in the observed relation between precision and sample size. The hard fact is that in sampling organisms with highly contagious distributions, achieving high precision will require intensive sampling.

So far we have been considering the precision of a single mean. It is of course possible to combine seasonal means into a single annual index which would have a smaller variance. For example if the means of two surveys were averaged, the standard deviation of the resulting mean would be reduced by approximately a factor of 0.7 (assuming homogeneous variances for the original means). Thus if the separate standard deviations were on the order of .2 (corresponding to a $\pm 50$ percent level of precision), the standard deviation of the combined mean would be about . 14 , corresponding to a $\pm 30$ percent precision level. Essentially the same precision would have been achieved by simply doubling the sampling effort on one cruise, and in that sense there would be no gain in accuracy through combination of two cruises. However by combining results of more than one season within a given year, there is less likelihood of bias due to variation in seasonal availability factors.

Finally it should be noted that in most cases it usually takes at least several years for major changes in stock size to occur. Given annual surveys, we then have a number of points in a time series with which to test for a significant slope or trend, and precision of such a test would be greater than that indicated for a single survey.

## Comparisons between research and commercial abundance indices

Returning now to the more general concept of accuracy, we need to consider further the problem of blas in conjunction with precision. In particular we are concerned about the possibility that the ratio of our relative abundance indices to the absolute (unknown) abundance may not be constant at difference levels of absolute abundance. We may gain some insight into this question by comparing abundance indices derived from both research and commercial catch data. However we must use care in making such comparisons because both types of data are subject to error. The commercial data are potentially more subject to serious bias, and research data are usually characterized by larger sampling errors.

The most serious biases in commercial abundance data arise from unknown changes in the effective unit of effort usually related to economic or technological factors. Even with standard gear however, bias can result simply from variations in avilability of fish. With proper sample design the research vessel index is free of the first bias, but still may be subject to bias from changes in availability. For example the catchability coefficient for a given species and research trawl may change due to a change in vertical distribution of the species, in response to some environmental factor or even as a function of absolute abundance itself. The possibility of a significant bias of this type intuitively would seem to be much greater for a species for which the trawl has a very low efficiency. We shall return to this point later in comparing joint US-USSR survey results.

From the standpoint of precision it is important to recognize that the commercial abundance index nearly always will be more precise than a research index simply becuase it is based on a very large number of hauls. However we seldom obtain variance estimates for commercial indices since at best it is a very complicated task involving many sources of error. It is a relatively simple matter to obtain statistically valid estimates of sampling error from surveys but unfortunately the errors are large.

With the above characteristics in mind we may now turn to some comparisons of research and commercial indices. Fourth quarter U.S. landings/day figures for cod, haddock, and yellowtall on Georges Bank, and U.S. fall survey abundance indices for strata 13-25, are tabulated for the period 1963-1969 in Table 7. The percentage deviations of each index from the '63-'69 mean are plotted in Figure 4, and it is clear that the two indices are correlated for haddock and yellowtall.

For yellowtail the commercial and research indices show quite similar trends in relative abundance; and the magnitude of changes indicated by the research indices was not much greater than that indicated by the commercial indices (Fig. 4). Correlation coefficients were . 95 (linear scale, survey) and . 81 (log scale, survey), and both are significant at the 95 percent probability level.

The correspondence is perhaps almost too good in this case. That is, if the research index is accurate to within only $\pm 50$ percent changes in abundance, then one might not expect such close correspondence from year to year when the actual yellowtail abundance (based on fairly reliable commercial indices) appeared to vary by no more than about 30 percent from the mean. In other words there may be some indication here that variance estimates may be inflated. More detailed study will be required to clarify this notion.

For Georges Bank haddock correlation coefficients are also significant at the 95 percent level - . 74 and . 84 for linear and $\log$ scales respectively. Corresponding trends in abundance are indicated but the research indices show a much greater magnitude of change in stock size than is indicated by the U.S. commercial index. In this case however the commercial indices are belleved to have been negatively biased particularly in the mid-1960's as has been described by Hennemuth (1968). Another feature is that the efficiency of vessels remaining in the fishery after 1967 probably was above average, which might be the explanation for the apparent discrepancy in trend between the two sets of indices in the late 1960's. It should be emphasized that changes in efficiency of commercial fleets are quite likely when stock levels change drastically.

There is less consistency between commercial and research indices for Georges Bank cod than for yellowtail and haddock. Up to 1967 there was a rough similarity in trends, but thereafter the correspondence is poor (Fig. 4). Correlation coefficients do not differ significantly from zero. In the later years it is possible that the scarcity of haddock may have resulted in a partial shift of effort toward cod, in which case the commercial index would have a positive bias. This too will require more detailed study.

Another set of comparisons is provided by U.S. commercial and research indices for haddock off western Nova Scotia (Table 8). The best comparison is afforded by the first quarter commercial indices vs. the spring research indices and these show quite a consistent picture both with respect to trend and magnitude of change (Fig. 5). Trends are basically similar between fall surveys and annual comercial indices, but an unusually large discrepancy occurred in 1967. Sampling error was not particularly high in that year (see Table 5) and so far I have no explanation for the apparent discrepancy.

Still another set of comparisons is avallable for red hake in southern New England. During the period 1965-1968 there was a rapid steady decline in abundance shown by both the catch per haul statistics of the USSR fleet and the U.S. survey (Table 9, Fig. 6). The commercial data suggest that by 1968 abundance had dropped to about one-quarter the 1965 level, and the survey data imply a decline to about one-third the 1965 level. Abundance appeared to increase again in 1969 as indicated by both comercial and research indices. In contrast to southern New England, comparisons for Georges Bank show poor correspondence between the commercial and research data for red hake (Table 9, Fig. 7). This may be partly due to the fact that after 1965 the principal fishing effort by the Soviet fleet on red hake occurred in southern New England, and Georges Bank effort was not directed specifically toward red hake.

To summarize briefly the comparisons among commercial and research indices, it appears that survey indices more often than not provide about the same trends and relative changes in stock size as do commercial indices. This I think is basically encouraging. The problem now is how to improve precision.

Table 7. Fourth quarter U.S. comercial abundance indices and Albatross IV fall survey indices for cod, haddock and yellowtail on Georges Bank. Commercial index: Landings/day ( $1 \mathrm{~b} \times 10^{-3}$ ) 5 Z east. Survey index: stratified mean catch/haul (1b, linear and $\log _{\mathrm{e}}$ ) strata 13-25.

| Year | COD |  |  |  | HADDOCK |  |  | YELLOWTAIL |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Comm. | Survey |  | Comm. | Survey |  | Comm. | Survey |  |
|  |  | 1 b | $e^{1 b}$ |  | 1b | $e^{1 b}$ |  | 1 l | $\mathrm{e}^{1 \mathrm{l}}$ |
| 1963 | 2.1 | 25.1 | 1.8 | 8.2 | 118.6 | 3.3 | 8.7 | 21.6 | 2.0 |
| 1964 | 1.4 | 15.6 | 1.3 | 10.8 | 193.6 | 3.9 | 8.0 | 22.3 | 1.4 |
| 1965 | 0.8 | 7.5 | 1.3 | 14.9 | 131.0 | 4.0 | 7.3 | 14.7 | 1.3 |
| 1966 | 1.8 | 8.8 | 1.2 | 9.4 | 51.4 | 2.4 | 4.5 | 6.5 | 1.0 |
| 1967 | 2.5 | 20.0 | 1.7 | 5.7 | 43.1 | 2.4 | 6.1 | 11.7 | 1.3 |
| 1968 | 2.8 | 10.8 | 1.0 | 6.1 | 19.2 | 1.2 | 7.7 | 17.7 | 1.4 |
| 1969 | 5.1 | 7.5 | 1.3 | 4.8 | 5.6 | 1.1 | 6.1 | 14.4 | 1.4 |

Table 8. Haddock abundance indices for 4X based on U.S. commercial data and Albatross IV surveys. Commercial index: U.S. landings per day, metric tons rd. fresh, Browns Bank survey: stratified mean catch per haul ( $\log _{e}$ pounds), strata 31-35, 41, 42.

|  | Commercial |  | Survey |  |
| :--- | :---: | :---: | :---: | :---: |
| Year | Annual | lst Qtr. | Fall | Spring |
| 1963 |  | 6.9 |  |  |
| 1964 | 7.5 | 6.9 | - | - |
| 1965 | 6.5 | 5.3 | 3.61 | - |
| 1966 | 4.7 | 6.8 | 3.22 | 3.72 |
| 1967 | 5.4 | 3.4 | 3.87 | - |
| 1968 | 4.5 | 3.3 | 2.93 | 3.13 |
| 1969 | 3.4 | 3.2 | 2.68 | 2.53 |
| 1970 |  |  | 2.82 | 2.99 |

Table 9. Abundance Indices for red hake in New England waters based on catch per haul statistics from USSR fleet ${ }^{1}$, and joint USA-USSR groundfish surveys ${ }^{2}$.

|  |  |  | RED |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | So. N (str | $\begin{aligned} & \text { ngland } \\ & 1-12 \text { ) } \end{aligned}$ |  | $\begin{aligned} & \text { George } \\ & \text { (strata } \end{aligned}$ | $\begin{aligned} & 3 \text { 3ank } \\ & 3-25) \end{aligned}$ |  |
|  | Fleet | Surv |  | Fleet | Surv |  |
| Year | USSR | USSR | USA | USSR | USSR | USA |
| 1965 | 2.44 |  | 1.85 | 1.32 |  | 0.78 |
| 1966 | 1.69 |  | 1.48 | 2.39 |  | 0.72 |
| 1967 | 0.96 | 2.07 | 1.05 | 0.96 | 0.84 | 0.46 |
| 1968 | 0.56 | 1.88 | 0.79 | 0.62 | 1.79 | 0.64 |
| 1969 | $1.75{ }^{3}$ | 2.20 | 1.18 | - | 1.03 | 0.85 |
| 1970 | - | 2.36 | 1.35 | - |  | 0.44 |

1 Catch per haul hour for red hake from ICNAF research document 70/39 by Richter, for "stocks I and II" which correspond approximately to strata sets $13-25$ and 1-12 respectively.

2 Stratified mean catch per haul (pounds, natural log scale).
3 Estimate provided on graph by Dr. Noskov at Working Group in Copenhagen, January 1971.

Table 10. Stratified mean catch per haul $\left(\log _{e}\right.$ pounds) of selected species in southern New England (strata 1-12). USA and USSR joint surveys.

RED HAKE

| Year | Strat. mean |  | Varlance |  | S.D. |  | S.D./mean |  | No. hauls |  |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
|  | U.S. | USSR | U.S. | USSR | U.S. | USSR | U.S. | USSR | U.S. | USSR |
|  | 1.05 | 2.07 | .0229 | .0554 | .1513 | .2354 | .14 | .11 | 65 | 40 |
| 1968 | 0.79 | 1.88 | .0238 | .0421 | .1543 | .2052 | .20 | .11 | 62 | 46 |
| $1969^{2}$ | 1.18 | 2.20 | .0236 | .0760 | .1536 | .2757 | .13 | .12 | 66 | 42 |
| 1970 | 1.35 | 2.36 | .0199 | .0314 | .1411 | .1772 | .10 | .08 | 64 | 56 |

SILVER HAKE

|  | U.S. | USSR | U.S. | USSR | U.S. | USSR | U.S. | USSR |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
|  |  |  |  |  |  |  |  |  |
| 1967 | 1.63 | 2.64 | .0202 | .0579 | .1421 | .2406 | .09 | .09 |
| 1968 | 1.80 | 3.62 | .0155 | .0404 | .1245 | .2010 | .07 | .06 |
| 1969 | 1.20 | 3.38 | .0142 | .0676 | .1192 | .2600 | .10 | .08 |
| 1970 | 1.35 | 3.71 | .0125 | .0273 | .1118 | .1652 | .08 | .04 |

YELLOWTAIL

|  | U.S. | USSR | U.S. | USSR | U.S. | USSR | U.S. | USSR |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 1967 | 2.25 | 1.70 | .0270 | .0514 | .1643 | .2267 | .07 | .13 |
| 1968 | 2.03 | 1.78 | .0380 | .0592 | .1949 | .2433 | .10 | .14 |
| 1969 | 2.00 | 1.75 | .0361 | .0708 | .1900 | .2661 | .10 | .15 |
| 1970 | 2.12 | 1.50 | .0420 | .0657 | .2049 | .2563 | .10 | .17 |

${ }^{1}$ No. hauls in stratum 10; sampling in strata 9, 11, 12 restricted to area west of $70^{\circ} \mathrm{W}$.
224.6 m trawl used by USSR vessel in 1969; 27.1 m trawl used by USSR vessels in all other surveys.

## Comparisons between USA and USSR survey indices

The larger USSR trawls appear to have up to 5 times the fishing power of the USA survey trawl for some species, as indicated by trawl comparison experiments and joint surveys since 1967 (Hennemuth, 1968; Grosslein and Sauskan, 1970). The question arises whether there is any significant relation between fishing power and accuracy of the abundance indices. We have been particularly concerned about the possibility that in the case of species for which our U.S. gear has relatively low fishing power (e.g. silver hake), relatively minor changes in behavior and especially vertical distribution might change avilability enough to obscure real changes in abundance. So far there is no clear evidence of any such disadvantage with the smaller trawl from the standpoint of accuracy.

Table 11. Stratified mean catch per haul $\left(\log _{e}\right.$ pounds) of selected species in Georges Bank (strata 13-25). USA and USSR joint surveys.

| COD |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| YEAR | Mean |  | Variance |  | S. D. |  | S.D./mean |  | No. hauls |  |
|  | USA | USSR | USA | USSR | USA | USSR | USA | USSR | USA | USSR |
| 1967 | 1.74 | - | . 0473 | - | . 2175 | - | . 12 | - | 67 | - |
| $1968{ }^{1}$ | 1.04 | 1.19 | . 0319 | . 0400 | . 1786 | . 2000 | . 17 | . 17 | 69 | 49 |
| $1969{ }^{2}$ | 1.32 | 1.59 | . 0254 | . 0178 | . 1594 | . 1334 | . 12 | . 08 | 73 | 37 |
| $1970{ }^{3}$ | 1.35 | 0.87 | . 0332 | . 0367 | . 1822 | . 1916 | . 13 | . 22 | 70 | 31 |
| HADDOCK |  |  |  |  |  |  |  |  |  |  |
|  | - USA | USSR | USA | USSR | USA | USSR | USA | USSR |  |  |
| 1967 | 2.45 | 1.07 | . 0521 | - | . 2282 | - | . 09 | - |  |  |
| 1968 | 1.15 | 1.07 | . 0296 | . 0248 | . 1720 | . 1667 | '. 15 | . 16 |  |  |
| 1969 | 1.10 | 1.65 | . 0215 | . 0649 | . 1466 | . 2548 | . 13 | . 15 |  |  |
| 1970 | 1.35 | 0.57 | . 0345 | . 0285 | . 1857 | . 1688 | . 14 | . 30 |  |  |
| YELLOWTAIL |  |  |  |  |  |  |  |  |  |  |
|  | USA | USSR | USA | USSR | USA | USSR | USA | USSR |  |  |
| 1967 | 1.32 | - | . 0277 | - | . 1664 | - | . 13 | - |  |  |
| 1968 | 1.40 | 1.01 | . 0382 | . 0340 | . 1954 | . 1844 | . 14 | . 18 |  |  |
| 1969 | 1.35 | 1.91 | . 0252 | . 0615 | . 1587 | . 2480 | . 12 | . 13 |  |  |
| 1970 | 0.96 | 1.80 | . 0204 | . 0878 | . 1428 | . 2963 | . 15 | . 16 |  |  |

1 No hauls by USSR vessel in 1968 in stratum 25 and only one haul each in strata 15,17 and 22.
224.6 m trawl used by USSR vessel in 1969; 27.1 m trawl used by USSR vessels in all other surveys.

3 No hauls in strata $23-25$ by USSR vessel in 1970.

With respect to sampling errors we find that variances of stratified means are fairly comparable for the two sizes of gear, and they appear to be rather independent of fishing power differentials. For example the fishing power differential is large for red hake but quite small for cod, haddock and yellowtail, and yet variances are quite similar for all these species and both types of trawl in New England waters (Tables 10, 11). Generally similar results were obtained in the 1970 surveys off Nova Scotia (Table 12).
Table 12. Stratified means (catch per haul, $\log _{\mathrm{e}}$ pounds) and measures of precision for selected species
in 1970 surveys in Division 4 X (sampling strata $31,32,41-49$ ); USSR ( 34 hauls), USA ( 45 hauls).

| Species | Stratified mean |  | Variance |  | S.D. |  | Mean $\pm 2$ S.D. |  | Factordifference |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | USA | USSR | USA | USSR | USA | USSR | USA | USSR | USA | USSR |
| Cod | 1.20 | 0.55 | . 0690 | . 0416 | . 2627 | . 2040 | .68-1.72 | . 15-. 95 | 2.9 | 2.2 |
| Haddock | 2.05 | 1.81 | . 0231 | . 0574 | . 1520 | . 2396 | 1.75-2.35 | 1.33-2.29 | 1.8 | 2.6 |
| Am. Dab | 0.83 | 0.21 | . 0214 | . 0041 | . 1463 | . 0640 | .53-1.13 | .09-. 33 | 1.8 | 1.3 |
| Yellowtail | 0.35 | 0.11 | . 0227 | . 0041 | . 1507 | . 0640 | .05-.65 | 0-. 23 | 1.8 | 1.3 |
| Silver hake | 0.85 | 1.02 | . 0216 | . 0941 | . 1470 | . 3068 | .55-1.15 | . $40-1.64$ | 1.8 | 3.4 |

With respect to comparability of trends we find very close correspondence between the indices for red hake in southern New England, in both direction and magnitude (Fig. 6). The correspondence is not as good for silver hake in the southern New England area but the direction of change is the same from year to year (Fig. 7). More variability between the two indices was encountered for both red and silver hake on Georges Bank, where they were less abundant, but again the correspondence was better for red hake (Figs. 6, 7). These data are difficult to interpret because the USSR trawl used in 1969 was not the same as that used in 1967 and 1968; the 1969 gear obviously had greater fishing power. Furthermore sampling with the USSR trawls on Georges Bank was less intensive (and not consistent in the sense that all areas were not fished each year) because of more frequent tear-ups and necissity of avoiding rough bottom.

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21. Commercial fishes taken on the shrimp grounds of the Northwest Atlantic ${ }^{1}$
by B. Fontaine
ISTPM, St. Pierre et Miquelon

## Introduction

Whether they are in coastal or open waters, shrimp grounds rarely provide exclusive catches of shrimp. Fish which appear in trawl hauls made on such grounds, at least those which are of commercial value, are among the species common to the whole Northwest Atlantic.

This paper attempts to show the importance and species composition in trawl hauls made by research vessels which provide a good yield of deep shrimp, Pandalus borealis.

The grounds described here are divided into different sections, starting from Georges Bank to the West Greenland coast; either coastal, such as the Burgeo trench or Esquiman Channel or offshore grounds as for all other sections.

## Fishing areas

To simplify matters, the grounds which provided a good shrimp yield are designated by the name of the most general area where they are located. The position and characteristics of these grounds for each section are specified below.

## 1. Georges Bank section

The deep shrimp was taken in the deep located to the west of Georges Bank; this deep is indicated on the charts as Georges Basin.

In May and June 1969, seven trawl hauls, made between 145 and 195 m , provided an average shrimp yield of $38 \mathrm{~kg} / \mathrm{hr}$ of trawling.

## 2. Nova Scotia section

The shrimp grounds are located in certain trenches dividing the banks, i.e., the trench of the LaHave Bank, that of Emerald Bank, of Canso Bank, etc.

In March and April 1967, eight trawl hauls, spread over these deeps between 170 and $250^{\circ} \mathrm{m}$, gave an average shrimp yield of $40 \mathrm{~kg} / \mathrm{hr}$ of fishing.

## 3. South Newfoundland section

This region comprises the complex formed by the Laurentian Channel, its slopes and canyons, i.e., the deeps located between the banks of the northeast region of Nova Scotia Shelf. The Burgeo Trench is dealt with separately.

Eight hauls, made in May 1970 at $190-320 \mathrm{~m}$, provided an average shrimp yield of $22.5 \mathrm{~kg} / \mathrm{hr}$ of fishing.

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## 4. Burgeo Trench

This well-defined trench has been fished as a typical coastal grounds, as is the Esquiman Channel. It extends along the south Newfoundland coast, from La Poile Bay to Ramea Islands, Inside Rose Blanche and Burgeo Banks. The soundings are in the order of $220-250 \mathrm{~m}$.

Ten hauls, made in December 1970, gave average shrimp yields of $42 \mathrm{~kg} / \mathrm{hr}$ of fishing.

## 5. Esquiman Channe1

This deep is located in the Gulf of St. Lawrence, along the west coast of Newfoundland; it is about 43 miles wide at the level of Anticosti Island and gradually decreases, disappearing near the Newfoundland coast opposite Port-au-Choix.

Six hauls in December 1970, at depths of 225-250 m, ylelded shrimp at an average of $83 \mathrm{~kg} / \mathrm{hr}$ of fishing.

## 6. Labrador section

The important grounds are located off the Labrador coast and the eastnortheast coast of Newfoundland, particularly northeast of Hopedale and off Bonavista Bay.

For eleven trawl hauls made at $350-400 \mathrm{~m}$, shrimp catches averaged about $30 \mathrm{~kg} / \mathrm{hr}$ of fishing.

## 7. West Greenland section

In July and August 1970, R/V Thalassa made good catches in different areas of the west coast of Greenland. These consisted of deeps located on the eastern edge of Fyllas Bank off Godthaab, east-southeast of Banana Bank, southsoutheast and north of Lille Hellefiske Bank.

Depths varied according to zone, from $250-550 \mathrm{~m}$; for 17 traw1 hauls the average deep shrimp yield was $40-80 \mathrm{~kg} / \mathrm{hr}$ of fishing.

## Method of capture

Two types of gear were used during the different cruises. For the south of Newfoundland, Burgeo Trench and Esquiman Channel sections, a shrimp trawl "balloon type", with a $60-\mathrm{mm}$ mesh in the wing, back and belly, and 40 mm in the "amorce" and codend (stretch mesh) was used. For all other sections, a bottom trawl of the "Lofoten" type, with a bobbin groundrope, $140-\mathrm{mm}$ mesh in the wings, back and belly and 50 mm in the codend was used. In the study of the Greenland section, this codend was lined wth a $30-\mathrm{mm}$ mesh net.

## Results

The results are given in three tables. Table 1 gives the number of hauls per 100 trawl hauls on the shrimp beds, in which each fish species of commercial value was observed. This table shows the rate of occurrence of a species and not the quantity of animals caught; the crosses indicate rates lower than

Table 1．Abundance index（percentage of hauls containing each fish species） of commercial species taken on the shrimp beds．Crosses（＋）indicate an index less than 30.

|  |  | W1 0 0 0 0 0 0 0 2 |  |  |  | $\begin{aligned} & \text { H } \\ & \text { 菏 } \\ & \text { 0⿹\zh26灬 } \end{aligned}$ |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Squalus acanthias | 57 | ＋ | ＋ | － | － | － | － |
| Raia radiata | 100 | 75 | 88 | 100 | 100 | 90 | ＋ |
| Raia senta | 71 | 75 | 75 | 100 | 83 | － | － |
| Raia spinicauda | － | － | － | － |  | ＋ | － |
| Raia laevis | ＋ | － | ＋ | － | － | － | － |
| Raia erinacea | － | － | ＋ | － | － | － | 70 |
| Clupea harengus harengus | 85 | ＋ | 75 | 100 | 100 | － | － |
| Mallotus villosus villosus | － | 50 | ＋ | ＋ | ＋ | 30 | ＋ |
| Argentina silus | ＋ | 37 | 37. | 60 | － | ＋ | － |
| Merluccius bilinearis | 100 | 37 | 62 | ＋ | － | － | － |
| Brosme brosme | 42 | － | － | － | － | － | － |
| Urophycis teruis | 71 | 75 | 88 | 100 | $+$ | － | － |
| Urophycis chesteri | － | 37 | 62 | － | － | － | － |
| Gadus morhua morhua | 85 | 100 | 37 | 100 | 100 | 100 | 65 |
| Gadus morhua ogac | － | － | － | － | － | － | ＋ |
| Boreogadus saida | － | － | － | － | － | － | ＋ |
| Melanogrammus aeglefinus | 85 | 87 | ＋ | ＋ | ＋ | － | － |
| Pollachius virens | 57 | 87 | － | 40 | － | － | － |
| Molva dipterygia | － | － | － | － | － | － | ＋ |
| Macrumis sp． | ＋ | 35 | 50 | － | － | 70 | 30 |
| Anarhichas lupus | － | ＋ | ＋ | － | ＋ | 80 | 40 |
| Anarhichas minor | － | ＋ | － | － | － | 90 | 30 |
| Sebastes sp． | 100 | 100 | 100 | 100 | 100 | 100 | 85 |
| Hippoglossoides platessoides | 100 | 100 | 100 | 100 | 100 | 80 | 95 |
| Hippoglossus hippoglossus | － | 50 | － | － | 66 | － | － |
| Reinhardtius hippoglossoides | ＋ | 87 | 62 | 100 | 100 | 90 | 95 |
| Glyptocephalus cynoglossus | 100 | 100 | 100 | 100 | 100 | 80 | － |
| Lophius americanus | 57 | 50 | 88 | 80 | ＋ | － | － |

30 which，according to us，represented an accidental occurrence．Table 2 gives the average weight in kg of different commercial fish caught with 100 kg of deep shrimp．Table 3 indicates the size composition of catches for the five most important commercial species．These tables require，however，explanation which will be given when the different species considered are reviewed．

Sharks appeared in some hauls in the Georges Bank section in very small numbers，or as isolated individuals．Rays are almost always present on shrimp

Table 2. Weight (in kilos) of the different comercial fishes taken with 100 kg of deep shrimp.

|  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Dogfish (Chien de mer) | 14 | - | - | - | - | - | - |
| Rays (Rales) | 154 | 87 | 103 | 45 | 37 | 67 | 12 |
| Herring (Hareng) | 31 | - | 30 | 58 | 56 | - | - |
| Argentine | - | - | 62 | 32 | - | - |  |
| Silver hake (Merlu) | 41 | 28 | 36 | 1 | - | - | - |
| Cusk (Brosme) | 15 | - | - | - | - | - | - |
| Red hake (Morue barbue) | 88 | 110 | 154 | 165 | 1 | - | - |
| Cod (Morue) | 253 | 390 | 29 | 62 | 218 | 322 | 28 |
| Haddock (Eglefin) | 84 | 135 | - | 1 | 1 | - | - |
| Pollock (Lieu noir) | 54 | 53 | - | 2 | - | - | - |
| Grenadiers (Rats) | - | - | - | - | - | 200 | - |
| Wolffish (Loups) | - | 27 | - | - | - | 105 | 24 |
| Redfish (Rougets) | 175 | 867 | 940 | 2703 | 1044 | 939 | 21 |
| American plaice <br> (Balai) | 131 | 208 | 62 | 15 | 24 | 59 | 125 |
| Halibut (Flétan) | - | 8 | - | - | 10 | - | - |
| Greenland halibut (Flétan noir) | - | 13 | 27 | 43 | 126 | 74 | 125 |
| Witch (Sole grise) | 32 | 272 | 83 | 272 | 72 | 54 | - |
| Angler (Lotte) | 19 | 5 | 35 | 21 | - | - | - |
| Total | 1091 | 2203 | 1561 | 3420 | 1589 | 1820 | 335 |

grounds; only the species Raia radiata is of some importance by weight, except in Greenland where it is replaced by Raia erinacea. Raia senta is of ten represented, but always by a very limited number of individuals.

Herring, even if not of great importance by weight, were caught more or less regularly in certain sections. In the Burgeo Trench, the maximum catch per hour was 86 kg . In the absence of pelagic detection, it is supposed they were dispersed on the bottom.
Table 3. Length composition of the catch of five of the most important coumercial species.

| Species <br> Fishing areas | Georges Bank | Nova Scotia | Labrador | Greenland | South of Newfoundland | Burgeo Trench | Esquiman Channel |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Cod $\quad$: range of lengths (cm) <br>  <br>  <br>  <br> $:$ less than 40 cm <br> between 40 and 60 cm | $\begin{gathered} 53-132 \\ 0 \% \\ 10 \% \end{gathered}$ | $\begin{gathered} 39-92 \\ 5 \% \\ 66 \% \end{gathered}$ | $\begin{gathered} 19-128 \\ 31 \% \\ 55 \% \end{gathered}$ | $\begin{gathered} 35-99 \\ 0 \% \\ 15 \% \end{gathered}$ | $\begin{gathered} 20-95 \\ 20 \% \\ 67 \% \end{gathered}$ |  |  |
| Redfish $\left.\begin{array}{l}\text { : range of lengths (cm) } \\ \\ :\end{array}\right)$ less than 25 cm <br>  between 25 and 35 cm | $\begin{array}{r} 14-42 \\ 77 \% \\ 18 \% \end{array}$ | $\begin{aligned} & 7-38 \\ & 51 \% \\ & 44 \% \end{aligned}$ | $\begin{gathered} 7-60 \\ 20 \% \\ 26 \% \end{gathered}$ | $\begin{aligned} & 7-43 \\ & 70 \% \\ & 25 \% \end{aligned}$ | $\begin{gathered} 11-43 \\ 83 \% \\ 15 \% \end{gathered}$ | $\begin{array}{r} 7-43 \\ 87 \% \\ 9 \% \end{array}$ |  |
| American : range of lengths (cm) <br> plaice <br> $:$ less than 30 cm <br> between 30 and 40 cm  | $\begin{array}{r} 13-68 \\ 40 \% \\ 46 \% \end{array}$ | $\begin{gathered} 10-52 \\ 61 \% \\ 32 \% \end{gathered}$ | $\begin{array}{r} 13-56 \\ 72 \% \\ 20 \% \end{array}$ | $\begin{gathered} 11-50 \\ 48 \% \\ 44 \% \end{gathered}$ | $\begin{aligned} & 15-57 \\ & 66 \% \\ & 19 \% \end{aligned}$ | $\begin{array}{r} 14-66 \\ 79 \% \\ 11 \% \end{array}$ | $\begin{gathered} 10-63 \\ 87 \% \\ 7 \% \end{gathered}$ |
| Greenland : range of lengths (cm) halibut $\left.\begin{array}{l}\text { less than } 40 \mathrm{~cm} \\ :\end{array}\right)$ between 40 and 60 cm | - | - | $\begin{array}{r} 13-84 \\ 43 \% \\ 44 \% \end{array}$ | $\begin{gathered} 12-78 \\ 79 \% \\ 10 \% \end{gathered}$ |  | $\begin{array}{r} 21-61 \\ 48 \% \\ 47 \% \end{array}$ |  |
| Witch : range of lengths $(\mathrm{cm})$ <br>  <br> $:$ <br>  <br> $:$ between 30 and 40 cm | $\begin{array}{r} 35-60 \\ 0 \% \\ 15 \% \end{array}$ | $\begin{gathered} 21-55 \\ 5 \% \\ 76 \% \end{gathered}$ | $\begin{array}{r} 26-66 \\ 13 \% \\ 27 \% \end{array}$ | - |  | $\begin{array}{r} 13-58 \\ 32 \% \\ 41 \% \end{array}$ | $\begin{gathered} 17-49 \\ 42 \% \\ 30 \% \end{gathered}$ |

Argentine were only caught on the shrimp grounds in the south of Newfoundland in May, and in the Burgeo Trench in December.

Silver hake were caught on Georges Bank. The catches consisted of specimens $12-52 \mathrm{~cm}$ in total length, and about $30 \%$ were less than 15 cm .

Cusk seem almost accidental on Georges Bank as they are represented only by isolated individuals.

Red hake, Urophycis tenuis, occur in catches from Georges Bank to the Burgeo Trench (Table 2). This species is represented by a limited number of Individuals whose sizes vary from $22-98 \mathrm{~cm}$, and only $31 \%$ are less than 40 cm total length.

Cod caught with the deep shrimp in the Burgeo Trench and Esquiman Channel were not measured; individuals of less than about 40 cm were weighted separately.

In the Burgeo Trench, the numbers of cod caught were very small but were always of a large size. Thus, one of the hauls provided two cod of 132 and 121 cm , weighing 19 and 17 kg respectively. The same observations are valid for the Georges Bank and Greenland sections.

In the Esquiman Channel, cod less than 40 cm formed $20 \%$ of total weight. This section can be compared with those of Nova Scotia, south of Newfoundland, and Labrador.

Redfish caught in Esquiman Channel were not measured; individuals less than 25 cm made up approximately $35-40 \%$ of total weight. Table 3 shows that the Labrador section and, to a lesser degree, the Nova Scotia section provided in association with the shrimp, good commercial catches of redfish. From Nova Scotia to Labrador, redfish in the catches constitutes a problem for the shrimp fishery by being a nuisance and by damaging the shrimp which are more fragile than the fish.

American plaice were present in nearly all hauls with smaller quantities being taken in coastal deeps. Approximately $90 \%$ of the individuals caught were less than 40 cm in length.

Greenland halibut were not measured in the areas south of Newfoundland, nor in Esquiman Channel or Greenland, where it has some importance in weight.

Witch, rare in Greenland where only one specimen was taken, are well represented in hauls from Nova Scotia and from the Burgeo Trench. Individuals of less than 30 cm are more numerous on coastal than offshore grounds. Catches from Newfoundland were not measured.

American angler were present fairly regularly in catches from Georges Bank to Burgeo Trench. However, the number of individuals caught by trawl was always small.

## Conclusion

Shrimp grounds located off the west coast of Greenland are of particular interest as shown clearly in Table 2. For all other sections, fishing of deep
shrimp involves taking a large quantity of fish of commercial value. These catches are unfortunately made up of a large number of individuals of small non-commercial size. This fact is particularly clear in the case of American plaice and redfish. The minimum sizes of fish caught are identical so we use a trawl with codend of 50 or 30 mm (stretch mesh); however, using a shrimp trawl catches a greater proportion of fish of smaller size.

The coastal shrimp grounds and, to a lesser degree, those offshore in the Northwest Atlantic are the areas of concentration of the young of certain species.

Data on fish catches accompanying the shrimp catch are accurate but little infqrmation exists about the importance of the quantity of rejected fish. A study of the influence of small mesh trawls on fish populations encountered on offshore shrimp grounds will be made and it is hoped that the coastal shrimp grounds can also be investigated.
22. Summary report of the ICNAF NORWESTLANT surveys, $1963^{1}$

by E. Smidt<br>Greenland Fisheries Investigations<br>Charlottenlund, Denmark

The NORWESTLANT surveys was the first large scale international scheme of environmental research sponsored by ICNAF. Not less than 8 member countries took part with 10 research vessels during April-July 1963 in the northwestern Atlantic. Three surveys were carried out, namely NORWESTLANT 1 in April, 2 in May-June, and 3 in July. The surveys were carefully planned during several meetings and a detalled Guide Book was prepared. Mr Arthur Lee (Lowestoft) was the enthusiastic and dynamic organizer.

The main aim of the surveys was to establish the distribution and drift of the eggs and larvae of cod and of the larvae of redfish, in relation to specific environmental factors in the area. Other fish species were also included. In addition, observations were made on adult fish and marine mamals.

Figure 1 shows the NORWESTLANT survey area and the grid stations occupied during NORWESTLANT 2. Much meteorological, hydrographical, and biological data was collected by the research vessels, and also by weather ships. All material collected was worked up by the institutions in the different collecting countries. The results were then sent to scientific coordinators who prepared the various reports.

This paper reviews the results which are relevant to the main aim of the surveys, and which are especially important to Subarea 1. The results were published in ICNAF Special Publication No. 7, Part I (Text) and II (Atlas) in 1968 and distributed in 1969. The original Oceanographical and Biological Data were distributed as Parts III and IV respectively.
I. Meteorological conditions and physical and chemical oceanography

Weather conditions were abnormal in 1963 and the surveys were carried out in particularly stormy weather which influenced the stability of the water layers. Ice condtions, however, were nearly normal along both the east and west coast of Greenland.

Surface water temperature anomalies were calculated in different areas in the different months. In the area north of $55^{\circ} \mathrm{N}$ lat., positive anomalies in April ranged from about $1^{\circ} \mathrm{C}$ off West Greenland to $0.4^{\circ} \mathrm{C}$ off East Greenland. South of $55^{\circ} \mathrm{N}$ lat., anomalies were slightly negative. In the months following the anomalies were generally decreasing, especially off West Greenland.

The stability of the surface water layers, which is important to primary production, was determined by Mr Arthur Lee as density differences between the surface and 50 m . During NORWESTLANT 1 (April) there was 1ittle stability over the area as a whole, except in certain zones where cold, low salinity water covered warmer, high salinity water. This was most extensive off West Greenland where north to northeast winds had driven the low salinity coastal water westwards over water of Irminger Current origin. During the months following the zones of stability expanded until in July the whole area had a certain amount of stability. However, off West Greenland, stability had decreased in July.

[^18]

Fig. 1. NORWESTLANT 2., stations and sections. (Chart 32 in the NWL publication.

The distribution of phosphates, silicates, and dissolved oxygen from $0-$ 100 m was in good agreement with the stability conditions. In April there was very little well-defined stratification of phosphates and silicates except in Davis Strait. Here, a high percentage of oxygen saturation was also found.

## II. Phytoplankton production

Investigations on the phytoplankton were carried out using various methods of which the most comprehensive was to give a picture of the seasonal and geographical distribution of the total phytoplankton as micrograms of carbon per litre, $\mu \mathrm{gC} / \ell$. Phytoplankton from water samples taken at all stations from $10-\mathrm{m}$ depth was treated by Dr M. Gillbricht (Hamburg). The main results are presented in charts showing the distribution during each survey.

During NORWESTLANT 1 (Fig. 2) phytoplankton was only abundant west of Greenland with a maximum of about $50 \mu \mathrm{gC} / \ell$ at $64^{\circ} \mathrm{N}$ lat. In almost all other areas production was less than $5 \mathrm{\mu g} / \ell$. This early (April) phytoplankton


Fig. 2. Phytoplankton at 10 m in $\mu \mathrm{gC} / \ell$ during NORWESTLANT 1. Contour levels at 5, 10, 20, 50, and $100 \mu \mathrm{gC} / \ell$. (Chart 180 in the NWL publication.)
maximum at West Greenland is in agreement with the early stability of the water masses in the area. During NORWESTLANT 2, the phytoplankton was at its maximum in most of the area, as the water had now become generally stabilized. During NORWESTLANT 3 there was an overall decline, especially at West Greenland.

Figure 3 shows the pattern of phytoplankton distribution during NORWESTLANT 3 (July), which is in good agreement with the distribution of the gross production around Greenland in July-August of 1954, as measured by the carbon-14 technique during the cruise of Dana. Figure 4 (after Steemann Nielsen, 1958) shows a high rate of production where fronts between two current systems are found and where deeper water ascends to the surface. Such productive areas were found: 1) at the front between the Atlantic Current and the water from the Irminger Sea; 2) outside the East Greenland Current; 3) at the front between the Labrador Current and the oceanic water in the centre of Davis Strait; and 4) off the West Greenland coast. An illustration of the hydrographic mechanism is given in the NORWESTLANT Report by Mr F. Hermann. The front area between the light water component of the West Greenland Current and the heavy Irminger water component is an area with horizontal turbulence causing vertical water movements. A section over Fyllas Bank (Fig. 5) in July shows the density conditions, the current velocity with a maximum in the front over the western slope of the Bank, and the upwelling of phosphate-rich and oxygen-poor water.


Fig. 3. Phytoplankton at 10 m in $\mu \mathrm{gC} / \ell$ during NORWESTLANT 3. Contour levels as in Fig. 2. (Chart 183 in the NWL publication.

In addition to the above-mentioned investigations on the distribution of phytoplankton in the whole area, less extensive research on the primary production was made by chemical methods between southwestern Greenland and Labrador from USSR vessels (M.V. Fedosov and I.A. Ermachenko). Phytoplankton abundance was estimated from the silks of the continuous plankton recorder surveys (G.A. Robinson). Estimates of chlorophyll-a concentration were made by 4 countries (J.H. Steele). The results of these additional investigations seem to be in agreement with those for the whole area. Further, the continuous plankton records allowed comparison of conditions in 1963 to those of other years from 1958 to 1964 for all areas, except West Greenland. Production off West Greenland was already abundant in May 1963, about a month earlier than in 1962, the only year for. which data are available for comparison from that area. Elsewhere, the production was relatively small, presumably due to the late stabilizing of the water layers, but it was close to its normal timing.
III. The zooplankton

Much zooplankton material was collected during the surveys using the different nets described in the Guide Book, namely, 1) Hensen net: diameter


Fig. 4. Gross production in the waters around Greenland in July-August 1954. The results from similar investigations in 1955, 1956, and 1957 agree in principle fully with those from 1954. (After Steemann Nielsen, 1958).

72 cm , hauled vertically from 100 m to the surface, or from a few metres off the bottom at stations less than 100 m in depth; 2) stramin net: diameter 2 m , hauled obliquely from about 50 m to the surface at 1.5 knots, duration about 30 min ; 3) Icelandic High Speed Sampler (IHSS): an aluminum body 1 m in length with a sampling aperture 90 mm in diameter, with a silk or nylon gauze bag inside, hauled obliquely from about 50 m to the surface at 5 knots, duration about 30 min .

The most complete coverage of the stations on all surveys was with the Hensen net and the stramin net, while the IHSS was only used regularly by few vessels. In addition to these standard nets, the Nansen net and the Helgoland larva net were used in some cases.

Displacement volumes were measured on board the vessels, while most of the sorting and counting was done at the different laboratories. At a meeting in October 1963 it was agreed which species caught by stramin net and IHSS


Fig. 5. Section over Fyllas Bank, 6-7 July 1963 (NORWESTLANT 3). A - Sigmatt; B - Currents ( $\mathrm{cm} / \mathrm{sec}$ ) normal to the section, all north-going; $C-\mathrm{PO}_{4}-\mathrm{P}$ (ug at/ $\ell$ ); D - Dissolved oxygen content (percent saturation); E - Vertical water movements. (Fig. 9 in the NWL publication.)
should be given priority for determining their total numbers. Animals in samples from vertical hauls were recorded as numbers under $1 \mathrm{~m}^{2}$ of the sea surface within the top 100 m . The material was coordinated by Mr V. Bainbridge (Edinburgh) and Mr J. Corlett (Lowestoft), who prepared distribution chart:s, made figures, and prepared the reports. In addition, continuous plankton records were published (Mr R.S. Glover and Mr G.A. Robinson, both Edinburgh).

In the Irminger and Labrador Seas the species composition showed considerable uniformity with Calanus finmarchicus as the dominant member of the zooplankton, followed by Euchaeta nomegica, Thysanoessa longicaudata, Conchoecia obtusata, Spiratella retroversa, Aglantha digitale and Eukrohnia homata. In the central part of Davis Strait, the plankton was somewhat the same but, in addition to Calonus finmarchicus, there were substantial numbers of c. glacialis

$\square$ Eggs Naupwlin Copapodites

Fig. 6. The mean biomass of various food organisms in redfish larvae of all length-groups from 6 to 13 mm during NORWESTLANT 1-3 (April-July). (Fig. 52 in the NWL publication.)
and $C$. hyperboreus. In the Icelandic coastal waters there was a greater diversity of species. Quantities of zooplankton were lowest in the cold waters of the East and West Greenland Currents.

In this brief review it is impossible to go into detail about all the selected species. Only animals important to the feeding and predation of fish larvae are mentioned hereafter.


Fig. 7. The mean biomass of various food organisms in cod larvae of all length-groups from 3 to 10 mm during NORWESTLANT 2 (May-June) at Iceland and at West Greenland. (Fig. 53 in the NWL publication.)

1. The feeding of cod and redfish larvae

An important part of the NORWESTLANT surveys was an excellent comparative study of the food of redfish and cod larvae by Mr V. Bainbridge and Mr B.J. McKay (both Edinburgh) over a wide area.

It was shown that Calanus finmarchicus is by far the most important food organism for both redfish and cod larvae.

Recently extruded redfish larvae in the Irminger Sea are principally feeding on Calanus eggs in April and May, while older larvae taken in June and July in the Irminger Sea and Davis Strait were eating mainly nauph1ii and copepodites of Calanus (Fig. 6).

The food of cod larvae is shown in Fig. 7. At West Greenland the larvae feed almost exclusively on different stages of Calanus, while in the Icelandic area the diet is more varied consisting also of other copepods, euphausids, Evadne, and some other organisms. The differences in the diet of cod larvae from the two areas reflect the diversity of the neritic zooplankton off Iceland compared with the more restricted species composition of the zooplankton in the West Greenland waters. It is also noted that cod larvae from the Icelandic area contained about twice the amount of food found in larvae from West Greenland waters, a difference which seems to be correlated with the availability of food in the plankton. Further in this connection, it should be noted that cod larvae off Iceland and in Denmark Strait seem to have a higher growth rate than at West Greenland.

Larvae of both redfish and cod showed a similar diurnal feeding pattern. The amount of food found in the gut decreased throughout the night to a minimum about sunrise.

In addition to the NORWESTLANT samples, a large number of cod larvae collected off West Greenland in 1950, 1957, 1958, and 1961 were also examined. In all but one of 18 samples, various stages of Calanus constituted more than $95 \%$ by weight of the food of the larvae.

## 2. Calanus finmarchicus

"Since Calanus is both the dominant herbivore and the principal food consumed by cod and redfish larvae in the seas around Greenland, the seasonal development of this species will be considered in detail. Average numbers of each copepodite stage per $\mathrm{m}^{2}$ of sea surface have been calculated for each survey in seven standard areas using data from the Hensen-net samples." The standard areas are shown in Fig. 8, averages obtained are shown as histograms in Fig. 9.

It is known (Marshall and Orr, 1955; Østved, 1955) that Calanus overwinters in deep water as late copepodite stages, and in cold waters there is only one generation per year. During spring, the copepodites ascend towards the surface, moult to become adults, spawn, and die soon afterwards.

Figure 9. shows that during April the Calanus population in the top 100 m was represented mainly by adults; while early copepodites were scarce. In June all three western areas had high numbers of early copepodites, but in the eastern oceanic areas, the Irminger Sea, there was a scarcity of young stages, and a persistence of adults in the northern Irminger Sea, which indicates that the main spring spawning had been delayed in these areas. In July early copepodites were reduced in numbers in the western areas, while they had become numerous in the eastern areas. In the western, as well as the eastern, areas, the stage composition in July indicates that the rate of development must have


Fig. 8. Standard areas for the zooplankton surveys. (Chart 192 in the NWL publication.)
varied throughout the survey areas from north to south with younger stages predominant in the north and older stages predominant in the south, possibly due to temperature conditions.

The late spawning of Calanus in the Irminger Sea may be explained by the late outburst of phytoplankton caused by the late stabilization of the surface water in the area. It is known that the reproduction of Calanus depends very closely on the food available (Marshall and Orr, 1964).

Seasonal and geographical variation in the development of other plankton animals, namely euphausiacea, Spiratelle retroversa, certain medusae, and chaetognatha; was studied, but it is not possible to go into detail here.

## 3. Potential predators of cod eggs and larvae

"It is generally accepted that the larval phase in the life history of the fish is critical and two of the main causes of larval mortality are thought to be starvation and predation, both of which will be directly influenced by the abundance and composition of the zooplankton.


Fig. 9. Average numbers of the adult and copepodite stages of Calanus under $1 \mathrm{~m}^{2}$ in the standard areas shown in Fig. 8, during NORWESTLANT 1-3. (Fig. 26 in the NWL publication.)


Fig. 10. Distribution and abundance of cod eggs and larvae during NORWESTLANT 1 (April). Numbers per $30-\mathrm{min}$ stramin-net haul; numbers per $1 \mathrm{~m}^{2}$ by Hensen-net haul; number per $\mathrm{m}^{3}$ by IHSS. (Chart 232 in the NWL publication.)
"Predation losses will depend on the abundance of predators and the length of time the fish larvae remain vulnerable members of the zooplankton ecosystem." This may, to a certain degree, be the explanation for the close relation between water temperatures at West Greenland and year-class strengths of the West Greenland cod stock.

Some observations on the abundance of predators within the area of distribution of cod eggs and larvae are mentioned in an Appendix to the zooplankton report. The following carnivores are known to be predators of fish larvae: Aglantha, ctenophores, siphonophores, chaetognaths, Tomopteris, cephalopods, Euchaeta, and hyperids. These predators were more abundant in oceanic areas than in coastal areas, and it could be shown that at West Greenland they were much more numerous in stramin-net samples (from $50-0 \mathrm{~m}$ ) taken at stations with cod larvae over deep water (more than 300 m ) than over shallow water (less than 300 m ). During NORWESTLANT 2 the average numbers per 30 min haul were 2,083 predators at deep water stations and only 101 predators at shallow water stations, and during NORWESTLANT 3 the respective numbers were 5,405 and 1,768.


Fig. 11. Distribution and abundance of cod eggs and larvae during NORWESTLANT 3 (July). Numbers per $30-\mathrm{min}$ stramin-net haul. (Chart 236 in the NWL publication.)

That means that cod larvae drifting northwards over the West Greenland fishing banks are less exposed to predation than larvae drifting westwards over deep water in Davis Strait.

## IV. Fish eggs and larvae

1. Cod eggs and larvae (Coordinator, Mr P. Hansen, Denmark)

During April cod eggs were distributed in a continuous zone from Iceland to East Greenland, around southern Greenland and along West Greenland as far north as the Arctic Circle, with some concentrations off Labrador. Figure 10 shows cod eggs taken with different nets, a gap in the grid of stramin-net stations being filled by Hensen-net stations. A special report on Labrador cod was presented by Mr A.I. Postilaky (USSR).

Figure 11 shows the distribution of cod eggs and larvae during July. Larvae were concentrated at West Greenland and at East Greenland-Iceland and

## Averoge number



Fig. 12. Average numbers of cod larvae per $30-\mathrm{min}$ stramin-net haul off West Greenland in different years from 1950 to 1963. (Fig. 35 in the NWL publication.)


Fig. 13. Year-class strength (calculated as percentage of the 1924 year-class) plotted against mean temperature, surface to 45 m over Fyllas Bank in June. (After Hermann, Hansen, and Horsted, 1965)
were absent around southern Greenland. There is some evidence that, during the critical period when eggs and larvae were drifting around southern Greenland, they were in lower temperatures and that food was scarce.

The occurrence of cod larvae in April and cod eggs in July in the East Greenland-Iceland area is remarkable and indicates a longer spawning period in the area compared with the West Greenland area, and consequently, a possibly higher production of larvae.

Figure 12 shows average numbers of cod larvae taken per $30-\mathrm{min}$ stramin-net hauls off West Greenland annually from 1950 to 1963. To judge from the quantity of larvae, as well as from the temperature conditions, the 1963 year-class was expected to be poor. Hermann, Hansen, and Horsted (1965) have shown a close relationship between variations in year-class strength of the West Greenland cod and the mean temperature in the 0-45-m layer over Fyllas Bank in June, good year-classes being associated with water warmer than $2^{\circ} \mathrm{C}$ (Fig. 13). Temperatures in 1963 indicated a relatively poor year-class. Nevertheless, some years later it became apparent that it was relatively good, as shown by Horsted in an Annex to the NORWESTLANT Report. In 1966 the 1963 year-class was numerous in the southernmost part of West Greenland inshore waters. From here it spread northwards in the following years. A suggested explanation for this is its transport by the currents from the East Greenland-Iceland larval area. In previous years, the good 1956 year-class and, partly, the 1961 yearclass occurred similarly and were also suggested to be of East Greenland-Iceland origin.


Fig. 14. Distribution and abundance of redfish larvae during NORWESTLANT 2 (May-June). Numbers per 30 -min straminnet haul. (Chart 238 in the NWL publication.)

## 2. Redfish larvae (Coordinator, Mr Jakob Magnusson, Iceland)

The widest distribution of redfish larvae was during NORWESTLANT 2 (MayJune), when they were reported from almost the entire Irminger Sea and were found in a zone along southwestern Greenland up to $65^{\circ} \mathrm{N}$ lat (Fig. 14).

Extrusion of larvae was at its peak in May in the Irminger Sea. From here they drifted to East Greenland and around Cape Farewell to off southwestern Greenland, where they mixed with those extruded locally.

Icelandic data showed that the larvae were less numerous in 1963 than in 1961 and 1962 and that in 1963 they were in deeper water than in 1961, possibly due to the stormy weather.

The West Greenland redfish stocks are recruited from the southwestern and southeastern areas. The adults migrate to these areas as has been shown by tagging fish released from pound-net catches in Godthab Fjord and recaptured off southwestern and southeastern Greenland.

Redfish larvae from Continuous Plankton Recorder samples were reported by Mr G.T.D. Henderson (Edinburgh).
3. Green1and halibut (Coordinator, Dr E. Smidt, Denmark)

Figure 15 shows larvae from stramin-net samples in July from both the NORWESTLANT survey and from all the Danish expeditions since 1908. The densest occurrence is in Davis Strait where investigations have been made for many years.


Fig. 15. Distribution and abundance of Greenland halibut larvae from all Danish expeditions since 1908 and from NORWESTLANT. Numbers per 30 -min stramin-net haul in July. (Chart 250 in NWL publication.)


Fig. 16. Average numbers of Greenland halibut larvae per $30-\mathrm{min}$ stramin-net haul off West Greenland in different years from 1950 to 1964. (Fig. 48 in the NWL publication.)

It has been known since 1908 that spawning takes place in Davis Strait in deep water south of the submarine ridge between Greenland and Canada almost along the Arctic Circle. The larvae drift north along West Greenland, but no larvae have been taken north of $68^{\circ} \mathrm{N}$ lat. in spite of very rich populations of adult fish in the northern inshore Greenland areas, where they are fished by the Greenlanders, especially in the Disko Bay and in Umanak district. Also, no larvae or eggs have been taken in the West Greenland fjords, where many stramin-net hauls have been made over several years and where dense populations of adult fish live. Figure 15 also shows that no larvae have been taken in the Polar Current zone near the coast of southeastern Greenland and the southernmost coast of West Greenland. In the East Greenland-Iceland area, the larvae are scarce.

Figure 16 shows average numbers of larvae per $30-\mathrm{min}$ stramin-net haul in different years in Davis Strait (ICNAF Div. 1B-1E) in July. Annual variations are slight, and no relation to hydrographic variations could be shown. The number of larvae was almost normal in 1963.
4. American plaice (Coordinator, Mr E. Wells, Canada)

Pelagic eggs and larvae were very numerous at West Greenland, the eggs mostly during NORWESTLANT 1-2, the larvae mostly during NORWESTLANT 3 (July). In other areas the eggs and larvae were scarce. Figure 17 shows numbers of pelagic eggs per $30-\mathrm{min}$ stramin-net haul during NORWESTLANT 1 (April).

The adult stock at West Greenland is very rich but it has not yet been fished commercially. In 1970, limited experimental fishing was carried out for industrial purposes.


Fig. 17. Distribution and abundance of American plaice eggs during NORWESTLANT 1 (April). Numbers per 30 -min stramin-net haul. (Chart 251 in the NWL publication.)

## 5. Capelin (Coordinator, Mr Jutta Magnusson, Iceland)

Figure 18 shows numbers of larvae per $m^{3}$ of water taken by IHSS during MayJune in the eastern area. Most larvae were caught in the Icelandic Shelf region. This is in agreement with earlier observations, which showed that capelin larvae are the most common of the fish larvae west of Iceland in the spring and early summer.

No larvae were observed in offshore samples from West Greenland in spite of the fact that adults are very numerous in the coastal area. However, in the fjords where spawning takes place in May-June, mainly in the tidal zone, the larvae were very numerous in the plankton samples, but all the NORWESTLANT samples were from the offshore region.


Fig. 18. Distribution and abundance of capelin larvae during NORWESTLANT 2 (May-June). Numbers per $\mathrm{m}^{3}$ by IHSS. (Chart 245 in the NWL publication.)

At West Greenland the species is very important as a food for cod, Greenland halibut, salmon, etc. For the Greenlanders, it is important for local consumption.

## 6. Other fish larvae

Larvae of halibut and wolffish species were also included in the surveys. Unfortunately, one very important fish species, the sandeel, was not included. This fish is a most important food-chain link over the offshore fishing banks at West Greenland, where it plays the same role as the capelin in the inshore area. It is an important food, especially for cod and salmon. The larva is one of the most common fish larvae over the northern West Greenland fishing banks.

Figure 19 is a reminder that most of us are still zoologists. In the middle of the figure is a cod larva, in the upper row some herbivores the early stages of which are eaten by fish larvae, in the bottom row some of the predators of fish larvae.

## V. Concluding remarks

The NORWESTLANT surveys have provided much experience and valuable information, but it has also left several open questions. In his Conclusion of the


Fig. 19. Some important North Atlantic plankton animals. 1. Calanus finmarchicus; 2. Meganyctiphanes norvegica; 3. Spiratella retroversa; 4. cod larva; 5. Euchaeta norvegica; 6. Themisto libellula; 7. Aglantha digitale; 8. Sagitta elegans. (From various authors).

Report Mr Lee says, "before any further ICNAF cooperative work is carried out in the NORWESTLANT area, this Report will be studied by its Research and Statistics Committee. The question will no doubt arise as to whether it is possible to carry out surveys which are much more sophisticated than those of the reconnoitering type which formed NORWESTLANT 1-3."

Several technical problems arose during the surveys. Considerable difficulty was met in comparing samples from the different types of plankton nets used and points to the need to use standard nets.

Indeed, I am not an expert in this matter, but I feel it would be useful to consider carefully what kind and amount of research should possibly be carried out in the future, for instance, surveys on phytoplankton production, on occurrences and quantities of fish larvae, etc.

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23. Summary of statistics on discards, $1969{ }^{1}$
by J.S. Beckett
ICNAF Secretariat
At their 1969 meeting, the Subcommittee on Statistics and the Standing Committee on Research and Statistics recommended
"that the discard portion of the summary document on discards and industrial fish continued to be pủblished annually in Redbook, Pt. III ......." (Redbook 1969, Pt. 1, p. 48)

Statistics on species and nominal catch discarded at sea for 1969 have been prepared from data submitted on ICNAF Statistics Form 4 by Canada (M), Canada (N), France (M), Germany (Fed. Rep.), Poland, Portugal, Spain and the USA.

Some discrepancies between STANA 1W and Statistics Form 4 submissions have been noted but where these appear to be obvious errors, the corrected information has been incorporated into the summaries. However, there are cases where no nominal catch can be assigned to a particular vessel class for which discard and industrial fish information is available.

Canada (M) notes that "it remains normal practice to discard 100\% of all silver hake, argentines, sculpins, lumpfish, sea robins, anglers, eelpouts and dogfish". However, these are species not sought for and, in general, are too small to be retained by a $41 / 2$ inch mesh. Skates are landed occasionally but are normally discarded. Discards of commercial species of groundfish by longliners and gill netters are negligible due to the selective properties of the gear. Discards of pollock, hake and cusk by otter trawlers are also low.

[^19]Abbreviations and Symbols Used (as in latest Statistical Bulletin)

| Species: | Had | - haddock |
| :---: | :---: | :---: |
|  | Red | - redfish |
|  | F10 | - flounders |
|  | Pol | - poilock |
|  | Her | - herring |
|  | Wol | - wolffish |
|  | Mix | - mixed |
|  | NK | - not known |
| Gear: | OT | - otter trawl |
|  | OT.Si | - otter trawl, side |
|  | OT. St | - otter trawl, stern |
|  | PT | - pair trawl |
|  | DS | - Danish seine |
| Tonnage Class: | 1 | $0-50 \mathrm{GRT}$ |
|  | 1b | 26 - 50 GRT |
|  | 2 | 51 - 150 GRT |
|  | 3 | 151-500 GRT |
|  | 4 | 501 - 900 GRT |
|  | 5 | 901-1800 GRT |
|  | 6 | over 1800 GRT |
| Country : | Can (M) | - Canada (Maritime and Quebec) |
|  | Can (N) | - Canada (Newfoundland) |
|  | Fr (M) | - France (Metropolitan) |
|  | Ger | - Germany |
|  | Pol | - Poland |
|  | Por | - Portugal |
|  | Spa | - Spain |

Source of information:

|  | Log <br> Int <br> Rep | - logbook <br> - dockside interview <br> - reports by captain |
| :--- | :--- | :--- |
| Symbols: | $\ldots$ | - not available or not reported |
|  | 0 | - magnitude known to be nil or zero |
|  | - | - magnitude known to be more than zero |
| but less than half the unit |  |  |

SUMMARY OF STATISTICS ON DISCARDS, 1969

SUMMARY OF STATISTICS ON DISCARDS， 1969 －continued

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[^20]
[^0]:    ${ }^{T}$ Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res. Doc. 71/22.

[^1]:    T Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/49, Part II.

[^2]:    T Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/82.

[^3]:    T Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res. Doc. 71/91.

[^4]:    Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/1.
    2 This applies to ICES only (C.Res. 1969/5:1,P.-v.Réun. Cons.Int. Explor.mer 1969, p. 120). The Standing Committee on Research and Statistics of ICNAF has not yet adopted the new standard polyamide, but it recomended that the Subcommittee on Assessments examine the requirements for further selection experiments in relation to adoption of the new standard (Rec.18, Redbook 1970, Part I, p. 18).

[^5]:    उThe nominal fineness is $\mathrm{R} 18,000$ tex.

[^6]:    T Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/86.

[^7]:    I Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/42.

[^8]:    T Common and scientific name as used in ICNAF Statistical Bulletin.
    2 Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/89.

[^9]:    To date, there has been no agreement among investigators about the Latin nomenclature of grenadiers. In one case, the genus is spelled Macrumus; in another, Macrourus. Sometimes Macrurus rupestris is called Coryphaenoides rupestris. We have adopted the scientific name Macrourus mpestris and the common name roundnose grenadier, as used in the ICNAF Statistical Bulletin.
    2 Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res. Doc. 71/93.

[^10]:    ${ }^{T}$ Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/65 (also ICES/ICNAF Salmon Doc. 71/9).

[^11]:    T Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/84.

[^12]:    T Submitted to 1971 Annual Meeting of ICNAF as LCNAF Res.Doc. 71/7.

[^13]:    T Norweglan ships' recoveries believed incompletely reported.

[^14]:    ${ }^{1}$ Revised version of paper submitted to 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/8.

[^15]:    I Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/87.

[^16]:    T Submitted to the 1971 Annual Meeting of ICNAF as ICNAF Res.Doc. 71/38.

[^17]:    T Revised version of paper submitted to 1971 Annual Meeting of ICNAF as ICNAF Res. Doc. 71/59.

[^18]:    T For published details, see ICNAF Special Publication, No. 7, 1968.

[^19]:    TSubmitted to the 1971 Annual Meeting of ICNAF as part of ICNAF Res.Doc. 71/27.

[^20]:    ${ }^{1}$ Motor trawler
    2 Steam trawler

