# THE NORTH-EAST ARCTIC COD STOCK'S CONSUMPTION OF VARIOUS PREY SPECIES 1984-1989 

by<br>Bjarte Bogstad and Sigbjern Mehl<br>Institute of Marine Research<br>P.O.Box 1870 Nordnes<br>N-5024 Bergen, Norway


#### Abstract

The North-East Arctic cod stock's consumption of different prey species has earlier been calculated by Mehl (1989) for the period 1984-1986. However, a new model to describe gastric evacuation in cod has now been presented by dos Santos (1990). This model is based on extensive feeding experiments with relevant prey items, done at relevant temperature conditions. This model is combined with the stomach content data to give consumption estimates for each of the years 1984-1989. The results are compared and judged against data on individual cod growth, daily rations and food conversion efficiency.


## INTRODUCTION.

One essential requirement for multispecies models is estimates of the food consumption by the main predator stocks over the area of distribution. In addition to data on the number and distribution of the different predator age groups, good estimates of the annual food requirements are needed. The latter can be estimated in two ways:

- from quantitative stomach content data and data on gastric evacuation rates (models)
-from consideration of energy requirements and data on diet composition
The first method has been chosen in several investigations, may be because it consists of more or less straight forward measurements. But these measurements which results in the consumption rates by/of the actual predator/prey stocks are important and sometimes also weak points in the models they are used in.
This is the situation for the MSVPA models both in the North Sea and in the Baltic, and it has the same importance for the multispecies modeling work in the Barents Sea. The first results of this work were presented in Bogstad and Tjelmeland (1990, 1991). In the Barents Sea stomach data were previously combined with temperature-correlated evacuation rates from the North Sea to calculate the cod's consumption rates (Mehl, 1989). Later a preliminary evacuation rate model based on feeding experiments done in Balsfjord, Northern Norway by Jorge dos Santos have been used (Mehl, 1989); (Mehl and Sunnanå, 1991). This evacuation rate model was compared to five other evacuation rate models in Bogstad and Mehl (1990), and the results were then compared with prey stock sizes and data from other investigations on daily ration and food conversion efficiency. The new evacuation rate model given by dos Santos (1990), which is based on experiments done at relevant temperature and with relevant prey, will be used for the consumption calculations made in this paper. The model takes into account both the temperature and the meal size/ body weight ratio in addition to the prey species. As we do not know the meal size from the stomach data, we will need a relationship between the meal size and the stomach content. We will suggest such a relationship, and compare the results with data from other investigations on daily ration and food conversion efficiency as was done in Bogstad and Mehl (1990).


## MATERIAL AND METHODS.

The evacuation rate model.
According to dos Santos (1990) the evacuation of a single meal for cod can be described in the following way for prey category i:

$$
S_{i t}=S_{i 0} e^{-\left(\frac{\frac{1}{2}}{H_{i} e^{-c T}\left(\frac{S_{i j}}{W}\right)^{b}}\right)^{K_{i}} \ln 2}
$$

, where $\mathrm{S}_{\mathrm{it}}$ is the stomach content at time t (hours), $\mathrm{S}_{\mathrm{i} 0}$ is the initial meal size, W the fish body weight and $H_{i}$ is the evacuation rate constant which is a theoretical value expressing for a given prey i the half-life(hours) of a meal of the same size as the fish body weight at $0^{\circ} \mathrm{C} . \mathrm{T}$ is the temperature in ${ }^{\circ} \mathrm{C}, \mathrm{b}$ and c are constants. The shape parameter $K_{i}$ determines the shape of the evacuation curve. We have here used a notation which differs slightly from the one used by dos Santos (1992).

For consumption calculation purposes, we assume a steady state, i.e that the amount of food eaten is equal to the amount digested. This means that the ration of prey item $i$ eaten per hour, $R_{i}$, becomes

$$
R_{i}=\frac{\ln 2 \cdot S_{i} \cdot K_{i}}{H_{i} \cdot e^{-c T} \cdot\left(\frac{S_{i 0}}{W}\right)^{b}} \cdot\left(\frac{t}{H_{i} \cdot e^{-c T} \cdot\left(\frac{S_{i}}{W}\right)^{b}}\right)^{K_{i}-1}
$$

In dos Santos (1990) it is shown that the shape parameter $K_{i}$ is not a main contributor to the goodness-of-fit of this single-meal model. He has estimated the parameters both when $K_{i}$ is allowed to vary and when $K_{i}$ is restricted to 1.0 for all prey. For all prey except shrimp, the $95 \%$ confidence interval for $K_{i}$ was found to contain the value 1.0 (exponential evacuation) when the model was applied in terms of wet matter of prey: Thus, we have chosen to use the parameter values obtained when $K_{i}$ is restricted to 1.0 for all prey because this is a more convenient formula to work with. Equation (2) can then be simplified to :

$$
R_{i}=\frac{\ln 2 \cdot S_{i} \cdot e^{c T}}{H_{i} \cdot\left(\frac{S_{i j}}{W}\right)^{b}}
$$

The parameter values are (wet weight): $b=0.54, c=0.11, H=205$ (krill), 533 (shrimp), 452 (herring) and 283 (capelin).
For sequential meals, the best fit was found when the term $\frac{S_{i 0}}{W}$ was changed following each new meal, assuming that the 'new' total stomach content is the stimulus driving evacuation. Thus, we will relate the initial meal size $S_{i 0}$ to the observed stomach content. We will start by assuming that the initial meal size is equal to twice the observed total average stomach content $S=\Sigma S_{i}$. The final formula for $R_{i}$ then becomes:

$$
R_{i}=\frac{\ln 2 \cdot S_{i} \cdot e^{c T}}{H_{i} \cdot\left(\frac{2 S}{W}\right)^{b}}
$$

The cod stock's consumption in tonnes of prey species $i$, prey size group $j$ in season $l\left(C_{i, j l l}\right)$ is then given by

$$
C_{i, j, l}=\sum_{k, n} R_{i, j, k, l, n} \cdot X N_{n, l} \cdot X P_{k, l, n} \cdot \text { const }
$$

where $R_{i, j, k, l, n}$ is the ration of prey species $i$, prey size group $j$ in area $k$ and season 1 for cod age group $n, \mathrm{XN}_{\mathrm{n}, 1}$ is the number of individuals of predator age group n in season 1 and $\mathrm{XP}_{\mathrm{k}, \mathrm{l}, \mathrm{n}}$ is the proportion of individuals of predator age group n in area k and season 1. Const is a scaling factor in order to get the consumption in tonnes per season.

It was decided to apply the evacuation rate for capelin also for amphipods. For cod, haddock and redfish the evacuation rate for herring was applied. For "other" prey species, the evacuation rate for krill was applied for cod age group 1 and 2, because for these age groups "other" consists mainly of smaller crustaceans and other easily digestable food. For the older cod age groups, the evacuation rate for herring was applied, because "other" for these age groups mainly consists of fish prey.

## Data used.

The temperatures used are calculated by a temperature model which is also used by the multispecies model for the Barents Sea (MULTSPEC) (Bogstad and Tjelmeland, 1990, 1991). Data from standard hydrographical sections (Fugløya-Bjørnøya, Vardø N, Kola section) are used. At different depths and over different parts of the sections Fourier analysis is used on the temperature data. The temperature is then integrated in time for a representative location in each area to give one temperature for each year, area and month. The procedure is documented in Alvarez and Tjelmeland (1989). A depth of 100 m has been used for these calculations. Due to problems with the temperature data base, the temperatures in 1988 have been set equal to the temperatures at the same position/time as in $1987+0.3^{\circ} \mathrm{C}$, and the temperatures in 1989 have been set equal to the temperatures at the same position/time as in $1987+0.9^{\circ} \mathrm{C}$. These differences of $0.3^{\circ} \mathrm{C}$ and $0.9^{\circ} \mathrm{C}$ are equal to the differences in the yearly mean at the Kola section between these years (PINRO, Murmansk, pers.comm.).
Stomach data for the years 1984-1989, collected by IMR and PINRO have been used in the calculations. Details about stomach sampling, analysis and aggregation of data are given in Mehl $(1986,1989)$ and Mehl and Yaragina (1991). The area distribution used in the multispecies model for the Barents Sea until 1990, which also is the basis for the division of the Barents Sea in areas for consumption calculation purposes, is shown on the map below.

Figure 1 Multispecies area division of the Barents Sea.


The consumption is calculated by 3 areas (II $+\mathrm{IV}, \mathrm{III}+\mathrm{V}$ and $\mathrm{VI}+\mathrm{VII}$ ) for each halfyear and predator age group. Because area VI+VII is sampled only once a year, the data
collected during the second half of the year are also used for the first half of the year. The data collected in 1984 were also used for the first part of 1984. During the second half of 1985 the sampling in the southern part of the Barents Sea (areas II+IV and III+V) was rather poor. Because of the large seasonal changes in the diet in these areas (Mehl, 1986), the data collected for the second part of 1984 in the same areas were used. But the content of shrimp was reduced to the same level (by percentage) as in area VI + VII during the second part of 1985, because of the all-over reduction of the shrimp stock from 1984 to 1985 (Tveranger and Øynes, 1985). In the second half of 1986, data for areas II +IV and III+V combined were used to give the diet for area II + IV due to poor sampling in area $\Pi+I V$.
Table 5 in the appendix presents the mean weight of cod in 1984-1989 in the first and second half of the year. The weights are calculated using the weight in stock used by the ICES Arctic Fisheries Working group (Anon., 1991a) for ages 3 and older and the average of Norwegian and Russian survey weights for ages 1 and 2 (all at January 1), and assuming that the growth in weight is the same in every quarter during the year. (For the 7-9 group, the weight is calculated as a weighted average of the weights of age groups 7-9). For some of these years, there is a large discrepancy between the Norwegian and Russian survey weights, and this should be kept in mind when evaluating the calculations of daily ration in $\%$ of body weight and food conversion efficiency. This discrepancy may to a large extent be due to differences in age reading, and this matter is now under investigation (Anon., 1993). The Working Group makes an arithmetic average of the weight at age from these two surveys.
VPA-data from the Arctic Fisheries Working Group (Anon., 1991a) are used when the total cod stock's consumption is estimated. The number in each age group in the stock in each season is set to be the number in the middle of the season. This is found by reducing the number at the beginning of the year, using one fourth of the yearly natural and fishing mortality for each quarter. For age groups 1 and 2, the number at the beginning of each season is found by back-calculating the number at age 3 (Anon., 1991a) using a yearly natural mortality (M) of 0.2. The numbers are presented in table 6 in the appendix.
The geographical and seasonal distribution of the different cod agegroups are based on survey data, both acoustic and bottom trawl data (Dalen et.al. (1984), Hylen et. al. (1985), Hylen et. al. (1986), Godø et. al. (1987), Hylen et. al. (1988a), Hylen et. al. (1988b), Hylen et. al. (1989), Jakobsen et. al. (1989), Godø et. al. (1989) and unpublished survey data). They are presented in table 7 in the appendix.

## DISCUSSION.

The calculated consumption by cod in the period 1984-1989 of the different prey species is shown in table 1.

Table 1 The Northeast Arctic cod stock's consumption in 1000 tonnes of the main prey species in 1984-1989 (\% of total consumption in parentheses).

| Year <br> Prey <br> species | 1984 | 1985 | 1986 | 1987 | 1988 | 1989 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| Amphi- <br> pods | $16(1)$ | $113(3)$ | $721(25)$ | $742(35)$ | $1029(44)$ | $646(27)$ |
| Shrimp | $417(19)$ | $156(5)$ | $125(4)$ | $177(8)$ | $128(5)$ | $137(5)$ |
| Capelin | $867(40)$ | $1938(56)$ | $956(33)$ | $226(11)$ | $511(22)$ | $783(31)$ |
| Herring | $66(3)$ | $161(5)$ | $131(4)$ | $30(1)$ | $1(0)$ | $4(0)$ |
| Cod | $33(2)$ | $47(1)$ | $103(4)$ | $32(2)$ | $10(1)$ | $8(0)$ |
| Haddock | $45(2)$ | $40(1)$ | $79(3)$ | $3(0)$ | $5(0)$ | $36(2)$ |
| Redfish | $331(15)$ | $205(6)$ | $251(8)$ | $299(14)$ | $190(8)$ | $213(9)$ |
| Others | $371(17)$ | $813(23)$ | $565(19)$ | $631(29)$ | $482(20)$ | $670(26)$ |
| Total | $2146(99)$ | $3475(100)$ | $2932(100)$ | $2141(100)$ | $2355(100)$ | $2497(100)$ |

From the table above, we see that capelin and amphipods were the most important prey species during the period, followed by redfish and shrimp. Young cod (cannibalism), haddock and herring contributed $5 \%$ or less each to the total consumption in all years. We see that amphipods, which were almost absent from the diet in 1984 and 1985, were the most important prey in 1987 and 1988. The proportion of capelin in the diet decreased from 1985 to 1987, in accordance with the collapse of the capelin stock, but after that the proportion rose again. The proportion of shrimp in the diet was relatively high in 1984, but stayed at a low level for the rest of the period. Compared to the other prey species, the percentage of redfish and "other" prey in the diet was relatively stable in the period. The cod stock's consumption was highest in 1985, both in tonnes and relative to the biomass of the cod stock at the beginning of the year.
The consumption of redfish, which is relatively stable, is high compared to the biomass of an age group at age 6 (10000-55000 tonnes for Sebastes mentella in the period in question, and less for Sebastes marinus (Anon., 1991a)). The main part of the redfish consumed is younger than 6 years, and the biomass of an age group of redfish younger than 6 years is less than that at age 6 . Thus, the predation by cod on redfish is quite significant.
The decrease in the consumption of shrimp from 1984 to 1985 seem to be consistent with the decrease in the biomass estimate (Tveranger and Øynes, 1985), and the relatively low consumption of shrimp in the following years is also consistent with the biomass estimate time series. The calculated consumption is of the same order as the biomass estimate in the period (Anon., 1991b), which is, however, thought to be an
underestimate. The catch in the period varies between 45 and 120 thousand tonnes, and the predation from cod is thus more important than the catch.
The consumption of mature capelin by cod has been studied by Bogstad and Tjelmeland (1990, 1991). It should be mentioned that for the first half of the year, most of the samples are taken in the first quarter, when there may be intensive predation on mature capelin, and this may lead to an overestimate of the total consumption of capelin.
The high consumption of cod (cannibalism) in the years 1984-1987 may be part of the explaination why the 1984-1986 year classes of cod, which seemed to be very strong at the 0-group stage, turned out to be rather poor. See also Skagen et.al. (1990). To a somewhat lesser extent, this also happened for haddock. The high consumption of haddock in 1989 compared to 1987 and 1988 may be due to an error in the data.
The consumption of herring is highest in the years 1984-1986, coinciding well with the time period when the strong 1983 year class of herring was present in the Barents Sea. For the age groups 1 and 2, krill and small crustaceans dominate the "other" group. For the older age groups, "other" consists of various crustaceans and small fish. The proportion of fish in "other" increases with age.
Table 2 summarizes the annual consumption per cod for the different age groups:
Table 2 Annual consumption per cod (grams) by age group in the Northeast Arctic cod stock in 1984-1989

| Year <br> Age group | 1984 | 1985 | 1986 | 1987 | 1988 | 1989 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 1 | 116 | 185 | 72 | 138 | 134 | 143 |
| 2 | 569 | 681 | 387 | 384 | 455 | 619 |
| 3 | 1501 | 1842 | 1094 | 879 | 864 | 1361 |
| 4 | 2716 | 3509 | 2532 | 1574 | 1855 | 2498 |
| 5 | 4823 | 5660 | 4785 | 3120 | 3452 | 4105 |
| 6 | 6829 | 9451 | 7232 | 5959 | 6228 | 6490 |
| $7-9$ | 11307 | 15996 | 13583 | 8811 | 13430 | 10641 |

Table 3 gives the daily consumption per cod in percent of the bodyweight in the middle of the year (daily coefficient).

Table 3 Daily coefficient by agegroup for Northeast Arctic cod in 1984-1989.

| Year <br> Age group | 1984 | 1985 | 1986 | 1987 | 1988 | 1989 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 1 | 0.54 | 0.95 | 0.42 | 0.84 | 0.55 | 0.48 |
| 2 | 0.56 | 0.91 | 0.73 | 0.82 | 0.68 | 0.66 |
| 3 | 0.57 | 0.78 | 0.65 | 0.84 | 0.68 | 0.76 |
| 4 | 0.51 | 0.77 | 0.63 | 0.71 | 0.83 | 0.82 |
| 5 | 0.54 | 0.73 | 0.71 | 0.60 | 0.87 | 0.87 |
| 6 | 0.52 | 0.76 | 0.66 | 0.67 | 0.81 | 0.86 |
| $7-9$ | 0.59 | 0.85 | 0.75 | 0.53 | 0.82 | 0.74 |

The new evacuation rate model gives a lower consumption estimate than found previously. The introduction of meal size dependence will mostly affect the consumption of capelin, because stomachs with large content usually contains capelin.
We see that the daily coefficient is in the interval $0.4-1.0$, which is lower than found in other investigations, see Bogstad and Mehl (1990). Daan (1973) found a drop in the daily coefficient for increasing age for cod in the North Sea. Our calculations do not show such a trend, which leads us to believe that the consumption by the youngest age groups is underestimated. We have relatively few samples of the youngest age groups, and the proportion of empty stomachs is high. Also, in the experiments done by dos Santos cod of sizes corresponding to age 3 and 4 for Barents Sea cod dominated. According to the latest experiments done by dos Santos (pers. comm.) it also seems that the consumption by the youngest fish is somewhat underestimated.
Table 4 presents the annual weight increment in percent of the annual consumption (food conversion efficiency - FCE).

Table 4 Food conversion efficiency (\%) by agegroup for Northeast Arctic cod in 1984-1989.

| Year <br> Age group | 1984 | 1985 | 1986 | 1987 | 1988 | 1989 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 1 | 55 | 29 | 53 | 32 | 70 | 98 |
| 2 | 65 | 34 | 34 | 33 | 51 | 46 |
| 3 | 25 | 26 | 16 | 17 | 41 | 28 |
| 4 | 19 | 19 | 13 | 13 | 27 | 26 |
| 5 | 22 | 14 | 11 | 11 | 22 | 21 |
| 6 | 14 | 9 | 13 | 10 | 17 | 18 |

The FCE for the 7-9 age group has not been calculated, because different age groups dominates this group from year to year. Also, the calculations for this group are based on relatively few samples, which leads to a large variation in calculated consumption from year to year.
For the youngest age groups, the FCE is obviously too high. The reason for this is the previously mentioned underestimation of the consumption for these age groups. The consumption estimates seems reasonable compared to the stock estimates of the prey species, while Mehl (1989) found that some prey species were "overconsumed". Still, the predation pressure on some prey year classes is considerable, see e.g. Skagen et. al. (1990).
For cod of medium size, the FCE in 1986/1987 is significantly lower than in the other years. This indicates that the low growth of cod in this period was mostly due to lower energy content in the food (less capelin!), and not so much lower total food consumption. There was, however, low abundance of capelin also in 1988, but a high FCE.
The FCE values above should be compared to values obtained from feeding experiments. Values of FCE obtained for fish in capture can be regarded as an upper limit. It would of course be very interesting to redo these calculations using energy content and not wet weight as done here.

Future improvements: The evacuation rate model could be improved by doing experiments with small and large cod and other prey species (redfish, amphipods, gadoids). Also, more experiments with multiple feeding should be performed. In order to make the best possible use of the new evacuation rate model, new calculations should be done based on individual stomach content and predator size data. It should then also be possible to use the actual temperature at the sampling site. The stomach sampling should be improved in order to get a better coverage in area and time. The differences in predator distribution, stomach content and temperature between bottom and pelagic should also be taken into account by trying to sample more representatively, The stomach content data have so far not been weighted by catch rate, due to data base problems. For the multispecies model in the Barents Sea, it is at the moment most important to get a precise estimate of the consumption of capelin by cod. Given a stock estimate of cod and of capelin, the uncertainty in the consumption estimate should not exceed the uncertainty in the stock size estimates.

## REFERENCES.

Anon. 1991a. Report of the Arctic Fisheries Working Group. ICES C.M. 1991/ Assess: 3.
Anon., 1991b. Ressursoversikt 1991. Fisken og havet, 1991. Særnummer 1. Institute of Marine Research, Bergen. (In Norwegian).
Anon. 1993. Report of the Arctic Fisheries Working Group. ICES C.M. 1993/ Assess: ?.
Alvarez, J. and Tjelmeland, S. 1989. MULTSPEC og oseanografi. IMR internal documentation. (In Norwegian).
Bogstad, B. and Mehl, S. 1990. The consumption rate of Northeast Arctic cod - a comparison of gastric evacuation models. ICES C.M. 1990/G:22. 12 pp .
Bogstad, B. and Tjelmeland, S. 1990. Estimation of predation mortalities on capelin using a cod-capelin model for the Barents Sea. ICES C. M. 1990/H:16. 33 pp.
Bogstad, B. and Tjelmeland, S. 1992. A method for estimation of predation mortalities on capelin using a cod-capelin model for the Barents Sea. In Bogstad, B. and Tjelmeland, S. (eds. ): Interrelations between fish populations in the Barents Sea. Proceedings of the fifth PINRO-IMR symposium, Murmansk 12-16 August 1991.
Daan, N. 1973. A quantitative analysis of the food intake of North Sea cod, Gadus morhua. Neth. J. Sea. Res., 6(4):479-517.
Dalen, J., Hylen, A., Jakobsen, T., Nakken, O. and Randa, K. 1984. Preliminary report of the Norwegian investigations on young cod and haddock in the Barents Sea during the winter 1984. ICES C.M. 1984/G:44.
dos Santos, J. 1990. Aspects of the eco-physiology of predation in Atlantic cod, Gadus morhua L., Dr. scient. thesis, University of Tromsø. 116 pp .
Godø, O.R., Hylen, A., Jacobsen, J.A., Jakobsen, T., Mehl, S., Nedreaas,K. and Sunnanå, K. 1987. Estimates of stock size of Northeast Arctic cod and haddock from survey data 1986/1987. ICES C.M. 1987/G:37.
Godø, O.R., Mehl, S. and Nedreaas, K. 1989. Investigations on cod, haddock, redfish and Greenland halibut in the Barents Sea and the Svalbard area autumn 1989. IMR internal survey-report.
Hylen, A., Jacobsen, J.A., Jakobsen, T., Mehl, S. Nedreaas, K. and Sunnanå, K. 1988a. Estimates of stock size of Northeast Arctic cod and haddock, Sebastes mentella and Sebastes marinus from survey data, winter 1988. ICES C.M. 1988/G:43.
Hylen, A., Jacobsen, J.A., Mehl, S. and Nedreaas, K. 1988b. Estimates of stock size of cod, haddock, redfish and Greenland halibut in the Barents Sea and Svalbard area autumn 1987. ICES C.M. 1988/ G:44.
Hylen, A., Jakobsen, T., Nakken, O. and Sunnanå, K. 1985. Preliminary report of the Norwegian investigations on young cod and haddock in the Barents Sea during the winter 1985. ICES C.M. 1985/G:68.
Hylen, A., Jakobsen, T., Nakken, O., Nedreaas, K. and Sunnanå, K. 1986. Preliminary report of the Norwegian investigations on young cod and haddock in the Barents Sea during the winter 1986. ICES C.M. 1986/G:76.
Hylen, A., Loeng, H., Mehl, S. and Nedreaas, K. 1989. Estimates of stock size of cod, haddock, redfish and Greenland halibut in the Barents Sea and Svalbard area autumn 1988. ICES C.M. 1989/ G:39.

Jakobsen, T., Mehl, S., Nakken, O., Nedreaas, K. and Sunnanå, K. 1989. Estimates of stock size of Northeast Arctic cod and haddock, Sebastes mentella and Sebastes marinus from survey data, winter 1989. ICES C.M. 1989/G:42.
Mehl, S. 1986. Stomach contents of North-East Arctic cod and possible changes in the diet the last years. ICES C.M. 1986/G:29. 29 pp.
Mehl, S. 1989. The North-East Arctic cod stock's consumption of commercially exploited prey species in 1984-1988. Rapp. P-v. Reun. Cons. int. Explor. Mer, 188: 185-205.
Mehl, S. and Sunnanå, K. 1991. Changes in growth of Northeast Arctic cod related to food consumption in 1984-1988. ICES mar. Sci. Symp., 193: 109-112.
Mehl, S. and Yaragina, N.A. 1992. Methods and results in the joint PINRO-IMR stomach sampling program. In Bogstad, B. and Tjelmeland, S. (eds. ): Interrelations between fish populations in the Barents Sea. Proceedings of the fifth PINRO-IMR symposium, Murmansk 12-16 August 1991.
Skagen, D.W., Bogstad, B. and Mehl, S. 1990. CEC- A model for cod-cód cannuibalism in the Barents Sea. Working paper no. 14, ICES Multispecies Assessment Working Group 1990. 9 pp. (unpubl.)
Tveranger, B. and Øynes, P. 1985. Results of stratified trawl surveys for shrimps ((Pandalus borealis) in the Barents Sea in May and in the Svalbard region in July-August 1985. ICES C.M. 1985/K:50.

## Appendix A Cod stock data used.

Table 5 Cod weight-at-age (kg)

| Cod age <br> Year/ <br> season | 1 | 2 | 3 | 4 | 5 | 6 | $7-9$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $1984-\mathrm{I}$ | 0.042 | 0.182 | 0.625 | 1.328 | 2.160 | 3.225 | 4.883 |
| $1984-\mathrm{II}$ | 0.074 | 0.367 | 0.815 | 1.583 | 2.680 | 3.855 | 5.468 |
| $1985-\mathrm{I}$ | 0.040 | 0.148 | 0.578 | 1.075 | 1.913 | 3.163 | 4.825 |
| $1985-\mathrm{II}$ | 0.067 | 0.262 | 0.813 | 1.405 | 2.318 | 3.608 | 5.415 |
| $1986-\mathrm{I}$ | 0.036 | 0.113 | 0.365 | 1.010 | 1.708 | 2.755 | 4.613 |
| $1986-\mathrm{II}$ | 0.055 | 0.178 | 0.455 | 1.170 | 1.983 | 3.225 | 5.238 |
| $1987-\mathrm{I}$ | 0.034 | 0.096 | 0.248 | 0.550 | 1.330 | 2.265 | 4.273 |
| $1987-\mathrm{II}$ | 0.056 | 0.159 | 0.323 | 0.650 | 1.498 | 2.555 | 4.718 |
| $1988-\mathrm{I}$ | 0.043 | 0.125 | 0.270 | 0.485 | 0.893 | 1.840 | 3.575 |
| $1988-\mathrm{II}$ | 0.090 | 0.242 | 0.430 | 0.735 | 1.278 | 2.360 | 4.565 |
| $1989-\mathrm{I}$ | 0.047 | 0.186 | 0.395 | 0.673 | 1.075 | 1.768 | 3.490 |
| $1989-\mathrm{II}$ | 0.117 | 0.328 | 0.585 | 0.998 | 1.505 | 2.363 | 4.370 |

Table 6 Number of individuals (million) of each age group by season and year.

| Cod age <br> Year/ <br> season | 1 | 2 | 3 | 4 | 5 | 6 | $7-9$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $1984-\mathrm{I}$ | 1121 | 572 | 363 | 125 | 73 | 42 | 34 |
| $1984-\mathrm{II}$ | 1014 | 517 | 326 | 107 | 57 | 29 | 17 |
| $1985-\mathrm{I}$ | 334 | 917 | 463 | 282 | 85 | 41 | 23 |
| $1985-\mathrm{II}$ | 302 | 830 | 410 | 237 | 64 | 28 | 13 |
| $1986-\mathrm{I}$ | 250 | 273 | 747 | 352 | 184 | 44 | 24 |
| $1986-\mathrm{II}$ | 226 | 247 | 668 | 292 | 131 | 27 | 13 |
| $1987-\mathrm{I}$ | 256 | 205 | 222 | 573 | 222 | 83 | 22 |
| $1987-\mathrm{II}$ | 231 | 185 | 198 | 471 | 155 | 47 | 12 |
| $1988-\mathrm{I}$ | 209 | 209 | 166 | 173 | 362 | 104 | 32 |
| $1988-\mathrm{II}$ | 190 | 190 | 149 | 146 | 260 | 67 | 17 |
| $1989-\mathrm{I}$ | 261 | 171 | 170 | 129 | 120 | 184 | 52 |
| $1989-\mathrm{II}$ | 236 | 155 | 152 | 107 | 96 | 128 | 33 |

Table 7 Proportion of individuals by age group, area, season and year for the Northeast Arctic cod stock in 1984-1989

| Area | Year- <br> Season | Age 1 | Age 2 | Age 3 | Age 4 | Age 5 | Age 6 | Age 7-9 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| II+IV | $1984-1$ | 0.002 | 0.109 | 0.322 | 0.417 | 0.573 | 0.576 | 0.782 |
| III+V | $1984-1$ | 0.980 | 0.827 | 0.653 | 0.544 | 0.398 | 0.409 | 0.208 |
| VI+VII | $1984-1$ | 0.018 | 0.064 | 0.025 | 0.039 | 0.029 | 0.015 | 0.010 |
| II+IV | $1984-2$ | 0.002 | 0.109 | 0.322 | 0.417 | 0.490 | 0.393 | 0.540 |
| III+V | $1984-2$ | 0.980 | 0.827 | 0.653 | 0.544 | 0.453 | 0.577 | 0.439 |
| VI+VII | $1984-2$ | 0.018 | 0.064 | 0.025 | 0.039 | 0.057 | 0.030 | 0.021 |
| II+IV | $1985-1$ | 0.000 | 0.000 | 0.057 | 0.288 | 0.664 | 0.840 | 0.923 |
| III+V | $1985-1$ | 0.654 | 0.917 | 0.854 | 0.479 | 0.302 | 0.058 | 0.000 |
| VI+VII | $1985-1$ | 0.346 | 0.083 | 0.087 | 0.233 | 0.033 | 0.102 | 0.077 |
| II+IV | $1985-2$ | 0.000 | 0.000 | 0.057 | 0.288 | 0.631 | 0.737 | 0.847 |
| III+V | $1985-2$ | 0.654 | 0.917 | 0.854 | 0.479 | 0.302 | 0.058 | 0.000 |
| VI+VII | $1985-2$ | 0.346 | 0.083 | 0.087 | 0.233 | 0.066 | 0.204 | 0.153 |
| II+IV | $1986-1$ | 0.000 | 0.000 | 0.057 | 0.332 | 0.660 | 0.883 | 0.848 |
| III+V | $1986-1$ | 0.991 | 0.868 | 0.846 | 0.550 | 0.295 | 0.066 | 0.000 |
| VI+VII | $1986-1$ | 0.009 | 0.132 | 0.096 | 0.119 | 0.045 | 0.050 | 0.152 |
| II+IV | $1986-2$ | 0.000 | 0.000 | 0.057 | 0.332 | 0.615 | 0.833 | 0.600 |
| III+V | $1986-2$ | 0.991 | 0.868 | 0.846 | 0.550 | 0.295 | 0.066 | 0.000 |

Table 7 (Continued) Proportion of individuals by age group, area, season and year for the Northeast Arctic cod stock in 1984-1989

| VI+VII | $1986-2$ | 0.009 | 0.132 | 0.096 | 0.119 | 0.090 | 0.100 | 0.400 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| II+IV | $1987-1$ | 0.000 | 0.110 | 0.220 | 0.592 | 0.830 | 0.712 | 0.800 |
| III+V | $1987-1$ | 0.333 | 0.292 | 0.206 | 0.298 | 0.080 | 0.086 | 0.000 |
| VI+VII | $1987-1$ | 0.666 | 0.599 | 0.574 | 0.110 | 0.090 | 0.203 | 0.200 |
| II+IV | $1987-2$ | 0.000 | 0.090 | 0.060 | 0.282 | 0.464 | 0.394 | 0.600 |
| III+V | $1987-2$ | 0.133 | 0.242 | 0.054 | 0.140 | 0.044 | 0.048 | 0.000 |
| VI+VII | $1987-2$ | 0.867 | 0.667 | 0.887 | 0.578 | 0.492 | 0.559 | 0.400 |
| II+IV | $1988-1$ | 0.000 | 0.000 | 0.307 | 0.480 | 0.514 | 0.477 | 0.500 |
| III+V | $1988-1$ | 1.000 | 0.590 | 0.475 | 0.275 | 0.268 | 0.114 | 0.000 |
| VI+VII | $1988-1$ | 0.000 | 0.410 | 0.218 | 0.245 | 0.218 | 0.409 | 0.500 |
| II+IV | $1988-2$ | 0.000 | 0.000 | 0.201 | 0.234 | 0.338 | 0.367 | 0.600 |
| III+V | $1988-2$ | 1.000 | 0.200 | 0.311 | 0.134 | 0.124 | 0.088 | 0.000 |
| VI+VII | $1988-2$ | 0.000 | 0.800 | 0.488 | 0.632 | 0.538 | 0.545 | 0.400 |
| II+IV | $1989-1$ | 0.222 | 0.286 | 0.164 | 0.286 | 0.407 | 0.601 | 0.639 |
| III+V | $1989-1$ | 0.667 | 0.643 | 0.639 | 0.586 | 0.483 | 0.324 | 0.250 |
| VI+VII | $1989-1$ | 0.111 | 0.071 | 0.197 | 0.128 | 0.110 | 0.075 | 0.111 |
| II+IV | $1989-2$ | 0.222 | 0.310 | 0.134 | 0.046 | 0.083 | 0.125 | 0.115 |
| III+V | $1989-2$ | 0.667 | 0.619 | 0.669 | 0.826 | 0.708 | 0.725 | 0.691 |
| VI+VII | $1989-2$ | 0.111 | 0.071 | 0.197 | 0.128 | 0.209 | 0.150 | 0.194 |

