# ANCHOVY EGG DISPERSAL AND MORTALITY AS INFERRED FROM CLOSE-INTERVAL OBSERVATIONS 

PAUL E. SMITH AND ROGER P. HEWITT<br>National Marine Fisheries Service<br>Southwest Fisheries Center<br>P.O. Box 271<br>La Jolla, California 92038


#### Abstract

Kill. Millions of tons of water come down that river. How do you know the day you made your tests there wasn't something unusual about the water?


Stockman. No, I took too many samples.
Kiil. How do you know? Why couldn't those little animals have clotted up only in the patch of water you souped out of the river? How do you know the rest of it wasn't pure?

Stockman. It's not probable . . . .
An Enemy of the People ${ }^{1}$

## ABSTRACT

The oceanic boundary currents and continental borderlands that anchovy inhabit comprise diverse spatial and temporal scales of motion. Repetitive sampling at close intervals has been conducted to explore the sources and intensity of variance of observations and maximize the effectiveness of a managementoriented sampling program. The critical scale of repetitive sampling occurs when adjacent observations are so interdependent that they do not improve the precision of the abundance estimates. Sampling studies indicate that the scale of anchovy egg interdependence is on the order of several hundred meters for some days and somewhat less than this in the first few hours after spawning. Spatial correlation appears to diminish rapidly at scales larger than $2,000 \mathrm{~m}$. The minimum distance between stations in the California egg production cruises is $7,500 \mathrm{~m}$; in the Peru egg production survey it was $5,500 \mathrm{~m}$.

In general, large structures in the ocean persist longer than small ones. One simple parameterization of this phenomenon is that a $600-\mathrm{m}$ structure persists for 6 hours in the ocean and that structures of 2,800 , $27,000,147,000 \mathrm{~m}$ persist for a day, a week, and a month, respectively. From simple considerations, if the average value of food abundance is too low for larval fish to survive, the useful scale of oceanic feature that contains sufficient food is about 27 km if the food is required for a week, or 147 km if the food is required for a month. The critical scale of these features becomes smaller if production within the

[^0]structure exceeds consumption and washout. The influence of persistence on survival may diminish as juveniles gain the ability to graze in schools and swim through larger intervening distances between favorable patches. We believe that experimental repetitive sampling supports the general scale for the passive dispersion of eggs: no obvious interrelations between interannual changes in these features and egg and larval survival were noted for the first 20 days of life. The consequences of a laterally incoherent and dissipated habitat may be delayed to the late larval and juvenile stages.

Will we be able to observe the later life stages with the same intensity and accuracy that we have observed the embryonic stages?

## RESUMEN

Las corrientes oceánicas de margen y los aledaños continentales que habita la anchoveta sufren movimientos de diferentes escalas temporales y espaciales. Se condujeron muestreos replicados a intervalos reducidos con el fin de explorar las fuentes e intensidad de la varianza de las observaciones y maximizar la eficacia de un programa de muestreo orientado hacia el manejo del recurso. La escala crítica del muestreo repetitivo ocurre cuando las observaciones adyacentes son tan interdependientes que no aumentan la precisión de las estimaciones de abundancia. Muestreos experimentales indican que la escala de interdependencia para los huevos de la anchoveta es del orden de varios centenares de metros durante algunos días, y algo menor en las primeras horas después del desove. La correlación espacial parece disminuir rápidamente a escalas mayores de 2000 m . La distancia mínima entre estaciones en las campañas californianas de producción de huevos es de 7500 m ; en los estudios peruanos de producción de huevos era de 5500 m . En general, las estructuras oceánicas mayores son más persistentes que las pequeñas. Una expresión cuantitativa sencilla de este fenómeno consiste en que estructuras de 600 m persisten en el oceano durante 6 horas, mientras que aquéllas de $2,800,27,000$ y $147,000 \mathrm{~m}$ persisten durante un día, una semana, y un mes, respectivamente. Partiendo de consideraciones simples se deduce que si la abundancia promedio de alimento es demasiado baja para la supervivencia de las larvas
de peces, la escala útil de la estructura oceánica que contiene alimento suficiente es de, aproximadamente, 27 km , si el alimento es requerido para una semana, o 147 km si lo es para un mes. La escala crítica de estas características disminuye si la producción dentro de la estructura excede al consumo y la exportación. La influencia de esta persistencia en la supervivencia puede disminuir a medida que los juveniles van adquiriendo la capacidad de alimentarse en cardúmenes y de nadar distancias mayores entre lugares favorables. Creemos que el muestreo repetitivo experimental concuerda con la escala general de dispersión pasiva de los huevos: no se observaron interrelaciones obvias entre cambios interanuales en estas características y la supervivencia de huevos y larvas durante los primeros 20 días de vida. Las consecuencias de un hábitat lateralmente incoherente y disipado pueden ser pospuestas hasta los estados larvales avanzados y los juveniles.
¿Podremos observar los estados de desarrollo tardíos con al misma intensidad y precisión con que observamos los estados embrionarios?

## INTRODUCTION

Demands on oceanic sampling have rapidly increased from the simple detection and description of resources (Ahlstrom 1968; Hempel 1973; Smith and Richardson 1977) to the testing of hypotheses on the causes of recruitment failure in dynamic coastal areas like the California Current (Lasker 1975; Vlymen 1977; Parrish and MacCall 1978; Lasker and Zweifel 1978; Parrish et al. 1981; Smith 1981; Hewitt 1981; Bakun and Parrish 1982). Smith (1981) described sampling strategies for testing several hypothesized sources of year-class failure; the sources include larval transport, critical period, predation including cannibalism, starvation, unfavorable distribution pattern, and parental deficiencies at the time of spawning. Success in the studies underway have encouraged international organizations (Bakun et al. 1982) to compare dynamic areas like the eastern boundaries of the oceans off South America and Africa with respect to the causes of massive fluctuations in fish reproductive success. These studies require additional work on efficient delineation of distribution and biomass estimation. Because of the nature of the survival mechanisms postulated, directed work on spatial pattern, turbulent diffusion, transport, and survival is also needed.

The spatial pattern of plankton may be considered from the aspects of interpreting existing samples and the strategy of future sampling (Silliman 1946; Sette and Ahlstrom 1948; Taft 1960; Zweifel and Smith 1981). Another aspect of spatial pattern of plankton is the interaction between predator and prey, in particu-
lar where food aggregations are necessary for sufficient feeding rates (Lasker 1975; Vlymen 1977; Lasker and Zweifel 1978; Hewitt 1981). Lastly, schooling coastal pelagic fishes proceed through a planktonic phase of weeks to months, and the eventual retention of viable concentrations of juveniles near the coast may be controlled by rates of turbulent diffusion and cross-shelf transport during the spawning season (Smith 1973; Smith and Lasker 1978; Parrish and MacCall 1978; Hewitt 1982; Bakun and Parrish 1982; Smith 1985).

The scale of spatial pattern contains additional information about the recent history and near future of the pelagic aggregation. For example, if individual ( 0.3 m ) spawning pelagic fish like sardines are found in populations ( $1,000 \mathrm{~km}$ ), school groups ( 10 km ), schools ( 100 m ), and spawning "cliques" ( 30 m ), one can infer from the rate of dispersal by turbulent diffusion that the major sources of variance in samples are from individuals, cliques, and schools rather than from school groups and populations based on the length-scale assumptions (Smith 1973). Since this pattern of eggs imposed by the spawning and fertilization behavior of the adults persists several days into the larval stage, one may postulate that similar features like diatom patches or grazed gaps in diatom layers would persist equally long in the pelagic environment (Okubo 1971).

It is the purpose of this symposium to explore similarities among the eastern boundary areas of the world oceans, because these contain the potential of tens of millions of tons annual catch of schooling coastal pelagic engraulids, clupeids, scombrids, and carangids. These fisheries are generally not managed and are subject to large natural fluctuations, which seem to be augmented by present fishing practices (Murphy 1977; Smith 1981; Bakun and Parrish 1982; Lasker 1985; Smith 1985). Our objective in this paper is to present a small-scale empirical study of anchovy egg dispersal in a coastal site. The study may then be considered in the context of large-scale environmental features and biological surveys. We believe that existing techniques can be modified to measure offshore drift and dispersal within the coastal habitat and thus describe the habitat of schooling coastal pelagic fish in eastern boundary habitats.

## METHODS

The methods used for close-interval sampling of the anchovy population off California are described in detail in Lasker (1984), Smith and Hewitt (1984), Smith, Flerx, and Hewitt (1984) and Moser and Ahlstrom (1984). We will only briefly describe the planning and conduct of the cruise, station activity, the


Figure 1. The survey area off southern California. Ordinate values are degrees north latitude, and the abscissa values are degrees west longifude. The lengths of the bars are proportional to the log of the number of anchovy eggs per 0.05 square meters as an average of eight replicates at each station. (See Table 5 for data.)
net, the procedures for sorting, staging and aging the eggs, and the data analysis.

## Cruise

The cruise was planned to encompass the spawning habitat of the central population of the northern anchovy (Engraulis mordax) with groups of eight samples at stations 20 miles apart in the cross-shelf direction and 40 miles apart parallel to the general run of the coast. Unfortunately, the $25-\mathrm{m}$ motor vessel (Scorpius) chartered for this cruise was not competent to occupy the preplanned stations in the weather of January 1979; therefore, stations in the general area of the spawning were occupied at haphazard positions that could be approached from safe harbors among the islands of the Southern California Bight. The stations that had at least one egg in eight tows are plotted in Figure 1 as "sticks" with height proportional to the log of the mean number of total anchovy eggs per observation.

On station we recorded position, cast a $10-\mathrm{m}$ Nansen bottle with reversing thermometer to estimate the temperature, and made eight vertical plankton tows to a wire length of 70 m and back. The positions, date, and time of arrival at the station and the $10-\mathrm{m}$ temperature are listed in Appendix Table 1. To maintain position on the station (relative to the water) as closely as possible, the officers of the vessel were instructed
to keep the $45-\mathrm{kg}$ weight on $10-\mathrm{m}, 3 / 16^{\prime \prime}$ cable vertical between casts and tows for the entire duration of the station. The usual interval between vertical tows was 12 minutes, but occasionally repeat tows and delays between tows lengthened that period. The distribution of intervals between adjacent tows is depicted in a histogram in Appendix Table 2; this average interval will be used to interpret all tows.

## The Tow

Of the 12 minutes between the initiation of tows, $3-5$ minutes were involved with lowering and raising the net, another 5 minutes was required to wash the net and concentrate the sample in the cod end, and 4-5 minutes was needed to preserve the sample in Formalin and label the sample bottle. The winch on the charter vessel was incapable of obtaining the $70-\mathrm{m}-$ per-minute towing rate specified for the project, and rates were commonly in the range of 25 to 40 m per minute.

The plankton net was based on the original design of anchovy egg net (Smith et al. 1968) and modified for vertical towing (Hewitt 1983; Smith et al. 1984). As the original net in the vertical tow series, it differed from the present design by being a single net, with mesh aperture area of $0.333-\mathrm{mm}$ nylon mesh. The mesh-aperture-to-mouth-aperture ratio was 8.7:1, of which $35 \%$ was in the terminal conical portion of the net and $65 \%$ was in the cylindrical portion to reduce length and facilitate self-cleaning under tow. Nets were washed from the outside by a moderate rate of flow.

## Laboratory Work

Anchovy eggs were sorted from samples with a dissecting microscope at a magnification of about $10 \times$ (Kramer et al 1972; Smith and Richardson 1977). The sorted eggs were subsequently staged using criteria of Moser and Ahlstrom (1984). Ages were estimated from the temperature-specific stage development rate (Ahlstrom 1943; Zweifel and Lasker 1976; Lo 1984), the actual time of tow, and the 10-m temperature on the station.

## Data Analysis

Descriptive statistics were performed on the UCSDVAX using Minitab. A correlation matrix on all ages of egg for all 60 tows was performed, yielding 7 correlation coefficients for adjacent columns, 6 for tows 24 minutes apart, 5 for tows 36 minutes apart, etc. Negative binomial parameter estimates were accomplished using the iterative techniques described in Southwood (1978). The BASIC program implemented on the UCSD-VAX to estimate the negative binomial parameters is listed in the Appendix.

TABLE 1
Replicate Observations of Anchovy Eggs < One Day Old

| Row | Replicate Number |  |  |  |  |  |  |  | Average |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 1 | 1 | 2 | 4 | 9 | 23 | 3 | 15 | 64 | 15.125 |
| 2 | 0 | 0 | 1 | 0 | 0 | 0 | 1 | 0 | 0.250 |
| 3 | 0 | 0 | 0 | 0 | 0 | 2 | 3 | 17 | 2.750 |
| 4 | 4 | 5 | 0 | 8 | 7 | 8 | 10 | 12 | 6.750 |
| 5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 6 | 4 | 2 | 7 | 0 | 0 | 0 | 0 | 0 | 1.625 |
| 7 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 8 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.125 |
| 9 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 10 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 11. | 1 | 0 | 0 | 0 | 1 | 0 | 1 | 0 | 0.375 |
| 12 | 0 | 0 | 0 | 0 | 1 | 3 | 0 | 5 | 1.125 |
| 13 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 14 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 15 | 30 | 39 | 51 | 0 | 0 | 0 | 0 | 0 | 15.000 |
| 16 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 17 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0.125 |
| 18 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 19 | 13 | 6 | 20 | 6 | 4 | 6 | 0 | 0 | 6.875 |
| 20 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 21 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 22 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |

## RESULTS

We will consider three aspects of the data. The first will be the data set itself. Secondly, we will describe the data and a possible probability-generating distribution from which it may have been drawn. Thirdly, we will consider one interpretation of the change of the data with time.

## The Data Set

The data set consists of four ages of egg, and the distribution of total eggs. The four ages of egg are those eggs less than 8 hours old, 1 day old, 2 days old, and 3 days old. These sets are from the same 8 replicates of 60 observations containing at least one anchovy egg per station.

Eggs produced in the first 8 hours after the onset of spawning are considered separately in the analysis of egg production because incidence and abundance are underestimated. (Smith and Hewitt 1984) In Table 1, it may be seen that of the 60 samples considered, only 22 were in the initial 8 -hour period. In no case does a large sample observation continue across the entire set of eight observations on the station. For example, in row 1, replicates 5, 7, and 8 indicate that the ship drifted into a patch, whereas rows 15 and 19 indicate that the ship drifted out of a patch.

The chief difference for one-day-old eggs (Table 2) is that in most observations, the set of replicates is usually taken entirely within or outside of a patch. There are still instances, such as row 49 , that indicate drifting out, but most show moderate changes within replicates on station. There are no visible differences in Tables 3 or 4 from Table 2 in terms of continuity of observations on station.

TABLE 2
Replicate Observations of Anchovy Eggs One Day Old

| Row | Replicate Number |  |  |  |  |  |  |  | Average |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 1 | 3 | 3 | 3 | 6 | 2 | 2 | 1 | 3 | 2.875 |
| 2 | 2 | 1 | 3 | 4 | 3 | 3 | 0 | 5 | 2.625 |
| 3 | 8 | 5 | 7 | 7 | 6 | 10 | 5 | 13 | 7.625 |
| 4 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 6 | 0 | 1 | 2 | 0 | 4 | 2 | 1 | 2 | 1.500 |
| 7 | 12 | 10 | 10 | 6 | 5 | 3 | 4 | 6 | 7.000 |
| 8 | 5 | 1 | 6 | 6 | 4 | 2 | 10 | 5 | 4.875 |
| 9 | 0 | 0 | 0 | 2 | 0 | 0 | 0 | 1 | 0.375 |
| 10 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 11 | 9 | 3 | 2 | 2 | 0 | 0 | 3 | 0 | 2.375 |
| 12 | 12 | 23 | 12 | 4 | 7 | 5 | 7 | 5 | 9.375 |
| 13 | 0 | 1 | 0 | 2 | 3 | 3 | 1 | 1 | 1.375 |
| 14 | 0 | 1 | 0 | 0 | 1 | 2 | 0 | 0 | 0.500 |
| 15 | 2 | 5 | 2 | 3 | 0 | 1 | 2 | 0 | 1.875 |
| 16 | 6 | 8 | 3 | 6 | 2 | 6 | 4 | 5 | 5.000 |
| 17 | 3 | 3 | 1 | 2 | 3 | 2 | 0 | 4 | 2.250 |
| 18 | 2 | 0 | 1 | 0 | 1 | 2 | 2 | 2 | 1.250 |
| 19 | 18 | 26 | 21 | 33 | 21 | 32 | 22 | 23 | 24.500 |
| 20 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 21 | 1 | 3 | 3 | 3 | 2 | 0 | 1 | 0 | 1.625 |
| 22 | 2 | 2 | 9 | 5 | 4 | 4 | 9 | 8 | 5.375 |
| 23 | 0 | 1 | 1 | 1 | 0 | 1 | 3 | 1 | 1.000 |
| 24 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 25 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 26 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 27 | 1 | 0 | 0 | 1 | 0 | 4 | 5 | 1 | 1.500 |
| 28 | 2 | 1 | 0 | 0 | 0 | 0 | 1 | 1 | 0.625 |
| 29 | 4 | 2 | 5 | 0 | 3 | 3 | 0 | 4 | 2.625 |
| 30 | 15 | 5 | 13 | 16 | 10 | 13 | 12 | 6 | 11.250 |
| 31 | 14 | 12 | 11 | 11 | 9 | 7 | 8 | 5 | 9.625 |
| 32 | 0 | 0 | 2 | 1 | 0 | 0 | 0 | 0 | 0.375 |
| 33 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 34 | 0 | 0 | 0 | 1 | 0 | 0 | 1 | 0 | 0.250 |
| 35 | 2 | 0 | 0 | 1 | 0 | 1 | 0 | 0 | 0.500 |
| 36 | 1 | 2 | 0 | 3 | 0 | 0 | 0 | 0 | 0.750 |
| 37 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 0.125 |
| 38 | 17 | 11 | 22 | 26 | 37 | 27 | 20 | 30 | 23.750 |
| 39 | 5 | 8 | 12 | 4 | 6 | 4 | 3 | 3 | 5.625 |
| 40 | 3 | 2 | 3 | 2 | 0 | 3 | 3 | 1 | 2.125 |
| 41 | 0 | 1 | 0 | 2 | 2 | 0 | 0 | 1 | 0.750 |
|  |  |  |  |  |  |  |  |  | (continued) |

TABLE 2 (continued)

| Row | Replicate Number |  |  |  |  |  |  |  | Average |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 42 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 43 | 0 | 1 | 0 | 0 | 0 | 0 | 0 | 1 | 0.250 |
| 44 | 1 | 4 | 0 | 3 | 5 | 3 | 4 | 3 | 2.875 |
| 45 | 7 | 4 | 4 | 1 | 3 | 2 | 2 | 3 | 3.250 |
| 46 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0.125 |
| 47 | 1 | 1 | 2 | 1 | 1 | 3 | 1 | 5 | 1.875 |
| 48 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 49 | 3 | 22 | 3 | 0 | 0 | 0 | 0 | 0 | 3.500 |
| 50 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 51 | 0 | 0 | 1 | 0 | 0 | 0 | 0 | 0 | 0.125 |
| 52 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 53 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 54 | 5 | 4 | 3 | 0 | 3 | 2 | 2 | 3 | 2.750 |
| 55 | 0 | 0 | 1 | 0 | 2 | 0 | 0 | 0 | 0.375 |
| 56 | 1 | 2 | 3 | 2 | 4 | 3 | 3 | 4 | 2.750 |
| 57 | 64 | 65 | 64 | 63 | 48 | 82 | 54 | 48 | 61.000 |
| 58 | 1 | 0 | 0 | 2 | 1 | 1 | 5 | 3 | 1.625 |
| 59 | 11 | 20 | 5 | 20 | 20 | 12 | 22 | 18 | -16.000 |
| 60 | 5 | 10 | 8 | 5 | 5 | 7 | 4 | 7 | 6.375 |

TABLE 3
Replicate Observations of Anchovy Eggs Two Days Old

| Row | Replicate Number |  |  |  |  |  |  |  | Average |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 1 | 0 | 1 | 0 | 1 | 0 | 2 | 1 | 1 | 0.750 |
| 2 | 0 | 2 | 0 | 1 | 1 | 1 | 1 | 1 | 0.875 |
| 3 | 0 | 1 | 2 | 0 | 1 | 1 | 0 | 1 | 0.750 |
| 4 | 2 | 0 | 2 | 1 | 1 | 0 | 2 | 1 | 1.125 |
| 5 | 0 | 2 | 1 | 1 | 3 | 0 | 1 | 3 | 1.375 |
| 6 | 3 | 5 | 5 | 6 | 8 | 15 | 14 | 12 | 8.500 |
| 7 | 2 | 2 | 0 | 4 | 2 | 2 | 1 | 0 | 1.625 |
| 8 | 6 | 12 | 11 | 8 | 12 | 9 | 10 | 9 | 9.625 |
| 9 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 10 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 11 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 0.125 |
| 12 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 13 | 1 | 1 | 3 | 0 | 0 | 0 | 3 | 2 | 1.250 |
| 14 | 2 | 3 | 5 | 0 | 6 | 7 | 4 | 10 | 4.625 |
| 15 | 1 | 2 | 2 | 0 | 4 | 2 | 1 | 2 | 1.750 |
| 16 | 5 | 2 | 6 | 2 | 2 | 1 | 3 | 2 | 2.875 |
| 17 | 0 | 0 | 0 | 2 | 4 | 4 | 4 | 4 | 2.250 |
| 18 | 1 | 1 | 2 | 0 | 1 | 1 | 1 | 2 | 1.125 |
| 19 | 6 | 14 | 11 | 12 | 12 | 9 | 5 | 9 | 9.750 |
| 20 | 1 | 2 | 2 | 1 | 0 | 3 | 1 | 1 | 1.375 |
| 21 | 7 | 5 | 7 | 2 | 2 | 5 | 1 | 5 | 4.250 |
| 22 | 26 | 32 | 46 | 30 | 17 | 9 | 14 | 13 | 23.375 |
| 23 | 1 | 2 | 2 | 2 | 0 | 0 | 2 | 0 | 1.125 |

TABLE 3 (continued)

| Row | Replicate Number |  |  |  |  |  |  |  | Average |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 24 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0.125 |
| 25 | 0 | 0 | 0 | 1 | 0 | 0 | 0 | 0 | 0.125 |
| 26 | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 0 | 0.125 |
| 27 | 1 | 0 | 6 | 4 | 2 | 2 | 10 | 3 | 3.500 |
| 28 | 3 | 2 | 6 | 4 | 1 | 0 | 3 | 6 | 3.125 |
| 29 | 34 | 35 | 40 | 30 | 36 | 39 | 35 | 35 | 35.500 |
| 30 | 7 | 3 | 1 | 2 | 3 | 4 | 1 | 7 | 3.500 |
| 31 | 22 | 10 | 13 | 11 | 15 | 21 | 9 | 12 | 14.125 |
| 32 | 3 | 3 | 2 | 3 | 3 | 3 | 3 | 0 | 2.500 |
| 33 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 34 | 4 | 1 | 1 | 3 | 1 | 0 | 0 | 0 | 1.250 |
| 35 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 36 | 0 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0.125 |
| 37 | 2 | 0 | 1 | 0 | 0 | 0 | 0 | 0 | 0.375 |
| 38 | 5 | 4 | 3 | 15 | 21 | 15 | 9 | 13 | 10.625 |
| 39 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 40 | 3 | 3 | 1 | 0 | 2 | 0 | 2 | 3 | 1.750 |
| 41 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 42 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.125 |
| 43 | 1 | 1 | 0 | 1 | 0 | 0 | 1 | 2 | 0.750 |
| 44 | 4 | 4 | 9 | 5 | 2 | 8 | 2 | 11 | 5.625 |
| 45 | 0 | 0 | 0 | 0 | 0 | 3 | 2 | 2 | 0.875 |
| 46 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 1 | 0.250 |
| 47 | 39 | 21 | 39 | 21 | 18 | 48 | 16 | 22 | 28.000 |
| 48 | 0 | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 0.125 |
| 49 | 0 | 2 | 1 | 0 | 2 | 0 | 2 | 0 | 0.875 |
| 50 | 6 | 4 | 4 | 10 | 7 | 4 | 8 | 5 | 6.000 |
| 51 | 2 | 4 | 6 | 4 | 1 | 2 | 4 | 2 | 3.125 |
| 52 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 53 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 54 | 27 | 22 | 21 | 0 | 8 | 11 | 4 | 4 | 12.125 |
| 55 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 56 | 22 | 14 | 6 | 9 | 6 | 4 | 6 | 1 | 8.500 |
| 57 | 96 | 117 | 94 | 89 | 84 | 119 | 111 | 91 | 100.125 |
| 58 | 12 | 3 | 9 | 6 | 3 | 4 | 5 | 9 | 6.375 |
| 59 | 8 | 6 | 2 | 4 | 4 | 2 | 9 | 5 | 5.000 |
| 60 | 1 | 5 | 6 | 4 | 4 | 5 | 12 | 10 | 5.875 |

Table 5 contains the values for all ages of anchovy egg within each station set. At these temperatures, one may expect three ages of egg within close proximity. In this set, only rows 33,52 , and 53 had none of the three ages identified: row 10 had neither one- nor two-day-old eggs, row 35 had neither two- nor three-day-old eggs, and rows $24,26,42$, and 50 had neither one- nor three-day-old eggs.

In summary, there is coherence among the replicate stations of anchovy eggs older than eight hours. This indicates that the scale of the pattern is large relative to the drift of the research vessel and the layers of water below over periods of one to two hours.

TABLE 4
Replicate Observations of Anchovy Eggs
Three Days Old

| Row | Replicate Number |  |  |  |  |  |  |  | Average |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 2 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 3 | 3 | 4 | 4 | 2 | 1 | 3 | 0 | 2 | 2.375 |
| 4 | 1 | 0 | 4 | 3 | 3 | 1 | 0 | 2 | 1.750 |
| 5 | 3 | 4 | 5 | 4 | 4 | 4 | 1 | 11 | 4.500 |
| 6 | 3 | 0 | 2 | 1 | 2 | 2 | 2 | 2 | 1.750 |
| 7 | 2 | 6 | 9 | 4 | 3 | 3 | 3 | 2 | 4.000 |
| 8 | 1 | 1 | 5 | 2 | 3 | 5 | 5 | 5 | 3.375 |
| 9 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 10 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0.125 |
| 11 | 5 | 4 | 0 | 3 | 1 | 1 | 4 | 9 | 3.375 |
| 12 | 9 | 11 | 8 | 5 | 8 | 1 | 5 | 5 | 6.500 |
| 13 | 1 | 2 | 2 | 1 | 1 | 3 | 3 | 2 | 1.875 |
| 14 | 2 | 4 | 5 | 3 | 2 | 7 | 2. | 4 | 3.625 |
| 15 | 1 | 0 | 1 | 0 | 0 | 0 | 0 | 1 | 0.375 |
| 16 | 8 | 5 | 3 | 5 | 10 | 6 | 8 | 3 | 6.000 |
| 17 | 9 | 3 | 3 | 0 | 1 | 0 | 1 | 1 | 2.250 |
| 18 | 1 | 0 | 3 | 0 | 1 | 0 | 3 | 0 | 1.000 |
| 19 | 17 | 16 | 15 | 20 | 10 | 10 | 11 | 5 | 13.000 |
| 20 | 4 | 5 | 8 | 4 | 3 | 7 | 5 | 2 | 4.750 |
| 21 | 13 | 7 | 6 | 4 | 9 | 7 | 7 | 3 | 7.000 |
| 22 | 24 | 16 | 16 | 10 | 8 | 9 | 4 | 6 | 11.625 |
| 23 | 1 | 3 | 1 | 2 | 0 | 4 | 1 | 2 | 1.750 |
| 24 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 25 | 0 | 3 | 0 | 0 | 0 | 0 | 0 | 0 | 0.375 |
| 26 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 27 | 5 | 0 | 9 | 6 | 6 | 5 | 11 | 10 | 6.500 |
| 28 | 1 | 1 | 0 | 1 | 1 | 0 | 1 | 0 | 0.625 |
| 29 | 21 | 17 | 15 | 23 | 14 | 11 | 14 | 17 | 16.500 |
| 30 | 20 | 6 | 18 | 22 | 9 | 11 | 3 | 9 | 12.250 |
| 31 | 6 | 4 | 5 | 4 | 5 | 2 | 7 | 1 | 4.250 |
| 32 | 1 | 0 | 0 | 1 | 0 | 1 | 1 | 0 | 0.500 |
| 33 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 34 | 0 | 0 | 1 | 0 | 0 | 1 | 0 | 0 | 0.250 |
| 35 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 36 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 37 | 0 | 1 | 1 | 1 | 1 | 0 | 0 | 0 | 0.500 |
| 38 | 3 | 4 | 5 | 4 | 3 | 2 | 4 | 5 | 3.750 |
| 39 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 40 | 0 | 0 | 0 | 1 | 0 | 0 | 1 | 1 | 0.375 |
| 41 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 42 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | ${ }^{\circ} 0$ | 0.000 |
| 43 | 4 | 8 | 4 | 7 | 8 | 5 | 3 | 11 | 6.250 |
| 44 | 2 | 5 | 5 | 2 | 5 | 5 | 6 | 12 | 5.250 |
| 45 | 0 | 1 | 0 | 0 | 1 | 0 | 2 | 2 | 0.750 |
| 46 | 0 | 0 | 0 | 0 | 2 | 1 | 2 | 0 | 0.625 |
| 47 | 25 | 29 | 32 | 43 | 41 | 53 | 52 | 45 | 40.000 |
| 48 | 1 | 2 | 4 | 1 | 0 | 1 | 1 | 0 | 1.250 |
| 49 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0.250 |
| 50 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |

TABLE 4 (continued)

| Row | Replicate Number |  |  |  |  |  |  |  | Average |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 51 | 61 | 69 | 75 | 84 | 76 | 62 | 61 | 68 | 69.500 |
| 52 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 53 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 54 | 10 | 10 | 12 | 0 | 5 | 8 | 2 | 5 | 6.500 |
| 55 | 120 | 86 | 119 | 107 | 136 | 114 | 104 | 102 | 111.000 |
| 56 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.000 |
| 57 | 22 | 26 | 16 | 34 | 16 | 28 | 42 | 26 | 26.250 |
| 58 | 1 | 2 | 2 | 6 | 1 | 5 | 3 | 3 | 2.875 |
| 59 | 3 | 8 | 5 | 2 | 6 | 3 | 3 | 3 | 4.125 |
| 60 | 7 | 12 | 12 | 14 | 9 | 12 | 15 | 17 | 12.250 |

TABLE 5
Replicate Observations of Anchovy Eggs All Ages Plus Disintegrated

| Row | Replicate Number |  |  |  |  |  |  |  | Average |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 1 | 4 | 6 | 7 | 16 | 25 | 7 | 17 | 68 | 18.750 |
| 2 | 2 | 3 | 3 | 5 | 4 | 4 | 1 | 6 | 3.500 |
| 3 | 14 | 11 | 13 | 10 | 8 | 14 | 5 | 16 | 11.375 |
| 4 | 4 | 0 | 8 | 5 | 4 | 1 | 4 | 3 | 3.625 |
| 5 | 3 | 6 | 6 | 6 | 7 | 4 | 2 | 14 | 6.000 |
| 6 | 6 | 6 | 9 | 7 | 14 | 19 | 17 | 16 | 11.750 |
| 7 | 16 | 18 | 19 | 14 | 10 | 10 | 11. | 25 | 15.375 |
| 8 | 17 | 19 | 22 | 24 | 26 | 24 | 35 | 31 | 24.750 |
| 9 | 0 | 0 | 0 | 2 | 0 | 0 | 0 | 1 | 0.375 |
| 10 | 0 | 0 | 0 | 0 | 0 |  | 0 | 2 | 0.375 |
| 11 | 15 | 7 | 2 | 5 | 1 | 2 | 7 | 10 | 6.125 |
| 12 | 21 | 34 | 21 | 9 | 16 | 5 | 12 | 10 | 16.000 |
| 13 | 2 | 4 | 7 | 4 | 4 | 11 | 8 | 8 | 6.000 |
| 14 | 4 | 8 | 10 | 3 | 9 | 16 | 6 | 14 | 8.750 |
| 15 | 4 | 7 | 5 | 4 | 4 | 3 | 3 | 3 | 4.125 |
| 16 | 19 | 15 | 12 | 13 | 14 | 13 | 15 | 10 | 13.875 |
| 17 | 19 | 9 | 13 | 7 | 15 | 7 | 5 | 11 | 10.750 |
| 18 | 9 | 9 | 18 | 12 | 8 | 13 | 10 | 7 | 10.750 |
| 19 | 56 | 74 | 63 | 74 | 53 | 64 | 48 | 44 | 59.500 |
| 20 | 6 | 7 | 12 | 6 | 3 | 11 | 7 | 3 | 6.875 |
| 21 | 22 | 15 | 16 | 9 | 13 | 12 | 9 | 8 | 13.000 |
| 22 | 57 | 58 | 76 | 50 | 32 | 27 | 30 | 30 | 45.000 |
| 23 | 2 | 6 | 7 | 5 | 0 | 10 | 7 | 4 | 5.125 |
| 24 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 0.125 |
| 25 | 1 | 3 | 0 | 2 | 0 | 0 | 0 | 0 | 0.750 |
| 26 | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 0 | 0.125 |
| 27 | 8 | 14 | 15 | 11 | 9 | 11 | 27 | 14 | 13.625 |
| 28 | 9 | 6 | 6 | 5 | 3 | 0 | 6 | 7 | 5.250 |
| 29 | 59 | 54 | 60 | 54 | 53 | 54 | 50 | 58 | 55.250 |
| 30 | 42 | 15 | 32 | 37 | 22 | 28 | 20 | 22 | 27.250 |
| 31 | 42 | 26 | 29 | 26 | 30 | 33 | 24 | 23 | 29.125 |
| 32 | 4 | 3 | 5 | 6 | 4 | 4 | 4 | 5 | 4.375 |
| 33 | 3 | 0 | 0 | 2 | 1 | 0 | 1 | 0 | 0.875 |

TABLE 5 (continued)

|  |  | Replicate Number |  |  |  |  |  |  |  |  |  |  |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | :---: | :---: | :---: |
| Row | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | Average |  |  |  |
| 34 | 4 | 1 | 2 | 4 | 1 | 1 | 1 | 0 | 1.750 |  |  |  |
| 35 | 2 | 0 | 0 | 1 | 0 | 1 | 0 | 0 | 0.500 |  |  |  |
| 36 | 1 | 3 | 0 | 3 | 1 | 4 | 3 | 2 | 2.125 |  |  |  |
| 37 | 2 | 1 | 2 | 1 | 1 | 0 | 1 | 0 | 1.000 |  |  |  |
| 38 | 56 | 58 | 81 | 48 | 64 | 49 | 34 | 50 | 55.000 |  |  |  |
| 39 | 5 | 8 | 12 | 4 | 6 | 4 | 3 | 3 | 5.625 |  |  |  |
| 40 | 6 | 5 | 4 | 3 | 2 | 3 | 6 | 6 | 4.375 |  |  |  |
| 41 | 0 | 1 | 0 | 2 | 2 | 0 | 0 | 1 | 0.750 |  |  |  |
| 42 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.125 |  |  |  |
| 43 | 5 | 10 | 4 | 8 | 8 | 5 | 4 | 14 | 7.250 |  |  |  |
| 44 | 7 | 13 | 14 | 10 | 12 | 16 | 12 | 26 | 13.750 |  |  |  |
| 45 | 20 | 11 | 24 | 7 | 8 | 11 | 6 | 7 | 11.750 |  |  |  |
| 46 | 0 | 1 | 2 | 1 | 2 | 3 | 4 | 2 | 1.875 |  |  |  |
| 47 | 66 | 52 | 73 | 65 | 60 | 104 | 72 | 73 | 70.625 |  |  |  |
| 48 | 1 | 2 | 4 | 1 | 0 | 2 | 2 | 0 | 1.500 |  |  |  |
| 49 | 4 | 25 | 4 | 4 | 8 | 4 | 4 | 1 | 6.750 |  |  |  |
| 50 | 6 | 4 | 4 | 10 | 7 | 4 | 8 | 5 | 6.000 |  |  |  |
| 51 | 63 | 73 | 82 | 88 | 77 | 64 | 65 | 70 | 72.750 |  |  |  |
| 52 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.125 |  |  |  |
| 53 | 0 | 0 | 1 | 0 | 0 | 0 | 0 | 1 | 0.250 |  |  |  |
| 54 | 47 | 39 | 43 | 1 | 18 | 23 | 9 | 13 | 24.125 |  |  |  |
| 55 | 120 | 87 | 122 | 109 | 138 | 116 | 104 | 102 | 112.250 |  |  |  |
| 56 | 23 | 16 | 9 | 11 | 10 | 8 | 10 | 6 | 11.625 |  |  |  |
| 57 | 182 | 208 | 174 | 187 | 148 | 229 | 208 | 165 | 187.625 |  |  |  |
| 58 | 15 | 6 | 11 | 16 | 8 | 10 | 13 | 16 | 11.875 |  |  |  |
| 59 | 29 | 39 | 13 | 27 | 31 | 20 | 37 | 29 | 28.125 |  |  |  |
| 60 | 13 | 27 | 27 | 23 | 19 | 24 | 32 | 34 | 24.875 |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |  |

## Data Summary

Owing to the wide range of numbers of eggs per sample, Tables 6,7 , and 8 are divided into classes whose boundaries encompass factors of four rather than unit frequencies. The general appearance of similarity among the replicate sets above is confirmed in the data summary table and the parameters calculated from them. The primary difference among the distributions of the eight replicates and the distribution of the mean of eight is the lower threshold ( 0.125 rather than 1 per $.05 \mathrm{~m}^{2}$ ) for the mean of eight samples on a station. Whereas one would expect the mean of the means to be the same as the mean of the replicates, one would expect the variance of the observations to be eight times the variance of the means of eight; instead, the variance of the means of eight is indistinguishable from the variances of the individual sets for one-, two- and three-day-old eggs.

The estimates of the negative binomial factor $k$ are the same from replicate to replicate and between the means of eight and the eight replicates (Tables 6, 7, and 8 ).Thus the estimation of the negative binomial parameters, mean, and $k$ from small sets of observations with high variance is relatively robust when small sample sizes are used. Furthermore, the negative binomial $k$ is well estimated from the fraction of zero observations, and even the "moments" estimate (from the mean and variance) is reasonably close to the others and relatively stable under these conditions.

The distribution of the total eggs (Table 9) differs

TABLE 6
Anchovy Egg Sample Frequency Distribution One-Day-Old Eggs

| Eggs/0.05 m2 | Replicate Number |  |  |  |  |  |  |  | Mean |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 0 | 26 | 24 | 27 | 25 | 29 | 27 | 27 | 25 | 13 |
| 0.0625-0.25 | - | - | - | - | - | - | - | - | 3 |
| 0.25-1 | - | - | - | - | - | - | - | - | 10 |
| 1-4 | 17 | 19 | 18 | 19 | 15 | 20 | 17 | 17 | 20 |
| 4-16 | 14 | 12 | 12 | 11 | 12 | 10 | 12 | 14 | 10 |
| 16-64 | 2 | 4 | 2 | 5 | 4 | 2 | 4 | 4 | 4 |
| 64-256 | 1 | 1 | 1 | - | - | 1 | - | - | - |
| Mean | 4.1 | 4.6 | 4.1 | 4.3 | 3.9 | 4.3 | 3.8 | 3.9 | 4.1 |
| Variance | 83.5 | 99.0 | 85.9 | 97.8 | 70.7 | 136.9 | 69.1 | 64.2 | 81.4 |
| Neg. binomial |  |  |  |  |  |  |  |  |  |
| $k(0)$ | . 321 | . 274 | . 277 | . 260 | . 283 | . 269 | . 327 | . 285 | . 312 |
| $k(1)$ | . 158 | . 195 | . 163 | . 157 | . 118 | . 145 | . 155 | . 170 | . 202 |
| $k(2)$ | . 264 | . 256 | . 254 | . 217 | . 234 | . 237 | . 252 | . 244 | . 296 |

$\boldsymbol{k}(0)$ is the parameter of the negative binomial distribution as estimated from the proportion of ' 0 ' values.
$\boldsymbol{k}(1)$ is the parameter of the negative binomial distribution as estimated from the sample mean and variance ('moments' estimate).
$k(2)$ is the maximum likelihood estimator (Southwood 1966).

TABLE 7
Anchovy Egg Sample Frequency Distribution Two-Day-Old Eggs

| Eggs/0.05 m2 | Replicate Number |  |  |  |  |  |  |  | Mean |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 0 | 24 | 22 | 24 | 27 | 24 | 26 | 20 | 22 | 10 |
| 0.0625-0.25 | - | - | - | - | - | - | - | - | 7 |
| 0.25-1 | - | - | - | - | - | - | - | - | 8 |
| 1-4 | 18 | 21 | 16 | 14 | 19 | 15 | 21 | 18 | 17 |
| 4-16 | 11 | 12 | 15 | 15 | 12 | 15 | 16 | 17 | 14 |
| 16-64 | 6 | 4 | 4 | 3 | 4 | 3 | 2 | 2 | 3 |
| 64-256 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| Mean | 6.1 | 5.9 | 6.3 | 5.0 | 5.0 | 6.1 | 5.4 | 5.4 | 5.7 |
| Variance | 213.2 | 265.7 | 222.0 | 163.8 | 148.8 | 292.4 | 225.0 | 163.8 | 198.8 |
| Neg. binomial |  |  |  |  |  |  |  |  |  |
| $k(0)$ | . 299 | . 347 | . 295 | . 268 | . 329 | . 262 | . 417 | . 363 | . 318 |
| $k(1)$ | . 180 | . 134 | . 184 | . 157 | . 174 | . 130 | . 133 | . 184 | . 157 |
| $k(2)$ | . 248 | . 276 | . 255 | . 241 | . 301 | . 227 | . 327 | . 316 | . 280 |

$k(0)$ is the parameter of the negative binomial distribution as estimated from the proportion of ' 0 ' values.
$k(1)$ is the parameter of the negative binomial distribution as estimated from the sample mean and variance ('moments' estimate).
$k(2)$ is the maximum likelihood estimator (Southwood 1966).
from that of any of the individual nights' spawning. In the individual age groups the frequency of observations decreases from the " 0 ' class to the " $64-256$ " eggs per observation class. In the total eggs observations, the frequency increases from the " 0 " class to the " $4-16$ " class and then descends. Even when each station has at least one egg in eight samples, the
number of observations with " 0 " eggs in each of the replicates varied from $10 \%$ to $18 \%$.

We concluded that nearly uniform results would have been obtained from any set of replicates. Also, the frequency distribution of the total is a composite of the day-class observations and may in fact exhibit a different type of distribution when three days' egg

TABLE 8
Anchovy Egg Sample Frequency Distribution Three-Day-Old Eggs

| Eggs/0.05 m2 | Replicate Number |  |  |  |  |  |  |  | Mean |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 0 | 22 | 25 | 24 | 25 | 24 | 25 | 23 | 24 | 15 |
| 0.0625-0.25 | - | - | - | - | - | - | - | - | 1 |
| 0.25-1 | - | - | - | - | - | - | - | - | 10 |
| 1-4 | 19 | 11 | 10 | 15 | 17 | 14 | 19 | 17 | 13 |
| 4-16 | 11 | 17 | 20 | 13 | 15 | 17 | 14 | 13 | 16 |
| 16-64 | 7 | 5 | 4 | 5 | 2 | 3 | 3 | 4 | 3 |
| 64-256 | 1 | 2 | 2 | 2 | 2 | 1 | 1 | 2 | 2 |
| Mean | 7.0 | 6.4 | 7.3 | 7.3 | 6.9 | 6.7 | 6.7 | 6.8 | 6.9 |
| Variance | 316.8 | 216.1 | 334.9 | 346.0 | 412.1 | 316.8 | 295.8 | 278.9 | 302.8 |
| Neg. binomial |  |  |  |  |  |  |  |  |  |
| $k(0)$ | . 321 | . 274 | . 277 | . 260 | . 283 | . 269 | . 327 | . 285 | . 283 |
| $k(1)$ | . 158 | . 195 | . 163 | . 157 | . 118 | . 145 | . 155 | . 170 | . 157 |
| $k(2)$ | . 264 | . 256 | . 254 | . 217 | . 234 | . 237 | . 252 | . 244 | . 257 |

$k(0)$ is the parameter of the negative binomial distribution as estimated from the proportion of ${ }^{\prime} 0$ ' values.
$k(1)$ is the parameter of the negative binomial distribution as estimated from the sample mean and variance ('moments' estimate).
$k(2)$ is the maximum likelihood estimator (Southwood 1966).

TABLE 9
Anchovy Egg Sample Frequency Distribution All Ages Plus Disintegrated

| Eggs/0.05 m2 | Replicate Number |  |  |  |  |  |  |  | Mean |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |  |
| 0 | 7 | 10 | 11 | 6 | 10 | 11 | 10 | 9 | - |
| 0.0625-0.25 | - | - | - | - | - | - | - | - | 4 |
| 0.25-1 | - | - | - | - | - | - | - | - | 7 |
| 1-4 | 12 | 9 | 6 | 12 | 11 | 9 | 9 | 12 | 7 |
| 4-16 | 21 | 24 | 25 | 27 | 23 | 23 | 25 | 20 | 26 |
| 16-64 | 17 | 13 | 12. | 10 | 12 | 12 | 12 | 14 | 12 |
| 64-256 | 3 | 4 | 6 | 5 | 4 | 5 | 4 | 5 | 4 |
| Mean | 19.1 | 18.9 | 20.1 | 17.8 | 17.1 | 18.5 | 17.1 | 18.3 | 18.4 |
| Variance | 979.7 | 1049.8 | 1049.8 | 1004.9 | 852.6 | 1303.2 | 1024.0 | 829.4 | 961.0 |

production is present. This may be expected to differ for eggs that hatch in a day or persist for weeks.

## Interpretation of Time Series

The correlation between recent observations of newly spawned eggs ends at intervals greater than 24 minutes, whereas one-day-old eggs have a persistent correlation over the entire 84 -minute interval between the first and eighth tows (Figure 2). The coherence among tows increases gradually with the age of eggs. Coefficients of correlation over .24 , or coefficients of determination over .06 are significant with 60 observations.

The time and space scales of controlled ship's drift in 1-2 hours are small in comparison with the change of distribution of a patch of eggs more than one day old. The scale of deposition of the eggs and their subsequent dispersal for a few hours is of significantly smaller scale. Lastly, since eggs are part of the totally passive plankton, these results suggest that other passive plankton patches, or gaps, would have been equally coherent and slowly changing with time, and that aggregating organisms would have had to expend relatively little energy in maintaining a patch under these conditions.

## DISCUSSION

The unusual circumstances of this set of observations have led to a new appreciation for the origin of small-scale distribution of anchovy eggs and the threeday persistence of this distribution. The scale and intensity of this pattern may have important consequences, both for comparative sampling of the habitats of the eastern boundary currents, and for interpreting predation. Because this set of samples was taken for other purposes, we will discuss the relations between these observations and similar observations, the implications of the persistence of pattern, and what
sampling design and effort are necessary to advance the pattern studies to an analytical level.

## Sampling Consequences

The high variability of plankton samples has been attributed to variations in towing procedure (Windsor and Walford 1936); heterogeneous water masses (Windsor and Clarke 1940; Cassie 1959); and aggregations of organisms (Ricker 1937; Langford 1938; Barnes and Marshall 1951). In particular, sardine eggs, which last in the plankton only a few days, are known to be aggregated at spawning (Silliman 1946; Taft 1960) and diffuse away from school-sized (tens to hundreds of meters diameter) patches, which persist for several days (Smith 1973). Aggregations of organisms and heterogeneous water masses occur at several scales and may persist over various time periods (Haury et al. 1978), and our perception of the pattern is profoundly influenced by frequency and duration of cruises, the spacing between stations, and the width and trajectory of the sampling instruments as well as the length of tow (Weibe 1972).

Number of samples. The required number of observations can be determined for a first approximation from an existing set (Santander et al. 1982) or from a pilot cruise. In general, pattern is small-scale relative to the distribution of the species, so a pilot study of spawning pattern can be conducted in a small area ( $100 \times 100 \mathrm{~km}$ ). The results of this study indicate that one must be separated from an area by either space ( km ) or time (days) to gather a valid replicate. versus a redundant sample. The same is true of the pilot study conducted on existing sample results.

If a study requires precise data on a single day's spawning, the variance of the total eggs will be an underestimate of the individual day's spawning by the ratio of the persistence of the egg and the sample variance of the total eggs: for an egg that persists 3


Figure 2. The persistence of correlation between tows as a function of time between tows. Points are from individual values of coelficient of determination for adjacent tows ( 7 sets), tows separated by 24 minutes ( 6 sets), tows separated by 36 minutes ( 5 sets), etc. The lines connect the medians of each interval. (For simplicity, only the nominal interval is graphed; the actual intervals varied from 6 to 36 minutes, with $92 \%$ of intervals between 9 and 17 minutes and $\mathbf{5 1 \%}$ between 11 and 13 minutes. For complete distribution see Appendix Table 2.)
days, the variance of the total eggs can be multiplied by three to estimate the required number of samples for a given standard error. If the study requires 200 observations to suitably estimate the total number of eggs, it would require 346 samples to estimate the number of eggs spawned in a single day. To describe the onset of spawning between 1800 and 0200 hours with equal precision, it would take a similar number of samples each hour.

Geographic position of samples. Anchovy spawning habitats seem to vary considerably in different eastern boundary currents. The spawning habitat off North America comprises broad regions in relatively
permanent gyrals. The spawning habitat off Peru is relatively narrow (Santander et al. 1982; Smith et al. 1983); the spawning habitat off South Africa appears to be a fast-moving coastal jet (Shelton and Hutchings 1982). The current practice is to make 1,000 observations per survey off California (Smith and Hewitt 1984). This may be the minimum required when one considers that the sample must encompass the spawning area, obtain a representative mean, and provide an estimate of egg mortality. Where the spawning habitat is too narrow to provide sufficiently independent adjacent samples, as in Peru, it may be necessary to occupy the habitat more than once. Where the habitat

TABLE 10

| Ocean and Atmospheric Predictability Time for <br> Different Scales of Motion |  |  |
| :---: | :---: | :---: |
| Scale | Predictability time |  |
|  | Atmosphere | Ocean |
| 10 m | - | 10 min. |
| 100 m | 3 min. | 1.5 hr |
| 1 km | 13 min. | 10 hr |
| 10 km | 1 hr | 3 days |
| 100 km | 4 hr | 3 weeks |
| 1000 km | 1 day | 4 months |
| 10000 km | 5.5 days | $\times$ years |

Modified from Platt et al. (1977)
is extremely dynamic, it may be necessary to expand the pattern with time to accommodate the rapidly moving jet and evaluate the survivors.

## Consequences for the Study of Pattern

The egg distribution of schooled, coastal, pelagic spawning fish sufficiently resembles coastal plankton blooms like red tide (Kierstad and Slobodkin 1953; Wroblewski 1984) to allow evaluation of the influence of turbulent diffusion and transport at spawning time (Lasker 1975; Bakun and Parrish 1982). The 60 samples reported here ( 480 observations in samples of 8 per station) are not sufficiently numerous to describe even mortality, so we may expect the number of samples necessary to exceed 200 if interannual variations are relatively large. The assembly of 5 years of egg distribution observations ( 1,666 positive samples) was adequate to estimate mortality and dispersal (Smith and Hewitt 1984). There was no evidence of offshore transport in the 3-day period for which eggs persist: transport may be inferred from the distribution of older versus younger larvae (Smith 1972 [sardine, Fig. 6; anchovy, Fig. 71; Hewitt and Methot 1982 [anchovy, Tab. 4, Fig. 15]). Because of this latter phenomenon, higher volumes will have to be filtered over longer distances than used for the vertical tows, to effectively describe transport. Also, the observations must be extended over time to describe transport in the mesoscale.

## Predictability in the Eastern Boundary

A new process of modeling, empirical measurements specified by models, and modification of models based on new measurements is beginning (Bakun et al. 1982). For eastern boundary currents the models will have an atmospheric component, an oceano-
graphic component, and a series of biological effects. We must remember that some causative factors in the air and ocean will remain unpredictable over certain space and time scales; thus biological responses will remain unpredictable. The nature of the problem of matching these time and space scales may be seen in Table 10. It seems reasonable to conclude from correlation of replicate samples for eggs less than 8 hours old that the pattern which yielded this result was on the order of hundreds of meters or the space scale of a fish school (perhaps in motion during spawning). Thus there is a new opportunity to design processoriented cruises with the goal of determining the most important time and space scales of organisms of known age and distributional heterogeneity.

## LITERATURE CITED

Ahlstrom, E.H. 1943. Influence of temperature on the rate of development of pilchard eggs in nature. In O.E. Sette, (ed.), Studies on the Pacific pilchard or sardine (Sardinops caerulea). U.S. Dept. of Interior Spec. Sci. Rep. 15. Reissued 1950.
. 1968. An evaluation of the fishery resources available to California fishermen. Univ. Wash. Publ. Fish. (New Series) 4:65-80.
Bakun, A., and R.H. Parrish. 1982. Transport, turbulence, and pelagic fish in the Califormia and Peru current systems. CalCOFI Rep. 24:99. 112.

Bakun, A., J. Beyer, D. Pauly, J.G. Pope, and G.D. Sharp, 1982. Ocean sciences in relation to living resources. Can. J. Fish. Aquat. Sci. 39:1059-1070.
Barnes, H., and S.M. Marshall. 195I. On the variability of replicate plankton samples and some applications of restricted periods. J. Mar. Biol. Assoc. U.K. 30:233-263.
Cassie, R.M., 1959. Micro-distribution of plankton. N.Z.J. Science 2:398-409.
Haury, L.R., J.A. McGowan, and P.H. Wiebe. 1978. Pattems and processes in the time-space scales of plankton distributions. In J.H. Steele (ed.), Spatial pattern in plankton communities. Plenum Publ. Corp. New York, p. 277-327.
Hempel, G. 1973. Fish eggs and larval studies (contribution to a manual). FAO Fish Tech. Paper. 122, 82 p.
Hewitt, R.P. 1981. The value of pattern in the distribution of young fish. Rapp. P.-v. Réun. Cons. Int. Explor. Mer 178:229-236.
_ 1982. Spatial pattern and survival of anchovy larvae: implications of adult reproductive strategy. Ph.D. dissertation, University of California, San Diego, 187 p.

- 1983. Roll, heave and vertical ichthyoplankton tows. Ocean Science and Engineering 8:41-51.
Hewitt, R.P., and R.D. Methot, Jr. 1982. Distribution and mortality of northern anchovy larvae in 1978 and 1979. CalCOFI Rep., 23:226-245.
Kierstad, H., and L.B. Slobodkin. 1953. The size of water masses containing plankton blooms. J. Mar. Res. 12:141-147.
Kramer, D., M.J. Kalin, E.G. Stevens, J.R. Thraillkill, and J. Zweifel. 1972. Collecting and processing data on fish eggs and larvae in the California Current region. NOAA Tech. Rep. NMFS Circ. 370.
Langford, R.R. 1938. Diumal and seasonal changes in the distribution of the limnetic crustacea of Lake Nipissing, Ontario. Publ. Ontario Fish. Res. Lab., 56:1-142.
Lasker, R. 1975. Field criteria for survival of anchovy larvae: the relation between inshore chlorophyll maximum layers and successful first feeding. Fish. Bull., U.S. 73:453-462.
_1. 1984. An egg production method for anchovy biomass assessment. In An egg production method for estimating spawning biomass of pelagic fish: application to the northern anchovy (Engraulis mordax). pelagic fish: application to the northern anchovy (E
Southwest Fisheries Center Admin. Rep. LJ-84-37.
- 1985. What limits clupeoid production?. Can. J. Fish. Aquat. Sci. 42 (Suppl. 1):31-38.

Lasker, R., and J.R. Zweifel. 1978. Growth and survival of patches containing different proportions of large and small prey. In J.H. Steele (ed.). Spatial pattem in plankton communities. Plenum Publishing Corp., New York, p. 329-354.
Lo. N.C.H. 1984. A model for temperature-dependent northem anchovy egg development and an automated procedure for the assignment of age to staged eggs. In R. Lasker (ed.). An egg production method for estimating spawning biomass of pelagic fish: application to the northern anchovy (Engraulis mordax) Southwest Fisheries Center Admin. Rep. LJ-84-37.
Moser, H.G., and E.H. Ahlstrom. 1984. Staging anchovy eggs. In R. Lasker, (ed.). An egg production method for estimating spawning biomass of pelagic fish: application to the northern anchovy (Engraulis mordax) Southwest Fisheries Center Admin. Rep. LJ-84-37.
Murphy, G.I. 1977. Clupeoids. In J.A. Gul]and (ed., Fish population dynamics. John Wiley \& Sons, New York, p. 283-308.
Okubo, A. 1971. Oceanic diffusion diagrams. Deep-Sea Res. 18:789-802.
Parrish, R.H., and A.D. MacCall. 1978. Climatic variation and exploitation in the Pacific mackerel fishery. Calif. Fish Game Fish Bull. 167:1110.

Parrish, R.H., C.S. Nelson, and A. Bakun. 1981. Transport mechanisms and reproductive success of fishes in the California Current. Biol. Oceanog. 1(2):175-203.
Platt, T., K.L. Denman, and A.D. Jassby, 1977. Modeling the productivity of phytoplankton. In E.D. Goldberg (ed.). The sea: ideas and observations on progress in the study of the seas. Vol. Vl. John Wiley \& Sons, New York, p. 807-856.
Ricker, W.E. 1937. Statistical treatment of sampling processes useful in enumeration of plankton organisms. Arch. Hydrobiol. 31:68-84.
Santander, H., P. E. Smith, and J. Alheit. 1982. Determinación del esfuerzo de muestreo requierido para el estimado de producción de huevos de anchoveta (Engraulis ringens), al frente al Peru. Bol. Inst. Mar. Peru Callao 7(1):1-18.
Sette, O.E., and E.H. Ahlstrom. 1948. Estimations of abundance of the eggs of the Pacific pilchard (Sardinops caerulea) off southem California during 1940 and 1941. J. Mar. Res. 7:511-542.
Shelton, P.A., and L. Hutchings. 1982. Transport of anchovy, Engraulis capensis Gilchrist, eggs and early larvae by a frontal jet current. J. Cons. Int. Explor. Mer 40:185-198.

Silliman, R.P. 1946. A study of variability in plankton tow-net catches of Pacific pilchard, Sardinops caerulea, eggs. J. Mar. Res. 6:74-83.
Smith, P.E. 1972. The increase in spawning biomass of northern anchovy, Engraulis mordax. Fish. Bull. U.S. 70:849-874.
. 1973. The mortality and dispersal of sardine eggs and larvae. Rapp. P.-v. Réun. Cons. Int. Explor. Mer 164:282-292.
1981. Sampling to determine anchovy larval mortality in the sea. In R. Lasker (ed.), Marine fish larvae. Univ. Wash. Press, Seattle, p. 12-21.

Can. J. Fish. Aquat. Sci. 42 (Suppl. 1):69-82.
Can. J. Fish. Aquat. Sci, 42 (Suppl. 1).69-82. egg production method of anchovy biomass assessment. In R. Lasker (ed.), An egg production method for estimating spawning biomass of pelagic fish: application to the northern anchovy (Engraulis mordax) Southwest Fisheries Center Admin. Rep. LJ-84-37.
Smith, P.E., and R. Lasker. 1978. Position of larval fish in an ecosystem. Rapp. P.-v. Réun. Cons. Int. Explor. Mer 173:77-84.
Smith, P.E., and S.L. Richardson. 1977. Standard techniques for pelagic fish egg and larva surveys. FAO Fisheries Tech. Pap. 175, 100 p.
Smith, P.E., R.C. Counts, and R.I. Clutter. 1968. Changes in filtering efficiency of plankton nets due to clogging under tow. J. Cons. Perm. Int. Explor. Mer 32:232-248.
Smith, P.E., W.C. Flerx, and R.P. Hewitt. 1984. The CalCOFI vertical egg tow (CalVET) net. /n R. Lasker (ed). An egg production method for estimating spawning biomass of pelagic fish: application to the northern anchovy (Engraulis mordax). Southwest Fisheries Center Admin. Rep. LJ-84-37.
Smith, P.E., H. Santander, and J. Alheit. 1983. Comparison of egg sample probability distributions of the anchovy (Engraulis ringens) and sardine (Sardinops sagax) off Peru and the anchovy (Engraulis mordax) and the sardine (Sardinops caerulea) off California. In G.D. Sharp and J. Csirke (eds.), Proceedings of the expert consultation to examine changes in abundance and species composition of neritic fish resources. FAO Fish. Rep. 291 Vol 3:1027-1038.
Southwood, T.R.E. 1966 . Ecological methods. Meihuen, London, 391 p. - 1978. Ecological methods. John Wiley \& Sons, New York, 524 p.
Taft, B.A. 1960. A statistical study of the estimation of abundance of sardine (Sardinops caerulea) eggs, Limnol. Oceanog. 5:254-264.
Vlymen, W.J. 1977. A mathematical model of the relationship between larval anchovy, Engraulis mordax, growth, prey microdistribution, and larval behaviour. Environ. Biol. Fishes 2:21|-233.
Wiebe, P.H. 1972. A field investigation of the relationship between length of tow, size of net and sampling error. J. Cons. Perm. Int. Explor. Mer 34:268-275.
Winsor, C.P. and G.L. Clarke. 1940. A statistical study of variation in the catch of plankton nets. J. Mar. Res. 3:1-34.
Winsor, C.P. and L.A. Walford. 1936. Sampling variations in the use of plankton nets. J. Cons. Int. Explor. Mer 11:190-204.
Wroblewski, J. 1984. Formulation of growth and mortality of larval northern anchovy in a turbulent feeding environment. Mar. Ecol. Prog. Ser. 20:1-12.
Zweifel. J.R., and R. Lasker. 1976. Prehatch and posthatch growth of fishes-a general model. U:S. Fish. Bull. 74:609-621.
Zweifel, J.R., and P.E. Smith. 1981. Estimates of abundance and mortality of larval anchovy (1951-75): application of a new method. Rapp. P.-v. Réun. Cons. Int. Explor. Mer 178:248-259.

## APPENDIX

TABLE 1
Station Data for Replicate Series

| $\begin{aligned} & \text { CalCOFI } \\ & \text { line } \end{aligned}$ | $\begin{aligned} & \text { CalCOFI } \\ & \text { station } \end{aligned}$ | Date | Time | 10 -meter temperature |
| :---: | :---: | :---: | :---: | :---: |
| 90.0 | 28.0 | 790122 | 2043 | 14.8 |
| 82.0 | 47.0 | 790127 | 0810 | 13.1 |
| 83.0 | 42.0 | 790127 | 1254 | 13.6 |
| 83.0 | 40.6 | 790127 | 1832 | 13.5 |
| 87.0 | 35.0 | 790128 | 0645 | 13.7 |
| 89.6 | 34.6 | 790128 | 1419 | 14.1 |
| 93.0 | 30.0 | 790129 | 1855 | 13.8 |
| 93.0 | 35.0 | 790129 | 2313 | 14.1 |
| 93.0 | 40.0 | 790130 | 0337 | 14.4 |
| 93.0 | 45.0 | 790130 | 0757 | 13.2 |
| 89.5 | 41.0 | 790131 | 0959 | 13.2 |
| 88.4 | 40.6 | 790131 | 1413 | 13.5 |
| 89.2 | 32.9 | 790201 | 0813 | 13.6 |
| 90.1 | 32.0 | 790201 | 1150 | 13.7 |
| 90.8 | 35.1 | 790201 | 1549 | 14.0 |
| 91.7 | 39.8 | 790201 | 2057 | 13.9 |
| 90.6 | 42.8 | 790202 | 0127 | 14.0 |
| 88.9 | 45.3 | 790202 | 0632 | 12.9 |
| 87.6 | 44.7 | 790202 | 1031 | 13.2 |
| 86.3 | 44.8 | 790202 | 1905 | 13.0 |
| 86.1 | 42.6 | 790203 | 0010 | 13.1 |
| 84.3 | 43.4 | 790203 | 1108 | 13.0 |
| 83.8 | 47.2 | 790203 | 1444 | 13.1 |
| 83.9 | 52.3 | 790203 | 1839 | 13.1 |
| 89.4 | 51.2 | 790204 | 0947 | 13.4 |
| 91.9 | 44.2 | 790204 | 1855 | 13.3 |
| 92.9 | 38.1 | 790204 | 2346 | 13.6 |
| 91.9 | 32.8 | 790205 | 0423 | 13.6 |
| 92.2 | 27.7 | 790205 | 0817 | 13.5 |
| 93.5 | 27.4 | 790205 | 1142 | 14.0 |
| -93.8 | 31.8 | 790206 | 1945 | 14.3 |
| 91.9 | 40.6 | 790207 | 0728 | 13.8 |
| 90.8 | 46.5 | 790207 | 1222 | 13.2 |
| 90.5 | 44.2 | 790207 | 2127 | 13.8 |
| 92.2 | 42.9 | 790208 | 0845 | 14.1 |
| 93.1 | 45.2 | 790208 | 1717 | 13.8 |
| 91.8 | 45.8 | 790208 | 2104 | 13.9 |
| 92.7 | 40.9 | 790209 | 0126 | 13.9 |
| 94.9 | 50.2 | 790209 | 1802 | 13.7 |
| 96.7 | 50.2 | 790209 | 2213 | 14.3 |
| 96.0 | 45.9 | 790210 | 0319 | 14.2 |
| 94.4 | 38.6 | 790210 | 1106 | 13.9 |
| 93.3 | 34.7 | 790210 | 1526 | 14.4 |

TABLE 1 (continued)

| CalCOFI <br> line | CaICOFI <br> station | Date | Time | 10-meter <br> temperature |
| :---: | :---: | :---: | :---: | :---: |
| 92.8 | 31.6 | 790210 | 1859 | 13.9 |
| 91.2 | 30.2 | 790211 | 0040 | 12.4 |
| 91.9 | 34.3 | 790211 | 0540 | 14.1 |
| 91.2 | 37.8 | 790211 | 0939 | 14.7 |
| 89.7 | 36.5 | 790211 | 1400 | 14.7 |
| 88.9 | 40.7 | 790211 | 1739 | 13.8 |
| 88.2 | 43.9 | 790212 | 1519 | 14.0 |
| 87.8 | 39.1 | 790212 | 1907 | 13.3 |
| 86.4 | 38.5 | 790212 | 2300 | 13.7 |
| 85.6 | 45.8 | 790213 | 0749 | 13.2 |
| 82.5 | 43.3 | 790214 | 0640 | 13.4 |
| 83.8 | 40.6 | 790214 | 1050 | 13.8 |
| 85.2 | 37.2 | 790214 | 1452 | 13.9 |
| 86.5 | 33.9 | 790214 | 1920 | 13.8 |
| 89.4 | 30.6 | 790215 | 0320 | 13.8 |
| 90.2 | 29.3 | 790215 | 0655 | 14.0 |
| 91.7 | 28.0 | 790215 | 1050 | 14.9 |

TABLE 2
Distribution of Intervals between Tows

| Interval <br> between <br> tows <br> (min) | Proportion <br> of <br> observations |  |
| :---: | :---: | :--- |
| 6 | .011 | $*$ |
| 8 | .017 | $* *$ |
| 10 | .089 | $* * * * * * *$ |
| 12 | .511 | $* * * * * * * * * * * * * * * * * * * * * * * * * * * * * * * * * *$ |
| 14 | .260 | $* * * * * * * * * * * * * * * * * *$ |
| 16 | .060 | $* * * * *$ |
| 18 | .026 | $* *$ |
| 20 | 0 |  |
| 22 | .017 | $* *$ |
| 24 | 0 |  |
| 26 | .003 | $*$ |
| 28 | 0 |  |
| 30 | 0 |  |
| 32 | .003 | $*$ |
| 34 | 0 |  |
| 36 | 0 |  |
| 38 | .003 |  |

SMITH AND HEWITT: ANCHOVY EGG DISPERSAL AND MORTALITY CalCOFI Rep., Vol. XXVI, 1985

TABLE 3
Program to Estimate Parameters of a Negative Binomial Distribution



[^0]:    'Henrik Ibsen. An Enemy of the People. An adaptation for the American stage by Arthur Miller. Dramatists Play Service, Inc., New York, 1951.

