# THE INCREASE IN SPAWNING BIOMASS OF NORTHERN ANCHOVY, ENGRAULIS MORDAX 

Paul E. Smith ${ }^{1}$


#### Abstract

The northern anchovy, Engraulis mordax, is a common fish off the west coast of North America. Its biomass has increased markedly since 1951. The various methods of deriving anchovy spawner biomass from the sardine spawner biomass and analogous census estimates of sardine and anchovy larvae are reviewed. A new compilation of anchovy and sardine larval data is presented for 1940, 1941, 1949, 1950, and 1951-69. The effect of several errors of estimate are examined and it is concluded that none is important enough to affect measurably the trend of increase between 1951 and 1966. Lastly, a current interpretation of the larval survey data is used to estimate the spawning biomass of both sardine and anchovy.


The northern anchovy (Engraulis mordax Girard) is a common pelagic schooling fish off the west coast of North America between British Columbia and Baja California (lat $53^{\circ} \mathrm{N}$ to $22^{\circ} \mathrm{N}$ ). The fishery for northern anchovy has usually been small relative to the Pacific sardine (Sardinops caerulea (Girard)) off California but the anchovy catch has been increasing since the late 1930's (Figure 1). Following the collapse of the sardine fishery in the early 1950 's, the catch of anchovy exceeded that of sardine. The further decline of sardine catch and the eventual moratorium on sardine, combined with a limited reduction fishery on anchovy, have again allowed the anchovy fishery to exceed that of sardine since the mid-1960's (Messersmith and Associates, 1969).

Routine planktonic larva sampling since 1951 shows an increase in the number of anchovy larvae (Murphy, 1966; Ahlstrom, 1966, 1968) and the present consensus estimate of the spawning population of northern anchovy is about 5 million tons for 1966, the last year for which complete data are available. The same sample data for the Pacific sardine now indicate an extremely small number of larvae and the spawning bio-

[^0]mass of the northern subpopulation of the sardine may now be less than 5,000 tons. All anchovy biomass estimates must now be referred to analogous sardine biomass estimates based on the sardine fishery and on sardine and anchovy egg and larva surveys because no fishery-based estimate of anchovy biomass has yet been made: this becomes increasingly difficult and imprecise as the sardine "reference" population diminishes.

The purposes of this paper are to:

1) review the estimates of spawner biomass of the anchovy;
2) present a summary of the incidence of larval anchovy by region and season since 1951;
3) examine the effects of three spawning behavior models on the estimates of spawning biomass;
4) establish a standard reference period for the determination of anchovy biomass without further consideration of the current size of the sardine population.

## PREVIOUS ESTIMATES OF ANCHOVY BIOMASS

Messersmith and Associates (1969, p. 9) tabulated all the estimates of anchovy spawning biomass for the years 1940-66. Ahlstrom (MRC,


Figure 1.-The annual catch of sardine (crosses) and anchovy (solid circles) in the California Current system since 1916. Whole fish reduction was essentially banned in 1919 and a limited whole fish reduction fishery was opened in 1965.
1964) ${ }^{2}$ in an informal statement, emphasized the changing ratio of anchovy larvae to sardine larvae from 3.1 to 1 in 1951 to 46.8 to 1 in 1959. In a similarly informal statement, MacGregor (MRC, 1964) described a method for estimating the spawning biomass of pelagic spawning species. That portion of his statement which is related to sardine and anchovy follows:

Egg and larval surveys conducted over the years have given us a basis for estimating the total numbers of eggs and larvae produced each year in the CalCOFI

[^1](California Cooperative Oceanic Fisheries Investigations) survey area for a number of fish species. We also have information on the fecundity of a number of these species and from the combined data can estimate the biomass of spawning adult fish on each species. The number of eggs a fish will produce at one spawning appears to be directly related to the weight of the fish. Thus, we can compute the numbers of eggs produced in one spawning by a ton of female fish as follows:
\[

$$
\begin{array}{lc}
\text { Species } & \text { Millions of eggs } \\
\text { Anchovy } & 525 \\
\text {. . . } & . \\
\text { Sardine } & 24 \dot{1} \\
\text {. . . . } & \text {. . }
\end{array}
$$
\]

The above figures represent female fish only. If we assume, as evidence indicates, that for each ton of adult females there is also present a ton of adult males, we would have to divide the number of eggs produced
by 2 to obtain the number of eggs produced [in one spawning] ${ }^{3}$ by one ton of adult fish. If we further assume that . . . . about half the [adult] sardine population spawns twice [a year] (average $11 / 2$ times) and the anchovy [adult] population spawns 2 or 3 times a year (average $21 / 2$ times), we would also multiply millions of eggs by $11 / 2$ for the sardine and $21 / 2$ for the anchovy. On the above basis the total number of eggs produced by one ton of adult fish of both sexes in one year would be as follows for each species:

| Species | Millions of eggs |
| :--- | :---: |
| Anchovy | 656 |
| Sardine | . |
| . . . . | .${ }^{\text {. }}$ |

It may be seen from the above that, although numbers of eggs or larvae in the plankton may be used as an index of adult abundance, a sardine egg in the plankton represents $31 / 2$ times . . . as much adult biomass as an anchovy egg. Estimates of the average biomass of each . . . species for the 3-year period 1955-57 and the average commercial landings for the same period are as follows:

|  | Biomass | Catch |
| :--- | :---: | :---: |
| Species | (tons) | (tons) |
| Anchovy* | 750,000 | 29,686 |
| - . . | Mardine | 254,000 |

- Anchovy larvae caught were used as an estimate of eggs. This may have resulted in underestimation by a factor of 2. Application of this actor brings this biomass of anchovies into line with the material in
${ }^{3}$ In brackets added for clarification.

In another informal statement Radovich (MRC, 1965) cited the change in anchovy to sardine larval ratios (Ahlstrom, MRC, 1964) and stated ". . . in this 8-year period the anchovy maintained its population level fifteen times better than the sardine . . ." He further pointed out that the necessary ratio of annual survival rates to attain this was 1.4. Should this difference in survival be in the larval stage, the MacGregor (MRC, 1964) biomass estimate should be divided by 1.4 or roughly 1.5 , resulting in an estimate of about 1 million tons.

Murphy (MRC, 1964) used an anchovy spawner biomass estimate which was essentially double MacGregor's (MRC, 1964) because he believed that the escapement of anchovy larvae through the mesh of the standard CalCOFI silk net exceeded the escapement of the sardine larvae enough to cancel the effect of anchovy fecundity exceeding sardine fecundity. Thus Murphy's calculation (1966, p. 60) of anchovy spawner biomass for the period 1955-57 was 3.3 million short tons as compared to MacGregor's 1.5 million short tons and Radovich's 1.0 million short tons for the same 3 -year period. An estimate by Murphy's method for 1958 would be 5.1 million short tons.

Ahlstrom (1968) estimated that the 1958 anchovy biomass was between 1.80 and 2.25 million short tons and observed further that the biomass had reached a plateau of 4.5 to 5.625 million short tons in the mid-1960's.

Table 1.-Comparison of data used for estimating the spawning biomass of northern anchovy.

|  | Standard houl summation ${ }^{1}$ |  | Summation of overage quarterly estimates 2 |  | Larval census estimates ${ }^{3}$ |  | Standard haul summations* |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Anchovy | Sardine | Anchovy | Sardine | Anchovy | Sardine | Anchovy | Sardine |
| 1951 | 29,551 | 1.1,068 | 9,826 | 3,689 | 15.101 | 5.774 | 29,552 | 11,066 |
| 1952 | 59,626 | 19,179 | 20,581 | 6,437 | 17,071 | 5,466 | 63,057 | 24,559 |
| 1953 | 99,160 | 14,400 | 34,314 | 4,924 | 23,680 | 4,020 | 103,928 | 15,055 |
| 1954 | 161,241 | 26,914 | 56,685 | 9,364 | 38,415 | 7,297 | 161,254 | 26,914 |
| 1955 | 140,183 | 14,121 | 51,096 | 5,554 | 37,658 | 4,341 | 140,183 | 14,121 |
| 1956 | 134,931 | 15,523 | 51,438 | 5.179 | 38,508 | 3,895 | 134,931 | 15,523 |
| 1957 | 146,631 | 9,833 | 53,921 | 3,415 | 40,441 | 2,432 | 146,631 | 9,833 |
| 1958 | 205,733 | 11,427 | 75,120 | 3,845 | 56,928 | 2,831 | 205,457 | 11,423 |
| 1959 | 206.753 | 5,374 | 72,732 | 2,072 | 54,168 | 1,159 | 206,000 | 5,308 |
| 1960 | 289,860 | 8,012 | 97,602 | 3,099 |  |  |  |  |
| 1961 | 97,103* | 1,708* | 97,103 | 1,708 |  |  |  |  |
| 1962 | 212,675* | 2,258** | 212,675 | 2,258 | ${ }^{1}$ Ahlstrom | ables 2 and | 1965, 1966 | ers from cur- |
| 1963 | 205,838* | 1,349** | 205,838 | 1,349 | rent unpublish | ). |  |  |
| 1964 | 166,517* | 2,757** | 166,517 | 2,757 | ${ }^{2}$ Ahlstrom. <br> a Ahlstrom. |  |  |  |
| 1965 | 258,781* | 3,573* | 258,781 | 3,603 | - Murphy. |  |  |  |
| 1966 | 380,420 | 5,640 | 161,333 | 2,211 | * Quarterly | rather than | thly. |  |

Table 2.-Stations and pooled areas within each region as used in this study.

| Regional name | Pooled areas | Stations | Regional name | Pooled areas | Stations |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Central California Instrore | 6.5 | 60.52 |  |  | 87.80 |
| 18 stations | 5,270 miles $^{2}$ | 60.55 |  | 8.9 | 80.90 |
| 4 pooled areas |  | 63.52 |  |  | 83.90 |
| 19,970 miles $^{2}$ |  | 63.55 |  |  | 87.90 |
|  |  | 67.50 |  | 9.7 | 9070 |
|  |  | 67.55 |  | 9.7 | 93.70 |
|  | 6.6 | 60.60 |  |  | 97.70 |
|  | 4,800 miles ${ }^{2}$ | 63.60 |  | 9.8 | 90.80 |
|  |  | 67.60 |  |  | 93.80 |
|  | 7.5 | 70.52 (51) |  |  | 97.80 |
|  | 5,100 miles ${ }^{2}$ | 70.55 (53) |  | 9.9 | 90.90 |
|  |  | 73.51 (50) |  |  | 93.90 |
|  |  | 73.55 (53) |  |  | 97.96 |
|  |  | $\begin{aligned} & 77.50(51) \\ & 77.55 \end{aligned}$ | Baja California inshore | 10.3 | 100.29 |
|  | 7.6 | 77.55 70.60 | 29 stations | 5,535 miles ${ }^{2}$ | 100.30 |
|  | 4,800 miles ${ }^{\text {a }}$ | 73.60 | 4 pooled areas |  | 100.35 |
|  |  | 77.60 | 21,089 miles ${ }^{\text {a }}$ |  | 103.30 |
|  |  |  |  |  | 103.35 |
| Central California offshore | 6.7 | 60.70 |  |  | 107.32 |
| 6 stations | 6.8 | 60.80 |  |  | 107.35 |
| 6 pooled areas | 6.9 | 60.90 |  | 10.4 | 100.40 |
| 28,800 miles $^{2}$ | 7.7 | 70.70 |  | 4,800 miles $^{2}$ | 103.40 |
|  | 7.8 | 70.80 |  |  | 103.45 |
|  | 7.9 | 70.80 |  |  | 107.40 |
| Southern California inshore | 8.4 | 82.47 |  | 11.3 | 110.33 (32) |
| 19 stations | 4,589 miles $^{2}$ | 83.40 |  | 5,956 miles ${ }^{\text {8 }}$ | 111.35 |
| 3 pooled areas |  | 83.43 |  |  | 113.30 |
| 15,348 miles $^{2}$ |  | 87.35 |  |  | 1173.35 |
|  |  | 87.40 |  |  | 117.26 |
|  |  | 87.45 |  |  | 117.30 |
|  | 9.3 | 90.28 |  |  | 117.35 |
|  | 5,989 miles ${ }^{2}$ | 90.30 (32) |  |  | 118.39 |
|  |  | 90.37 |  |  | 119.33 |
|  |  | 93.27 (28) |  |  | 120.25 |
|  |  | 93.30 |  |  | 120.30 |
|  |  | 97.30 |  |  | 120.35 |
|  |  | 97.32 |  | $\frac{11.4}{4.4}$ | 110.40 |
|  |  | 97.35 |  | 4,798 miles ${ }^{2}$ | 110.45 |
|  | 9.4 | 90.45 |  |  | 113.40 |
|  | 4,770 miles ${ }^{2}$ | 93.40 |  |  | 113.45 |
|  |  | 93.45 |  |  | 117.40 |
|  |  | 97.40 |  |  | 117.45 |
|  |  | 97.45 | Baia California offshore | 10.5 | 100.50 |
| Southern California offshore |  |  |  | 4,800 miles ${ }^{\text {a }}$ | 103.50 |
| 18 stations | $4,747 \text { miles }^{2}$ | $\begin{aligned} & 80.51 \\ & 80.55 \end{aligned}$ | 4 pooled areas 19.200 miles ${ }^{\text {a }}$ ( |  | 107.50 |
| 4 pooled areas | 4,747 miles ${ }^{2}$ | $\begin{aligned} & 80.55 \\ & 83.51 \end{aligned}$ | 19,200 miles ${ }^{\text {a }}$ | ${ }_{\text {cen }} 10.6$ | 100.60 |
| 19,147 miles $^{2}$ |  | 83.55 |  | 4,800 miles ${ }^{2}$ | 103.60 |
|  |  | 87.50 |  |  | 107.60 |
|  |  | 87.55 |  | 11.5 | 110.50 |
|  | 8.6 | 80.80 |  | 4,800 miles ${ }^{8}$ | 110.55 |
|  | 4,800 miles ${ }^{2}$ | 83.60 |  |  | 113.50 |
|  |  | 87.60 |  |  | 113.55 |
|  | 9.5 | 90.50 |  |  | 117.50 |
|  | 4,800 miles ${ }^{2}$ | 90.55 (53) |  |  | 117.55 |
|  |  | 93.50 |  | 11.6 | 110.60 |
|  |  | 93.55 |  | 4,800 miles ${ }^{2}$ | 1113.60 |
|  |  | 97.50 |  |  | 117.60 |
|  |  | 97.55 | Baja California seaward | 10.7 | 100.70 |
|  | 9.6 | 90.60 | 19 stations | 4,800 miles ${ }^{\text {a }}$ | 103.70 |
|  | 4,800 miles $^{3}$ | 93.60 | ${ }^{6}$ 6poled areas |  | 107.70 |
|  |  | -97.60 | 28,800 miles ${ }^{\text {a }}$ | 10.8 | 100.80 |
| Southern California seaward | 8.7 | 80.70 |  | 4,800 miles ${ }^{3}$ | 103.80 |
| 18 stations |  | 83.70 |  |  | 107.80 |
| 60 pooled areas |  | 87.70 |  | ${ }^{10.9}$ | 100.90 |
| 28,800 miles ${ }^{\text {a }}$ | 8.8 | 80.80 |  | 4,800 miles ${ }^{\text {s }}$ | 103.90 |
|  |  | 83.80 |  | 11.7 | 107.90 110.70 |

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Table 2.-Continued.

| Regional name | Pooled areas | Stations | Regional name | Pooled areas | Stations |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | 4,800 miles $^{2}$ | 113.70 |  | 4,800 miles $^{2}$ | 130.45 |
|  |  | 117.70 |  |  | 133.40 |
|  |  | 117.75 |  |  | 133.45 |
|  |  | 110.80 |  |  | 137.40 |
|  | 4,800 miles ${ }^{2}$ | 113.80 |  |  | 137.45 |
|  |  | 117.80 |  |  |  |
|  | $11.9$ | 110.90 | South Baia offshore | 12.5 | 120.50 |
|  | $4,800 \text { miles }^{2}$ | 113.90 | 16 stations | 4,800 miles ${ }^{2}$ | 120.55 |
|  |  | 117.90 | 4 pooled areas |  | 123.50 |
|  |  |  | 19,200 miles $^{2}$ |  | 123.55 |
| South Baja inshore | $12.3$ | 123.37 |  |  | 127.50 |
| 22 stations | $4,459 \text { miles }^{2}$ | 127.34 |  |  | 127.55 |
| 5 pooled areas | 12.4 | 1220.40 |  | 12.6 | 120.60 |
| 22,790 miles $^{3}$ | 4,800 miles ${ }^{\text {a }}$ | 120.45 123.42 |  | 4,800 miles $^{2}$ | 123.60 |
|  |  | 123.45 |  |  | 127.60 |
|  |  | 127.40 |  | 4,800 miles ${ }^{\text {a }}$ | 130.55 |
|  |  | 127.45 |  |  | 133.50 |
|  | 13.2 | 133.25 |  |  | 137.50 |
|  | 3,931 miles ${ }^{\text {a }}$ | 137.23 |  | 13.6 | 130.60 |
|  | 13.3 | 130.30 |  | 4,800 miles ${ }^{2}$ | 133.60 |
|  | 4,800 miles ${ }^{\text {a }}$ | 130.35 |  |  | 137,60 |
|  |  | 133.30 |  |  |  |
|  |  | 133.35 | South Baja seaward | 12.7 | 120.70 |
|  |  | 137.30 | 3 stations | 12.8 | 120.80 |
|  |  | 137.35 | 3 pooled areas | 12.9 | 120.90 |
|  | 13.4 | 130.40 | 14,400 miles ${ }^{2}$ |  |  |

Murphy (1966) estimated the biomass of the spawning stocks of sardine from 1932 to 1959. He also compared the anchovy:sardine larval ratio from 1951 to 1959 and graphically compared this with the larval ratio over a portion (about $20 \%$ ) of their joint range in 1940 and 1941 (p.65). The striking decline of the sardine, and the increase of the anchovy biomass, has stimulated speculation on the biological interactions of these species and the MRC (Marine Research Committee of California) has a standing recommendation that 200,000 tons of anchovies and 10,000 tons of sardines be harvested in an experimental attempt to foster the recovery of sardines. It is the primary goal of this recommendation to restore and maintain the balance of sardines and anchovies in the California Current system by manipulation of fishing effort.

The estimates of spawning biomass of the northern anchovy have been based on the spawning biomass of the Pacific sardine as derived from the fishery (Murphy, 1966; Ahlstrom, 1968) and on the assumed relationship between sardine and anchovy fecundity, survival, and escapement of larvae through the meshes of the CalCOFI standard silk survey net (Lenarz, 1972). For convenience, the larva data for all
estimates to date are listed in Table 1. A series of stations off southern California was occupied in 1940 and 1941, and the anchovy: sardine ratios were 1.18 ( 13,962 anchovy to 11,862 sardines) and 1.66 ( 12,560 anchovy to 7,564 sardines) respectively. The 1940, 1941, 1949, and 1950 data will be referred to later in a section on interaction of sardine and anchovy.

## METHOD OF DATA ASSEMBLY

The method of assembling the estimates of larval abundance for this paper differs from that of Sette and Ahlstrom (1948) and Ahlstrom (1954, 1966, 1967, 1968). Two methods of assembly were used previously: the "census estimate" and the "standard haul summation." In the "census estimate" each larva sample count was weighted by the area of a polygon formed by construction of "perpendicular bisectors of lines drawn from the station to each of all surrounding stations" (Sette and Ahlstrom, 1948, p. 521; Ahlstrom, 1968).

$$
\begin{equation*}
C_{k}=10 \sum_{j=1}^{n}\left[\sum_{i=1}^{m} A_{s}\left(a_{i}^{-2} b_{i}^{-1} c_{i} d_{i}\right)\right]_{j} \tag{1}
\end{equation*}
$$



Figure 2.-The selected stations of the CalCOFI grid (see Tables 2 and 3 ) and the regional boundaries defined for this paper. Columns of stations are indicated .30 , $.40 \ldots .90$, and rows of lines are indicated $60,70, \ldots 130$. The first two letters of each three-letter group stand for coastal bands, i.e., central California (CC), Southern California (SC), Baja California (BC), South Baja (SB) while the last letter of the three stands for offshore zones roughly parallel to the coast, inshore ( $0-80$ miles from the coast, " $I$ ", offshore ( $80-160$ miles from the coast, " $O$ "), and seaward (160-280 miles from the coast, "S"), so that the three letters SCI stand for the southern California inshore region.

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Table 3.-Description of CalCOFI regions used in this analysis.

| Regional name | Stations | Pooled <br> areas | Square <br> miles | Area <br> factor |  |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Central California inshore | 18 | 4 | 19,970 | $6.85 \times 10^{9}$ |  |  |  |  |  |
| Central California offshore | 6 | 6 | 28,800 | 9.88 |  |  |  |  |  |
| Southern California inshore | 19 | 3 | 15,348 | 5.26 |  |  |  |  |  |
| Southern California offshore | 18 | 4 | 19,147 | 6.57 |  |  |  |  |  |
| Southern California seaward | 18 | 6 | 28,800 | 9.88 |  |  |  |  |  |
| Baja California inshore | 29 | 4 | 21,089 | 7.23 |  |  |  |  |  |
| Baia California offshore | 15 | 4 | 19,200 | 6.58 |  |  |  |  |  |
| Baia California seaward | 19 | 6 | 28,800 | 9.88 |  |  |  |  |  |
| South Baja inshore | 22 | 5 | 22,790 | 7.82 |  |  |  |  |  |
| South Baja offshore | 16 | 4 | 19,200 | 6.58 |  |  |  |  |  |
| South Baia seaward | 3 | 3 | 14,400 | 4.94 |  |  |  |  |  |
| Totals |  |  |  |  |  | 183 | 49 | 237,544 | 81.48 |

Table 4.-Equivalent station list off southern California for 1940, 1941, and 1950-72.

| Southern California inshore |  |  | Southern California offshore |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Station no. used in 1940 | $\begin{aligned} & \text { Station } \\ & \text { no. } \\ & \text { used in } \\ & 1941 \end{aligned}$ | Nearest present station no. | $\begin{aligned} & \text { Station } \\ & \text { no. } \\ & \text { used. in } \\ & 1940 \end{aligned}$ | Station no. used in 1941 | Nearest present station no. |
| 02 | 12 | 82.47 | 04 | 15 | 80.51 |
| 01 | 11 | 83.40 | 05 | 16 | 80.55 |
| 08 | 23 | 83.43 | 03 | 13 | 83.51 |
| 10 | 21 | 87.35 | 13 | 14 | 83.55 |
| 17 | 22 | 87.40 | 24 | 24 | 87.50 |
| 11 | 34 | 87.45 | 12 | 25 | 87.55 |
| 21 | 30 | 90.28 | 06 | 28 | 80.60 |
| 19 | 31 | 90.30 | 09 | 27 | 83.60 |
| 20 | 32 | 90.37 | 18 | 26 | 87.60 |
| 22 | 41 | 93.27 | 23 | 35 | 90.50 |
| 28 | 51 | 90.30 | 31 | 36 | 90.55 |
|  | 60 | 97.30 | 39 | 45 | 93.50 |
| 35 | 61 | 97.32 | 40 | 46 | 93.55 |
| 36 | 62 | 97.35 | 41 | 64 | 97.50 |
| 30 | 33 | 90.45 | 14 | 55 | 97.55 |
| 29 | 52 | 93.40 | 32 | 37 | 90.60 |
| 34 | 54 | 93.45 | 25 | 47 | 93.60 |
| 37 | 63 | 97.40 | 33 | -- | 97.60 |
| 38 | 53 | 97.45 |  |  |  |

where $A_{s}=$ area of a polygon constructed of perpendicular bisectors of lines between station " $i$ " and all adjacent stations expressed as number of $10 \mathrm{~m}^{2}$ areas
$C_{k f}=$ estimate of abundance of larvae in year " $k$ " and " $f$ " takes the value of the number of the equation
$C_{i}=$ number of larvae in " $i$ th" sample
$a_{i}=$ area of mouth of the net used at the "ith" station
$b_{i}=$ length of tow in meters estimated from a calibrated flow meter at station " $i$ "
$d_{i}=$ tow depth in meters estimated
from the wire angle at maximum wire out at station " $i$ "
$m=$ number of stations $n=$ number of monthly cruises.
In the "standard haul summation" approach the sums of all tows on regularly occupied stations for each monthly cruise were totaled for the year without weighting for represented area.

$$
\begin{equation*}
C_{k}=10 \sum_{j=1}^{n}\left[\sum_{i=1}^{m}\left(a_{i}^{-2} b_{i}^{-1} c_{i} d_{i}\right)\right]_{j} \tag{2}
\end{equation*}
$$

To accommodate the quarterly cruises Ahlstrom $(1966,1968)$ established a modification of this in which the quarterly averages of 1 to 3 monthly cruise summations were added to make an
annual total (Ahlstrom, 1966, Table 8; 1968, Table 3).

$$
\begin{align*}
& C_{k}=10 \sum_{q=1}^{4}\left\{\sum _ { j = 1 } ^ { n } \left[\sum_{i=1}^{m}\right.\right. \\
& \left.\left.\times\left(a_{i}^{-2} b_{i}^{-1} c_{i} d_{i}\right)\right]_{j} n^{-1}\right\} . \tag{3}
\end{align*}
$$

Murphy (1966) used the monthly version of Ahlstrom's standard haul summation (Table 1).

The method used here is called the "regional census estimate"

$$
\begin{equation*}
C_{k 4}=10 A_{r} m^{-1}\left[\sum_{i=1}^{m}\left(a_{i}^{-2} b_{i}^{-1} c_{i} d_{i}\right)\right] \tag{4}
\end{equation*}
$$

where $C_{k 4}=$ estimate of abundance of larvae in region " $r$ " in each quarter
$A_{r}=$ Area of region " $r$ " in numbers of $10 \mathrm{~m}^{2}$ areas.

This method of assembly combines the simplicity of the "standard haul summation" and the areal weighting of the previously used census estimates. The regional census estimate consists of the mean number of larvae per standard area ( $10 \mathrm{~m}^{2}$ sea surface) of all stations taken within a region for a quarter of the year times the number of standard areas within the region. The same 183 stations or nearby alternates were used within the routinely occupied area as defined by Smith, Ahlstrom, and Casey (1970, Figure 1). The selected stations and acceptable alternates are listed in Table 2 and the regions are illustrated in Figure 2 and defined by area and area factor in Table 3.

The 1940 and 1941 stations have been assigned to the two southern California regions (Table 4) in which these cruises were conducted. The number of stations per unit area has been held nearly constant by the elimination of excess stations in the early cruises. The 1949 station equivalents are listed in Table 5. From 1950 to 1972 stations have been standard in placement.

Table 5.-Equivalent station list for the CalCOFI survey pattern for 1949 and 1950-72.

| Area | $\begin{gathered} \text { Station no. } \\ \text { Used in. } \\ 1949 \end{gathered}$ | Nearest present station nc. |
| :---: | :---: | :---: |
| Central | 601 | 63.58 |
| California inshore | 701 | 72.56 |
| Central | 602 | 62.68 |
| California | 603 | 61.78 |
| offshore | 604 | 61.87 |
|  | 702 | 71.66 |
|  | 703 | 71.76 |
|  | 704 | 71.85 |
| Southern Colifornia inshore | 901 | 92.39 |
| Southern | 801 | 82.57 |
| Colifornia | 902 | 92.48 |
| offshore | 903 | 92.58 |
| Southern | 802 | 82.67 |
| Californio | 803 | 82.77 |
| seaward | 804 | 82.87 |
|  | 904 | 92.68 |
|  | 905 | 92.78 |
| Baja | 1001 | 101.34 |
| California | 1002 | 101.44 |
| inshore | 1101 | W11.38 |
|  | 1102 | 111.48 |
| Baja | 1003 | 102.54 |
| California | 1004 | 102.64 |
| offshore | 1103 | 111.58 |
| Baia | 1005 | 102.74 |
| California | 1006 | 102.84 |
| seaward | 1007 | 102.94 |
|  | 1104 | 111.68 |
|  | 1105 | 112.78 |
|  | 1106 | 112.88 |
|  | 1107 | 112.98 |
| Southern | 1201 | 122.44 |
| Baja inshore |  |  |
| Southern 'Baja Offshore | 1202 | 122.53 |
| Southern | 1203 | 123.63 |
| Baja | 1204 | 123.73 |
| seaward | 1205 | 124.83 |

Extra cruises, additional lines, and stations on lines have been added periodically, resulting in some increase in station density in some areas. To stabilize sampling effort and remove possible effects of added effort nearshore, all but 183 stations and their nearby equivalents have been eliminated from further consideration for the 16 -year compilation of regional census estimates. The number of stations eliminated each year are summarized in Table 6.


Figure 3.-The size frequency of sardine larvae caught in the CalCOFI standard silk net $(0.55 \mathrm{~mm})$. The triangle dot-dash curve represents the 1951-60 catch from the selected stations in all eleven regions; the black dot-dash curve represents the 1940 catch from selected stations in the southern California inshore and offshore region and the clear dot solid line represents the 1941 catch from the same two regions. Relative catch refers to the catch at a standard size group (3.00, 3.25 (194041), $4.75,5.75 \ldots 15.75 \mathrm{~mm}$ ), divided by the average catch from size groups 8.75 to 15.75 .

## ESCAPEMENT AND AVOIDANCE BY ANCHOVY AND SARDINE LARVAE

The chief errors causing underestimates of $C$ for both anchovy and sardine larvae are those attributable to larvae "escaping" through the meshes of the standard plankton net (see Lenarz, 1972, p. 839) and larvae "avoiding" the mouth of the net (Silliman, 1943; Ahlstrom, 1954, 1959; Clutter and Anraku, 1968). While "escapement" and "avoidance" are important biases to consider in the study of larva growth and mortality, I consider them beyond the scope of this paper. The estimate of anchovy biomass is based on a relative estimate of the number of anchovy larvae and sardine larvae. I must treat escapement and avoidance briefly, since they act differently on the anchovy and sardine and vary from season to season and with changes in sampling gear.

Lenarz (1972) found no appreciable difference in size-specific escapement of sardine and anchovy through net apertures. However, newly hatched anchovy larvae are considerably smaller than sardine larvae. This leads to a variation in the degree of bias (Murphy, 1966) to such escapement. In Figure 3 the catch of all standard sizes of sardine is related to the average catch between 8.75 and 15.75 mm , a size range I assume to be completely retained on $0.55-\mathrm{mm}$ mesh width silk. The primary line is the average size composition for the period 1951-60 for the sardine. The size composition of the 1940

Table 6.-Stations eliminated from data assembly for regional census estimates.

| Year | No. of <br> stations <br> occupied | 183 stations <br> reported <br> eoch cruise | Stations <br> not <br> reported | Percent <br> reported |
| :---: | :---: | :---: | :---: | :---: |
| 1951 | 1,436 | 1,026 | 410 | 71.4 |
| 1952 | 1,376 | 1,167 | 209 | 84.8 |
| 1953 | 1,346 | 1.137 | 209 | 84.5 |
| 1954 | 1,473 | 1,217 | 256 | 82.6 |
| 1955 | 1,425 | 1,121 | 304 | 78.7 |
| 1956 | 1,399 | 1,129 | 270 | 80.7 |
| 1957 | 1,493 | 1,165 | 328 | 78.0 |
| 1958 | 1,851 | 1,276 | 575 | 68.9 |
| 1959 | 2,182 | 1,574 | 608 | 72.1 |
| 1960 | 1,810 | 1,305 | 505 | 72.1 |
| 1961 | 953 | 699 | 254 | 73.3 |
| 1962 | 920 | 659 | 261 | 71.6 |
| 1963 | 881 | 659 | 222 | 74.8 |
| 1964 | 877 | 680 | 197 | 77.5 |
| 1965 | 1,099 | 658 | 441 | 59.9 |
| 1966 | 1,979 | 1,487 | 492 | 75.1 |



Figure 4.-The size frequency of sardine and anchovy larvae. The black dot-dash line is sardine larvae caught in the standard CaICOFI silk net between 1951 and 1960 in all regions from January to June inclusive. The clear dot solid line is anchovy larvae caught in the standard CalCOFI silk net between 1951 and 1960, in all regions from January to June inclusive. The triangle dot-dash line is anchovy larvae caught in the new CalCOFI nylon net $(0.505-\mathrm{mm}$ nylon rather than $0.55-\mathrm{mm}$ silk) from January to June 1969 in all regions.
and 1941 sardine catches are included for later reference. In Figure 4, the 1951-60 size composition of sardine is compared to the anchovy for the same period. Anchovy retention in a new net with smaller, more regular meshes ( $0.505-\mathrm{mm}$ nylon) and more mesh area is included for comparison. All samples reported in the size frequency graphs are from the first half of the year to facilitate comparison with earlier samples and eliminate the effects of poor sampling coverage in the latter half of the year.

Ahlstrom (1954) acknowledged that larger larvae may avoid capture at night as in the daytime. Avoidance is more pronounced in
the daylight. In the sardine the night-to-day ratio of catches increases 0.6971 per mm of growth after 4.75 mm (Ahlstrom, 1954, p. 129). Similarly, the anchovy night-to-day catch ratio increases 0.64 for each millimeter growth after 3.5 mm (Ahlstrom, 1959, p. 136). Lenarz (in press) described the annual and diurnal variation in size specific catch rate for sardine, anchovy, hake, and jack mackerel. An important source of variability in avoidance bias is a shift of spawning season. For example, length of day varies from 9.6 hr in winter to 14.8 hr in summer at the latitude of San Francisco near the northern boundary of the survey grid and from 10.6 to 13.7 hr off south Baja California at the southern boundary of the sample grid ( $54 \%$ and $29 \%$ respectively). For the 10 -year period, 1951-60, the ratio of night-caught to day-caught larvae was 1.72 for sardines and 2.45:1 for anchovy. When the average catch per positive tow by month is corrected for day length at San Francisco the anchovy:sardine ratio changes from 2.96: 1 to 3.75:1 for the same decade. A proportionate shift of anchovy spawning toward June would for example, accentuate this difference.

The importance of avoidance and escapement


Figure 5.-The correlation coefficient between the annual total regional census estimates of sardine eggs and larvae by size class with the Murphy (1966) sardine biomass estimates. The first point is for total eggs, 1951-59; the second point is for total larvae, 1951-59; the third point is for total larvae with 1953 and 1959 censored. The succeeding points are for standard-size classes for all years 1951-59.

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Table 7.-A comparison of the regional census estimate (A), separately weighted nearshore (B), and 1 month per quarter sampling (C), in the southern California inshore region, 1951-60.

|  | Winter |  | Spring | Summer | Fall | Total | Annual rank |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1951 | A | 135 | 609 | 11 | 540 | 1,395 | 9 |
|  | B | 175 | 879 | 92 | 441 | 1,587 | 9 |
|  | C | 289 | 251 | 76 | 276 | 892 | 10 |
| 1952 | A | 87 | 261 | 314 | 134 | 796 | 10 |
|  | B | 70 | 229 | 336 | 119 | 754 | 10 |
|  | C | 52 | 315 | 531 | 122 | 1,020 | 9 |
| 1953 | A | 578 | 184 | 763 | 1,904 | 3.429 | 6 |
|  | B | 519 | 139 | 734 | 1,409 | 2,801 | 6 |
|  | C | 163 | 339 | 1,089 | 728 | 2,319 | 8 |
| 1954 | A | 3,123 | 516 | 519 | 237 | 4,395 | 3 |
|  | B | 1,006 | 445 | 412 | 180 | 2,043 | 7 |
|  | C | 2,577 | 332 | 934 | 149 | 3,992 | 5 |
| 1955 | A | 2,012 |  |  | 150 |  | 4 |
|  | B | 1,682 | $1,006$ | 737 | 117 | 3,542 | 4 |
|  | C | 1,941 | 930 | 1,111 | 55 | 4,037 | 4 |
| 1956 | A | 236 |  |  |  |  | 8 |
|  | B | 186 | 1,009 | 635 | 1.17 | 1,947 | 8 |
|  | C | 302 | 1,632 | 708 | 129 | 2,771 | 7 |
| 1957 | A |  |  |  |  |  |  |
|  | B | $3,650$ | $3,264$ | 933 | 238 | $8,085$ | I |
|  | C | 4,951* | 4,060 | 791 | 154 | 9,956 | 1 |
| 1958 | A | 2,316 | 2,912 | 403 | 63 | 5,694 | 2 |
|  | B | $2,186$ | $3,226$ | 417 | 36 | $5,865$ | $2$ |
|  | C | 1,753 | 4,246 | 554 | 51 | 6,604 | 2 |
| 1959 | A | 972 | 2,303 | 108 | 101 | 3,484 | 5 |
|  | B | 907 | 2,590 | $111$ | 72 | 3,640 | 3 |
|  | c | 413 | 5,354 | 218 | 19 | 6,004 | 3 |
| 1960 | A | 1,5/9 | 1,448 | 198 | 34 | 3,259 | 7 |
|  | 8 | 1,692 | 1,564 | 189 | 36 | 3,481 | 5 |
|  | C | 263 | 2,394 | 296 | 34 | 2,987 | 6 |
| Februc $\begin{aligned} & r_{s}= \\ & r_{s}= \end{aligned}$ |  |  |  |  |  |  |  |

to biomass estimation is solely a function of the variability these biases cause with respect to year-to-year differences between the capture and retention of sardine and anchovy larvae. For example, in Figure 5 the product moment regression coefficients, $r$ for sardine egg or larval census estimate versus sardine biomass are plotted by total eggs, total larvae (1951-59), total larvae with outlyer censored (1953-59), and each larval size interval through 15.75 mm . All the coefficients are high and positive with pronounced minima at the egg stage and at the 9.75 mm stage. One might ascribe the minimum associated with eggs to the effect of "patchy" distribution on precision of estimate. I have no ready explanation of the 9.75 mm minimum or the 12.75 mm maximum which follows it.

Sette and Ahlstrom (1948, p. 521) discussed the concept of "area of station" relative to sardine eggs. Trial calculations of the same kind suggest that with respect to sardine larvae, assigning equal areas to stations is also close to the more exact method of erecting perpendicular bisectors to each nearest station and using the area of the polygon so formed. For anchovy larvae, however, there may be an important problem. To study this problem, the southern California inshore region was divided into "nearshore", i.e., the standard station closest to shore on each line, and "nearshore-excluded" segments. In the 40 quarters of the years 1951-60, the mean concentration of eggs per positive station in the "nearshore" region exceeded that of the "near-shore-excluded" section by $35 \%$. Similarly the
proportion of positive stations was $87 \%$ "nearshore" and $68 \%$ in the "nearshore-excluded" region with respect to anchovy larvae.
In Table 7 I have compared the annual estimates of anchovy larvae using the quarterly regional census estimate of the entire region with the same estimate using the sum of the nearshore and nearshore-excluded segments of the region. The latter estimate is $8 \%$ lower than the regional census estimate, but it is not likely to be a bias since of the 10 annual estimates, the single estimate exceeds the partitioned estimate 5 times while the reverse is true an equal number of times. The Spearman rank coefficient of correlation is 0.85 . Also, a comparison of the annual estimates from 1951 to 1959 regional census estimates and those from a census estimate by Ahistrom (1967, Table 2) shows a Spearman rank correlation coefficient of 1.00 . Thus, we may conclude that errors of the kind involving nearshore gradients in the incidence and intensity of anchovy spawning and larval survival, while important for some applications, do not measurably affect the regional census estimates.

One may reasonably ask whether estimates generated from monthly cruises are comparable to larval abundance estimates from a single cruise in each quarter. In Table 7 I have compared the annual estimates of anchovy larvae using all regular occupied stations within the quarter and a similar estimate using only January, April, July, and October, with February used in 1957 because the January cruise was incomplete. The Spearman rank coefficient for the comparison is 0.90 : the mean value of the estimates from monthly cruises is $10 \%$ below the estimates from one cruise per quarter but in 5 years the monthly derived estimates exceed the quarterly and in 5 years the reverse is true. Thus, I conclude that differences which may arise from comparing estimates from quarterly and monthly cruises are not large enough to affect this study.
Technical errors should be relatively small and affect the catches of sardine and anchovy similarly. For example, the factor $a_{i}$, the area of mouth of the net is usually known to within $5 \%$, the factor $b_{i}$ is modified by the flow through the mouth of the net so that the length of tow is


Figure 6.-The size frequency of sardine larvae captured in 1951-60 in the southern California inshore (clear dot, solid line), offshore (black dot, small dash), and seaward (triangle, large dash) regions. The size frequency effect is attributed to offshore transport (see text).
underestimated by $13-15 \%$ : an error as great as $5-10 \%$ results from the flowmeter being centered in the backwash of the bridle apex hardware (Smith and Clutter, 1965; Tranter and Smith, 1968; Mahnken and Jossi, 1967).

Two biases, which have yet to be evaluated, are likely to be important. In cold water, the larvae may tend to grow more slowly. Thus the regional census estimate for a cold quarter or cold year could prolong the period for which the larvae are vulnerable to sampling. Similarly, larvae may persist without food for extended periods yielding the same kind of error mentioned for temperature. No estimate has been made for either bias for anchovy or sardine larvae.

All the sardine and anchovy larvae collected by the CalCOFI net have been subjected to trans-
port with the wind-driven layer of the ocean (Sette, 1943). While the entire problem of transport is beyond the scope of this paper, a comparison of size frequencies of larvae in the southern California inshore, offshore, and seaward regions shows a disproportionately lower number of younger larvae in samples from farther offshore (Figures 6 and 7). The most likely explanation of these data is that older larvae transported to the offshore regions are present in excess of the numbers that have been spawned and hatched there. Further, this implies that the larva size frequency slope in the spawning area is biased for the same reason. If the usual sampling grid encloses the spawning area and the areas to which larvae are transported no overall bias should ensue.

The MacGregor estimate of anchovy and sardine biomass cited above, mentioned specific numbers of spawnings per year and sex ratios. Neither of these has been suitably evaluated as yet. At present, there is no way of calculating the number of batches spawned per year. In this paper, I assume that the anchovy is twice as fecund as sardine. An attempt will be made to evaluate the sensitivity to multiple spawning by proposing three spawning models, below. The sex ratio is presently derived from the fishery which is conducted over a very small proportion of the anchovy range. Since males are somewhat smaller, they may be expected to be recruited to


Figure 7.-The size frequency of anchovy larvae captured in 1951-60 in the southern California inshore (clear dot, solid line), offshore (black dot, small dash), and seaward (triangle, large dash) regions. The size frequency effect is attributed to offshore transport (see text).
the fishery somewhat later than females. In this report, the simple assumption of equality of biomass will be maintained. This will tend to overestimate the adult biomass by the degree in which the male biomass is overestimated.

Table 8.-Sample size frequency distribution for sardine and anchovy. Total larvae per $10 \mathrm{~m}^{2}$ sea surface.

| Mid-range $x$ | Sardine I | $f x$ | $f x / \Sigma / x$ | Cumulative percent | Anchovy $f$ | $f x$ | $f x / \Sigma / x$ | Cumulative percent |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0.7 | 4 | 2.8 | 0.000 | 0.0 | 4 | 2.8 | 0.000 | 0.0 |
| 1.4 | 51 | 71.4 | 0.000 | 0.1 | 79 | 110.6 | 0.000 | 0.0 |
| 2.8 | 453 | 1,268.4 | 0.009 | 0.9 | 886 | 2,480.8 | 0.002 | 0.2 |
| 5.6 | 320 | 1,792.0 | 0.012 | 2.2 | 707 | 3,959.2 | 0.003 | 0.4 |
| 11.2 | 312 | 3,494.4 | 0.024 | 4.6 | 814 | 9,116.8 | 0.006 | 1.0 |
| 22.4 | 289 | 6,473.6 | 0.045 | 9.1 | 789 | 17,673.6 | 0.012 | 2.2 |
| 44.7 | 257 | 11,487.9 | 0.079 | 17.0 | 752 | 33,614.4 | 0.022 | 4.4 |
| 89.1 | 179 | 15,948.9 | 0.110 | 28.0 | 725 | 64,597.5 | 0.043 | 8.7 |
| 177.8 | 127 | 22,580.6 | 0.156 | 43.7 | 618 | $109,880.4$ | 0.072 | 15.9 |
| 354.8 | 78 | 27,674.4 | 0.191 | 62.8 | 579 | 205,429.2 | 0.135 | 29.4 |
| 707.9 | 44 | 31,147.6 | 0.216 | 84.4 | 369 | 261,215.1 | 0.172 | 46.6 |
| 1,412.5 | 14 | 19,775.0 | 0.137 | 98.0 | 247 | 348,887.5 | 0.230 | 69.6 |
| 2,818.4 | 1 | 2,818.4 | 0.019 | 100.0 | 94 | 264,929.6 | 0.174 | 87.1 |
| 5,623.4 |  |  |  |  | 23 | 129,338.2 | 0.085 | 95.6 |
| 11,220.2 |  |  |  |  | 4 | 44,880.8 | 0.030 | 98.5 |
| 22,387.2 |  |  |  |  | 1 | 22,387.2 | 0.015 | 100.0 |
| $\boldsymbol{N}$ | 2,129 |  |  |  | 6,891 |  |  |  |
| $\bar{X}$ | 67.88 |  |  |  | 226.95 |  |  |  |
| 2fx |  | 144,535.4 |  |  |  | 1,518,503.7 |  |  |

Table 9.-Regional census estimates of total anchovy larvae (0-: no sampling).

| Year | Quar. ter | Regions ${ }^{\text { }}$ |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | CCl | 0 CO | SCl | SCO | SCS | BCI | BCO | BCS | SBI | SBO | SBS | Total |
| 1940 | 1 | 0- | $0-$ | 771 | 196 | $0-$ | 0 | 0- | $0-$ | $0-$ | 0 | $0-$ | 967 |
|  | 2 | $0-$ | $0-$ | 258 | 214 | 0- | $0-$ | $0-$ | $0-$ | $0-$ | 0 | $0-$ | 472 |
|  | 3 | $0-$ | $0-$ | $0 \rightarrow$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 0 | $0-$ | $0-$ | $0-$ |
|  | 4 | $0 \rightarrow$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 0 | $0-$ | $0-$ | $0-$ |
| 19.41 | 1 | $0-$ | $0-$ | 184 | 351 | $0-$ | $0-$ | 0- | $0-$ | $0-$ | $0-$ | 0 | 535 |
|  | 2 | 0 | $0-$ | 422 | 579 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 1,001 |
|  | 3 | 0 | $0-$ | 27 | 8 | $0-$ | $0-$ | 0- | $0-$ | $0-$ | $0-$ | $0-$ | 35 |
|  | 4 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 0 | 0- | $0-$ | 0- |
| 1950 | 1 | 0 | 0 | 10 | 9 | 4 | 43 | 17 | 1 | 13 | 5 | 0 | 102 |
|  | 2 | 0 | 0 | 232 | 319 | 27 | 115 | 23 | 0 | 152 | 13 | 0 | 881 |
|  | 3 | 994 | 265 | $338$ | $5$ | 2 | $9$ | 1 | $0$ | 2 | 3 | 0 | $1,619$ |
|  | 4 | $0-$ | 0- | 0- | $0-$ | $0-$ | $0-$ | $0-$ | 0 | 0 | 0 | $0-$ | 0- |
| 1951 | 1 | 5 | 0 | 135 | 8 | 0 | 289 | 17 | 150 | 2,537 | 147 | 1 | 3,289 |
|  | 2 | 4 | 6 | 609 | 24 | 5 | 203 | 66 | 78 | 289 | 179 | 80 | 1,543 |
|  | 3 | 25 | 53 | 111 | 21 | 0 | 454 | 0 | 0 | 49 | 57 | 0 | 770 |
|  | 4 | 18 | 0 | 540 | 21 | 0 | 266 | 2 | 0 | 55 | 0 | 0 | 902 |
| 1952 | 1 | $0-$ | $0-$ | 87 | 33 | 0 | 1,372 | 111 | 0 | $1,762$ | 40 | $0$ | $3,405$ |
|  | 2 | 0 | 0 | 261 | 7 | 1 | 732 | 32 | 0 | 1,665 | $145$ | 0 | $2,843$ |
|  | 3 | 7 | 69 | 314 | 16 | 3 | 462 | 1 | 2 | 240 | 12 | 2 | 1,128 |
|  | 4 | 0 | 0 | 134 | 23 | 0 | 558 | 31 | 0 | 8 | 2 | 0 | 756 |
| 1953 | 1 | 0 | $0-$ | 578 | 67 | 6 | 1,849 | 247 | 2 | 4,160 |  | $227$ | 7.014 |
|  | 2 | 0 | 0 | 184 | 17 | 0 | 535 | 85 | 2 | 424 | 238 | $14$ | 1,499 |
|  | 3 | 2 | 0 | 763 | 38 | 7 | 373 | 2 | 0 | 150 | 33 | 0 | 1,368 |
|  | 4 | $0-$ | $0-$ | 1,904 | 882 | 0 | 831 | 48 | 0 | 84 | 2 | 0 | 3,751 |
| 1954 | 1 | 654 | $0-$ | 3,123 | 723 | 6 | 1,090 | 97 | 6 | 4,116 | 474 | 615 | 10,904 |
|  | 2 | 1 | 0 | 516 | 98 | 2 | 878 | 28 | 3 | 1,872 | 498 | 350 | 4,246 |
|  | 3 | 451 | 0 | 519 | 307 | 426 | 182 | 0 | 0 | 231 | 191 | 0 | 2,307 |
|  | 4 | 58 | $0-$ | 237 | 73 | 11 | 234 | 153 | 0 | 301 | 9 | 0 | 1,076 |
| 1955 | 1 | $0-$ | $0-$ | 2,012 | 283 | 6 | 7,152 | 499 | 11 | 750 | 312 | 2 | 11,027 |
|  | 2 | 2 | 0 | 1,074 | 246 | 6 | 540 | 266 | 296 | 787 | 380 | 24 | 3,621 |
|  | 3 | 7 | 0 | 849 | 184 | 35 | 257 | 3 | 0 | 602 | 9 | 27 | 1,973 |
|  | 4 | 5 | 10 | 1.50 | 19 | 0 | 210 | 15 | 0 | 59 | 7 | 4 | 479 |
| 1956 | 1 | $0-$ | $0-$ | 236 | 203 | 46 | 1,140 | 832 | 409 | 1,786 | $1,107$ | 2 | 5,761 |
|  | $2$ | 0 | 0 | $1,166$ | 63 | 14 | $2,239$ | 37 | 1 | $1,737$ | $154$ | 2 | $5,413$ |
|  | 3 | 247 | 0 | 708 | 562 | 21 | $1,413$ | 0 | 0 | 807 | 8 | 0 | $3,766$ |
|  | 4 | $0-$ | $0-$ | 156 | 115 | 4 | 0- | $0-$ | $0-$ | 0- | $0-$ | $0-$ | 275 |
| 1957 | $1$ | $0-$ | $0-$ | 3,314 | 1.085 | 969 | 2,720 | 102 | 3 | 2,683 | 69 | 0 | 10,945 |
|  | 2 | 20 | 21 | 3,601 | 814 | 69 | 853 | 104 | 8 | 426 | 44 | 13 | 5,973 |
|  | 3 | 5 | 20 | 791 | 592 | 64 | 380 | 1 | 0 | 358 | 234 | 17 | 2,462 |
|  | 4 | 64 | 0 | 267 | 229 | 4 | 86 | 0 | 0 | 10 | 0 | 0 | 680 |
| 1958 |  | 1,620 | $1,227$ | 2,316 | 2,407 | 268 | 4,680 | 618 | 16 | 4,606 | 788 | 5 | 18,551 |
|  | 2 | 252 | 15 | 2,912 | 1,163 | 497 | 1,099 | 384 | 2 | 165 | 57 | 44 | 6,590 |
|  | 3 | 413 | 387 | 403 | 1,508 | 84 | 143 | 5 | 0 | 15 | 0 | 0 | 2,958 |
|  | 4 | 11 | 5 | 63 | 34 | 0 | 54 | 2 | 0 | 4 | 0 | 0 | 173 |
| 1959 | 1 | 1,722 | 0 | 972 | 4,452 | 654 | 701 | 51 | 1 | 1,595 | 51 | 0 | 10,199 |
|  | 2 | 1,101 | 318 | 2,303 | 3,074 | 498 | 3,207 | 651 | 1 | 464 | 453 | 0 | 12,090 |
|  | 3 | 232 | 128 | 108 | 242 | 33 | 62 | 1 | 1 | 197 | 9 | 0 | $1,033$ |
|  | 4 | 5 | 0 | 101 | 2 | 0 | 9 | 4 | 0 | 20 | 0 | 0 | 1.41 |
| 1960 | 1 | 185 | 0 | 1,579 | 1,393 | 279 | 4,647 | 1,697 | 592 | 6,821 | 2,327 | 686 | 20,206 |
|  | 2 | 127 | 84 | 1.448 | 2,694 | 1,065 | 2,679 | 626 | 259 | 582 | 472 | 11 | 10,027 |
|  | 3 | 155 | 105 | 198 | 70 | 22 | 106 | 69 | 6 | 294 | 7 | 0 | $1,032$ |
|  | 4 | 1 | 0 | 34 | 9 | 0 | 67 | 0 | 0 | 21 | 2 | 15 | 149 |
| 1961 | 1 | 7 | 0 | 144 | 45 |  | $2,145$ | 143 | 0 | 13,436 | 234 | 0 | $16.154$ |
|  | 2 | 10 | 0 | 1,513 | 955 | 2,258 | $2,186$ | $516$ | $215$ | $3,068$ | $1,683$ | $115$ | $12,519$ |
|  | 3 | 677 | 310 | 359 | 1,127 | 1,070 | 114 | 1 | 0 | 27 | 9 | 0 | 3,694 |
|  | 4 | 35 | 15 | 18 | 25 | 4 | 32 | 3 | 2 | 30 | 7 | 0 | 171 |
| 1962 | 1 | 56 | 0 | 2,285 | 825 | 3 | 2,413 | 1,021 | 24 | 12,570 | 2,248 | 0 | 21.445 |
|  | 2 | 877 | 0 | 6,555 | 5,953 | 5,900 | 15,483 | 1,516 | 228 | 1,208 | 1,63 | 186 | 39,537 |
|  | 3 | $0-$ | $0-$ | 1,214 | 120 | 40 | 505 | 4 | 5 | 162 | 92 | 0 | 2,142 |
|  | 4 | 41 | 5 | 134 | 65 | 8 | 124 | 1 | 0 | 256 | 0 | 0 | 634 |

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Table 9.-Continued.

| Year | Quarter | Regions ${ }^{1}$ |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | CCl | CCO | SCI | SCO | SCS | BCI | BCO | BCS | SBI | SBO | SBS | Total |
| 1963 | 1 | 7 | 0 | 7,543 | 1,481 | 73 | 8,088 | 5,446 | 688 | 8,730 | 381 | 184 | 32,621 |
|  | 2 | 1,395 | 175 | 3.291 | 6,132 | 12,821 | 1,810 | 389 | 42 | 507 | 182 | 86 | 26,830 |
|  | 3 | $0-$ | $0-$ | 180 | 649 | 133 | 187 | 8 | 8 | 305 | 21 | 7 | 1,498 |
|  | 4 | $0-$ | $0-$ | 279 | 161 | 2 | 92 | 3 | 2 | 45 | 0 | 0 | 584 |
| 1964 | 1 | 1,420 | 0 | 4,137 | 3,552 | 7,213 | 2,895 | 133 | 66 | 6,678 | 148 | 5 | 26,247 |
|  | 2 | 3,056 | 82 | 4,524 | 6,223 | 3,656 | 1,063 | 1 | 0 | 384 | 73 | 77 | 19,139 |
|  | 3 | 538 | 125 | 353 | 1,814 | 1,516 | 1,058 | 26 | 15 | 204 | 6 | 0 | 5,655 |
|  | 4 | $0-$ | $0-$ | 945 | 110 | 3 | 71 | 0 | 2 | 81 | 0 | 0 | 1,212 |
| 1965 | 1 | 16 | 0 | 10,145 | 3,803 | 669 | 6,158 | 1,136 | 1.013 | 7,171 | 396 | 5 | 30,512 |
|  | 2 | 1,160 | 0 | 6,870 | 7,561 | 6,2,19 | 2,257 | 424 | 1,112 | 6,205 | 91 | 17 | 31,916 |
|  | 3 | 75 | 5 | 6,304 | 3,130 | 3,042 | 1,056 | 286 | 4 | 866 | 5 | 0 | 14,773 |
|  | 4 | $0-$ | $0-$ | 1,037 | 733 | 121 | 185 | 0 | 0 | 15 | 0 | 0 | 2,09.1 |
| 1966 | 1 | 2,973 | 0 | 7,078 | 4,285 | 409 | 4,426 | 275 | 9 | 3,289 | 189 | 15 | 22,948 |
|  | 2 | 181 | 0 | 7.823 | 9,902 | 1,164 | 1.065 | 63 | 5 | 1,023 | 2 | 0 | 21,228 |
|  | 3 | 738 | 56 | 2,427 | 960 | 214 | 881 | 36 | 2 | 454 | 90 | 0 | 5,858 |
|  | 4 | 51 | 2 | 1,022 | 62 | 14 | 181 | 4 | 0 | 83.0 | 0 | 0 | 2,186 |
| 1967 | 1 | $0-$ | $0-$ | O- | $0-$ | $0 \rightarrow$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | O- |
|  | 2 | $0-$ | $0-$ | 5,646 | 2,565 | 367 | 1,901 | 59 | 0 | 6.18 | 23 | 0 | 11.179 |
|  | 3 | $0-$ | 0 | $0-$ | 0 | $0-$ | 0- | 0 - | $0-$ | 0- | 0- | 0- | 0 |
|  | 4 | 0 | $0-$ | 0- | $0-$ | 0- | 966 | 150 | 8 | 224 | 605 | 0 | 1,958 |
| 1968 | 1 | 458 | $0$ | 3,318 | 3,259 | 406 | 2,342 |  | 0 | 0 | $0-$ | 0 | 11,634 |
|  | 2 | 4 | 0 | 4,020 | 1.194 | 697 | 960 | 397 | 0 | 303 | 361 | 0 | 7,936 |
|  | 3 | $0 \rightarrow$ | $0-$ | O- | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 0 | 0 | 0 | 0 |
|  | 4 | $0-$ | $0-$ | $0-$ | 0- | $0 \rightarrow$ | 0- | $0-$ | $0-$ | $0-$ | 0- | 0 | $0-$ |
| 1969 | 1 | 9.19 | 0 | 16,404 | 11,163 | 1,028 | $5,712$ | 1,671 | 5 | 4,198 | 588 | 38 | 41,726 |
|  | 2 | 3 | 0 | 5,982 | 2,364 | 294 | 3,459 | 350 | 196 | 612 | 237 | 3 | 13,500 |
|  | 3 | 28 | 91 | 3,101 | 678 | 92 | 282 | 13 | 19 | 178 | 13 | 0 | 4,495 |
| $\begin{array}{r} 1 \mathrm{CCl}= \\ \mathrm{CCO}= \\ \mathrm{SCl}= \\ \mathrm{SCO}= \\ \mathrm{SCS}= \\ \mathrm{BCl}= \end{array}$ |  | nia insho nia offsho rnia insh rnia offsh rnia seaw inshore |  |  |  |  | BCO <br> BCS <br> SBI <br> SBO <br> SBS | $\begin{aligned} & =\text { Baia } \\ & =\text { Baa } \\ & =\text { South } \\ & =\text { South } \\ & =\text { South } \end{aligned}$ | California <br> California <br> Baja insh <br> Baja off <br> Baja sea | offshore seaward ore hore ward |  |  |  |

The anchovy spawns over a small portion of its range apparently while still schooled; it also spawns over a small portion of the day ( $25 \%$ ); it now appears that most spawning takes place just after the full moon, ca. $20 \%$ of the lunar month (Smith).* In addition, the spawning behavior is highly seasonal with most spawning occurring in the first half-year. Biases result from sampling during a period in which spawning is occurring (Sette and Ahlstrom, 1948, p. 520 ), and similar errors may result from missing or oversampling spawning peak periods or dense small patches. Although patches would tend to disperse toward randomness, this process appears to be very slow relative to the duration of the larval stage. Patchiness persists and may

[^2]be augmented by predation. For this reason, the samples have been pooled to reduce variance wherever possible. Nevertheless, sample sizes range from zero to tens of thousands of larvae, and a pooled summary may still be based on a chance large sample. In all cases, the above error analysis was done with a sample frequency plot in view and all "outlyers" were examined for effect on the sampling question. All samples taken between 1951 and 1960 are listed by frequency distribution in Table 8 for sardine and anchovy total larvae.

Anchovy larvae abundance (Table 9) and sardine larvae abundance (Table 10) for 11 regions and the years 1940, 1941, 1949-69, are listed for each region and quarter in which sampling took place at standard stations. Figure 8 shows the comparison of the regional census estimates and standard haul summations from 1951 to 1966 (Ahlstrom, 1968). The close agree-

Table 10.-Regional census estimates of total sardine larvae (0-: no sampling).

| Year | Quarter | Regions ${ }^{1}$ |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | CCl | CCO | SCl | SCO | SCS | BCl | BCO | SCS | SBI | SBO | SBS | Total |
| 1940 | 1 | $0-$ | $0-$ | 81 | 343 | $0-$ | $0-$ | $0 \rightarrow$ | $0-$ | $0-$ | $0-$ | $0-$ | 424 |
|  | 2 | $0-$ | $0-$ | 141 | 210 | $0-$ | $0-$ | $0-$ | $0-$ | 0 | $0-$ | $0-$ | 351 |
|  | 3 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ |
|  | 4 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 0- |
| 1941 | 1 | $0-$ | 0 | 154 | 211 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 0 | $0-$ | 365 |
|  | 2 | $0-$ | $0-$ | 277 | 313 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 590 |
|  | 3 | $0-$ | $0-$ | 1 | 5 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 6 |
|  | 4 | $0 \rightarrow$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ |
| 1949 | 1 | 0 | 0 | 0 | 0 | 0 | 9 | 2 | 0 | $0-$ | 20 | 3 | 34 |
|  | 2 | 0 | 0 | 2 | 11 | 14 | 259 | 1 | 1 | 300 | 2 | 3 | 593 |
|  | 3 | 304 | 7 | 5 | 12 | 5 | 0 | 0 | 0 | 0 | 0 | 1 | 334 |
|  | 4 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 0 |
| 1950 | 1 | 0 | 0 | 0 | 0 | 0 | 92 | 1 | 0 | 117 | 128 | 2 | 340 |
|  | 2 | 0 | 2 | 116 | 206 | 128 | 60 | 91 | 3 | 1,951 | 402 | 0 | 2,959 |
|  | 3 | 9 | 4 | 3 | 2 | 2 | 1 | 2 | 0 | 10 | 11 | 0 | 44 |
|  | 4 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0 \rightarrow$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ |
| 1951 | 1 | 0 | 0 | 3 | 0 | 0 | 19 | 4 | 0 | 235 | 63 | 0 | 324 |
|  | 2 | 0 | 0 | 3 | 89 | 18 | 101 | 103 | 8 | 1313 | 107 | 17 | 1,759 |
|  | 3 | 0 | 0 | 4 | 1 | 0 | 270 | 0 | 0 | 94 | 7 | 0 | 376 |
|  | 4 | 0 | 0 | 0 | 0 | 0 | 146 | 0 | 0 | 79 | 1 | 0 | 228 |
| 1952 | 1 | $0 \rightarrow$ | $0-$ | 1 | 5 | 0 | 117 | 90 | 0 | 464 | 87 | 2 | 766 |
|  | 2 | 0 | 0 | 27 | 51 | 1 | 95 | 194 | 0 | 427 | 73.1 | 0 | 1,526 |
|  | 8 | 0 | 0 | 8 | 39 | 0 | 49 | 12 | 0 | 168 | 5 | 0 | 281 |
|  | 4 | 0 | 0 | 0 | 0 | 0 | 37 | 0 | 0 | 19 | 4 | 0 | 60 |
| 1953 | 1 | $0-$ | $0-$ | 0 | 0 | 0 | 46 | 199 | 0 | 342 | 94 | 40 | 723 |
|  | 2 | 0 | 0 | 1 | 0 | 0 | 73 | 509 | 0 | 123 | 370 | 4 | 1,080 |
|  | B | 0 | 0 | 1 | 0 | 0 | 86 | 0 | 0 | 100 | 22 | 0 | 209 |
|  | 4 | $0-$ | $0-$ | 0 | 0 | 0 | 62 | 0 | 0 | 113 | 2 | 0 | 177 |
| 1954 | 1 | 0 | $0-$ | 0 | 1 | 0 | 20 | 185 | 93 | 681 | 51 | 110 | 1,141 |
|  | 2 | 1 | 0 | 80 | 320 | 204 | 355 | 63 | 37 | 290 | 140 | 9 | 1,499 |
|  | 3 | 0 | 0 | 4 | 0 | 0 | 59 | 2 | 0 | 166 | 75 | 0 | 306 |
|  | 4 | 0 | $0-$ | 0 | 0 | 0 | 5 | 0 | 0 | 239 | 3 | 0 | 247 |
| 1955 | 1 | $0-$ | $0-$ | 1 | 2 | 8 | 179 | 278 | 285 | 142 | 110 | 0 | 1,005 |
|  | 2 | 0 | 0 | 35 | 146 | 85 | 72 | 99 | 70 | 14 | 8 | 0 | 529 |
|  | 3 | 0 | 0 | 1 | 31 | 34 | 228 | 4 | 0 | 34 | 9 | 0 | 341 |
|  | 4 | 0 | 0 | 0 | 0 | 0 | 18 | 0 | 0 | 64 | 2 | 0 | 84 |
| 1956 |  |  |  |  |  |  |  |  |  |  |  |  | 546 |
|  | $2$ | $0$ | $0$ | 2 | 173 | 48 | 52 | 11 | 14 | 70 | 93 | 0 | 463 |
|  | 3 | 0 | 0 | 2 | 43 | 9 | 307 | 0 | 0 | 335 | 0 | 0 | 696 |
|  | 4 | $0-$ | $0-$ | 1 | 0 | 0 | $0-$ | $0 \rightarrow$ | $0-$ | $0-$ | $0-$ | $0-$ | 1 |
| 1957 | 1 | $0-$ | 0 | 0 | 0 | 0 | 177 | 1 | 2 | 144 | 12 | 0 | 336 |
|  | 2 | 0 | 0 | 3 | 188 | 4 | 19 | 3 | 1 | 1 | 0 | 0 | 219 |
|  | 3 | 0 | 20 | 15 | 0 | 0 | 303 | 0 | 0 | 108 | 0 | 0 | 446 |
|  | 4 | 0 | 0 | 0 | 38 | 0 | 80 | 0 | 0 | 15 | 3 | 0 | 136 |
| 1958 | 1 | 0 | 0 | 83 | 52 | 0 | 259 | 0 | 0 | 224 | 21 | 0 | 639 |
|  | 2 | 3 | 0 | 52 | 27 | 4 | 9 | 2 | 0 | 12 | 1 | 67 | 177 |
|  | 3 | 0 | 0 | 26 | 94 | 2 | 287 | 1 | 0 | 217 | 0 | 0 | 627 |
|  | 4 | 0 | 0 | 0 | 0 | 0 | 9 | 0 | 0 | 1 | 0 | 0 | 10 |
| 1959 |  |  |  |  | 15 | 3 | 24 | 0 | 0 | 35 | 5 | 0 | 156 |
|  | 2 | 0 | 0 | 33 | 31 | 1 | 15 | 0 | 0 | 19 | 1 | 0 | 100 |
|  | B | 2 | 0 | 2 | 44 | 0 | 82 | 0 | 0 | 78 | 0 | 0 | 208 |
|  | 4 | 0 | 0 | 19 | 0 | 0 | 67 | 0 | 0 | 20 | 0 | 0 | 106 |
| 1960 | 1 | 0 | 0 | 33 | 13 | 0 | 6 | 0 | 0 | 207 | 0 | 0 | 259 |
|  | 2 | 0 | 0 | 40 | 7 | 0 | 2 | 0 | 0 | 6 | 0 | 0 | 55 |
|  | 8 | 0 | 0 | 13 | 0 | 0 | 228 | 81 | 0 | 121 | 2 | 0 | 445 |
|  | 4 | 0 | 0 | 1 | 0 | 0 | 138 | 0 | 5 | 72 | 0 | 0 | 216 |
| 1961 | 1 | 0 | 0 | 0 | 29 | 0 | 1 | 0 | 0 | 54 | 0 | 0 | 84 |
|  | 2 | 0 | 0 | 7 | 4 | 6 | 0 | 8 | 0 | 73 | 0 | 0 | 98 |
|  | 3 | 0 | 0 | 14 | 2 | 0 | 90 | 0 | 0 | 17 | 0 | 0 | 123 |
|  | 4 | 0 | 0 | 15 | 0 | 0 | 8 | 0 | 0 | 295 | 19 | 0 | 337 |

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Table 10.-Continued.

| Year | Quarter | Regions ${ }^{\text {a }}$ |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | CCl | CCO | SCl | Sco | SCS | BCl | BCO | BCS | SB! | SBO | SBS | Total |
| 1962 | 1 | 0 | 0 | 10 | 0 | 0 | 1 | 0 | 0 | 204 | 0 | 0 | 215 |
|  | 2 | 0 | 0 | 7 | 7 | 0 | 18 | 0 | 0 | 9 | 2 | 0 | 43 |
|  | 3 | $0 \rightarrow$ | $0-$ | 26 | 1 | 0 | 240 | 0 | 0 | 56 | 23 | 0 | 346 |
|  | 4 | 0 | 0 | 1 | 1 | 0 | 14 | 0 | 0 | 109 | 2 | 0 | 127 |
| 1963 | 1 | 0 | 0 | 50 | 0 | 0 | 62 | 0 | 0 | 3 | 0 | 0 | 115 |
|  | 2 | 3 | 0 | 1 | 0 | 0 | 0 | 3 | 0 | 10 | 0 | 0 | 17 |
|  | 3 | $0-$ | $0-$ | 1 | 0 | 3 | 54 | 0 | 0 | 49 | 0 | 0 | 107 |
|  | 4 | $0-$ | $0-$ | 7 | 5 | 0 | 96 | 1 | 0 | 31 | 0 | 0 | 140 |
| 1964 | 1 | 0 | 0 | 1 | 0 | 0 | 0 | 0 | 0 | 229 | 0 | 0 | 230 |
|  | 2 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 9 | 1 | 0 | 10 |
|  | 3 | 0 | 0 | 4 | 0 | 0 | 22 | 0 | 0 | 159 | 0 | 0 | 185 |
|  | 4 | $0-$ | $0-$ | 19 | 0 | 0 | 3 | 0 | 0 | 58 | 0 | 0 | 80 |
| 1965 | 1 | 0 | 0 | 2 | 0 | 0 | 5 | 0 | 0 | 7 | 0 | 0 | 14 |
|  | 2 | 1 | 15 | 0 | 0 | 0 | 3 | 0 | 0 | 0 | 0 | 0 | 19 |
|  | 3 | 0 | 0 | 7 | 0 | 0 | 4 | 0 | 0 | 429 | 1 | $0$ | 441 |
|  | 4 | $0-$ | $0-$ | 1 | 0 | 0 | 406 | 0 | 0 | 216 | 1 | $0-$ | 624 |
| 1966 | 1 | 3 | 0 | 1 | 0 | 0 | 19 | 0 | 0 | 49 | 0 | 0 | 72 |
|  | 2 | 0 | 0 | 10 | 0 | 0 | 1 | 0 | 0 | 2 | 0 | 0 | 13 |
|  | 3 | 0 | 0 | 1 | 0 | 0 | 50 | 4 | 0 | 262 | 1 | 0 | 318 |
|  | 4 | 10 | 0 | 1 | 0 | 0 | 238 | 4 | 0 | 87 | 2 | 0 | 332 |
| 1967 | 1 |  |  |  |  | $0 \rightarrow$ |  | 0 |  |  | $0-$ | $0-$ | 0- |
|  | 2 | $0-$ | $0-$ | $5$ | $0$ | $0$ | $105$ | 0 | $0$ | $1,111$ | $0$ | 0 | 1,221 |
|  | $3$ | $0-$ | $0-$ | $0-$ | $0-$ | 0 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 0- |
|  | 4 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ | 11 | 0 | 0 | 59 | 0 | 0 | 70 |
| 1968 | 1 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | $0-$ | $0-$ | $0-$ | 0 |
|  | 2 | 0 | 0 | 2 | 0 | 0 | 5 | 1 | 0 | 0 | 0 | 0 | 8 |
|  | 3 | $0 \rightarrow$ | $0-$ | 0 | $0-$ | $0 \rightarrow$ | 0 | 0 | $0-$ | $0-$ | $0-$ | $0-$ | $0-$ |
|  | 4 | $0-$ | $0-$ | $0-$ | $0-$ | $0 \rightarrow$ | 0 | 0 | $0-$ | $0-$ | 0 | $0-$ | $0-$ |
| 1969 | 1 | 1 | 0 | 1 | 1 | 0 | 5 | 1 | 0 | 0 | 0 | 0 | 9 |
|  | 2 | 0 | 0 | 3 | 0 | 0 | 10 | 3 | 0 | 30 | 0 | 0 | 46 |
|  | 3 | 10 | 0 | 1 | 0 | 0 | 48 | 0 | 0 | 25 | 3 | 0 | 77 |
| $\mathrm{BCl}=$ Baja California in$\mathrm{BCO}=$ Baia California o$\mathrm{BCS}=$ Baja California se$\mathrm{SBI}=$ South Baia inshorSBO $=$ South Baja offsho$\mathrm{SBS}=$ South Baia seawar |  |  |  |  |  |  |  |  |  |  |  |  |  |

ment between these values lends support to the idea that the method of data assembly causes little bias relative to size of the major fluctuations in anchovy larva abundance. Both the Spearman rank difference and the product-moment correlation coefficients are 0.99 .

MacGregor (1968) suggested that the problem of determining the number of batches of eggs spawned per year per female is the major source of imprecision and bias in the adult biomass estimates from egg census and fecundity data. He further suggested that the best strategy for egg census would be to conduct an intensive cruise over a brief period in which no female is likely to spawn more than once. For the purpose of this paper, a preliminary judgment as to the importance of multiple spawnings in the larva ra-
tio estimate of anchovy biomass may be made by comparing three models of spawning behavior. The first model is that the adult biomass is proportional to the regional census estimate. This model assumes that the product of the number of eggs spawned per ton of female, the number of spawnings per year, and the mortality rate of the larvae is stable. The second model is that each anchovy spawns once in the winter quarter. The third model is that each anchovy spawns in the single maximum quarter. Restated, the adult biomass is proportional to the 1) annual average regional census estimate of larval abundance, 2) winter quarter regional census estimate, or 3) annual maximum quarterly regional census estimate. The 1951-66 data for the Ahlstrom standard haul summation, the


Figure 8.-A 16-year comparison of the regional census estimate derived for this paper and the standard haul summation quarterly average (Ahlstrom, 1968).
average regional census estimate, the winter spawning, and seasonal maximum are found in Figure 9. It is clear that the same trend and magnitude of increase are seen from each model. Since there is no obvious difference, the annual average regional census estimate of larva abundance will be used for the current biomass estimates.

It may be seen from Table 9 that the anchovy larvae have increased in number, particularly in the southern California section. Figure 10 illustrates the subdivision of the regional census estimate annual totals by section of the coast from 1951 to 1966. Within the southern California section, the inshore region was the dominant producer of anchovy larvae for the first 7
years. In the next 3 years, the southern California offshore became equally important in numbers of anchovy larvae. Between 1962 and 1965, the seaward region ( $160-280$ miles off the coast) was a very important locality for anchovy larvae and this influence dropped radically to pre-1961 values in 1966 (Figure 11). This change and a smaller one inshore resulted in an appreciable lowering of the overall total in 1966 (Figure 10). The subpopulations of the northern anchovy have been assigned to coastal sections (Vrooman ${ }^{\text {; }}$ Vrooman and Smith, 1972),

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Figure 9.-A time-series comparison of the annual quarterly average standard haul summation (Ahlstrom, 1968 , double cross-hatch) and the annual quarterly average regional census estimate ("C", clear bar including line C if any).
and the major portion of increase of northern anchovy is ascribed to the "central" subpopulation which spawns off southern California.

The seasonal contribution of anchovy larva abundance from 1951 to 1966 is illustrated in Figure 12. From 1951 through 1959, wintercaught larvae dominated the total. Beginning in 1960 , spring larva abundance became increasingly important and, in 1962, substantially exceeded winter larva abundance. As mentioned above, the changing day-length in the spring quarter makes an appreciable difference for which the data have not yet been corrected. The spring-caught larvae are underestimated as a result. No correction has been applied for tem-perature- or food-specific growth rate changes, by season.

The increase of numbers of anchovy larvae shown in the foregoing regional census estimates could result from the traditional spawning areas being more completely covered with larvae, from new spawning areas being covered, or there being more larvae per unit area. There are instances of all three in the 1951-66 period. In particular, in the southern California section of the coast, where most of the increase took place, the primary effect has been the additional cov-
ering of traditional spawning areas. For example, in the southern California offshore region, April through June, 7 of 50 tows contained anchovy larvae in 1953, 25 of 48 tows contained larvae in 1955, and 51 of 53 tows had anchovy larvae in 1959 ( $14.0 \%, 52.1 \%$, and $96.2 \%$ respectively). If one fits a regression line to the increasing proportions of positive stations with respect to anchovy larvae off southern California the slopes in percent per year from 1952 to 1966 follow:

|  | Winter | Spring | Summer | Fall |
| :--- | :---: | :---: | :---: | :---: |
| Inshore (0-80 miles) | 1 | 2 | 1 | 1 |
| Offshore ( $80-160$ miles) | 4 | 6 | 4 | 1 |
| Seaward (160-280 miles) | 2 | 7 | 2 | 1 |

Also, there are more anchovy larvae per positive station than before. In the years 1962-66 there are 5-8 times as many larvae per positive station as in 1951 (Table 11). The number of larvae per positive station is not independent of the proportion of positive stations. One interpretation of samples of larvae is that they are drawn from relatively small ( 100 's of meters, Smith, in press) patches of larvae which are dispersing in such a way that they tend to overlap. For example, for randomly distributed


Figure 10.-A time-series comparison of the annual total regional census estimates of anchovy larvae as grouped by coastal section. The highest line (solid circle) is comparable to the quarterly average of "A" in Figure 9. The space between the top line and the line with open dots represents the contribution to the total by stations in the south Baja section. The space between the open dot line and the line connecting the solid squares represents the contribution of the Baja California section to the total. The space between the line connecting solid squares and the line connecting open squares represents the contribution of the southern California section. The space between the lines connecting open squares and the abscissa represents the contribution of the central California section to the total. For relative areas of the sections refer to Table 3 and Figure 2.
patches covering 0.1 of the area, the chance of encountering two patches is 0.01 ; with patches covering 0.9 of the area, the chance of encountering two patches simultaneously is roughly 0.8 . In the southern California inshore and offshore regions, the proportion of positive stations is related to the number of larvae per positive station in the following way:

$$
\begin{array}{cl}
\log _{10} N_{1}=3.20 P_{1}-0.218 & t=11.1737 \mathrm{df} \\
\log _{10} N_{2}=2.13 P_{2}+0.748 & t=9.9537 \mathrm{df} \\
\log _{10} N_{3}=2.32 P_{3}+0.573 & t=14.4076 \mathrm{df}
\end{array}
$$

where $N_{i}=$ number of larvae per $10 \mathrm{~m}^{2}$ per positive station in the $i$ th region, when $i=1=$ southern California inshore region, when $i=2=$ southern California offshore region, when $i=3$ the inshore and offshore regions are combined;

$$
\begin{aligned}
P_{i}= & \text { proportion of positive stations in } \\
& \text { winter and spring in offshore }(i=
\end{aligned}
$$ 2) or inshore areas ( $i=1$ ), or both $(i=3)$.

If the patchy model applies, this set of equations is good only for the CalCOFI standard tow, in these two regions. For example on theoretical grounds, if the standard tow were to sample under $20 \mathrm{~m}^{2}$ rather than the average $3 \mathrm{~m}^{2}$ as at present, one would expect the proportion of positive tows to rise and the number per positive tow to decrease for any given average of anchovy larvae per total unit area.

In addition to an increasing proportion of positive tows in traditional spawning areas and increasing numbers of larvae per positive tow, the southern California seaward region is an example of a new spawning area being invaded in


Figure 11.-A time-series comparison of the annual total regional census estimates of anchovy larvae as grouped by distance from shore within the southern California section. Lines connecting open triangles represent the southern California inshore, closed circles, the southern California offshore, and open circles, the southern California seaward. For relative areas of the regions refer to Table 3 and Figure 2.
the early sixties and becoming largely unused by the mid-1960's. The incidence of larvae has continued at less than $10 \%$ of the 1962 peak year in the southern California seaward region (Table 9 ).

## THE CURRENT ESTIMATE OF ANCHOVY SPAWNING BIOMASS

The practice adopted in this paper for estimating anchovy spawning biomass consists of 1) regression estimates of the relationship between sardine larval abundance and Murphy's estimate (1966) of sardine biomass; 2) the ratio of anchovy: sardine larva abundance until 1958; 3) the regression estimate of the relationship between anchovy larva abundance between 1951 and 1958 and the sardine-derived anchovy bio-
mass; and 4) the use of regression estimates to calculate anchovy and sardine biomass estimates outside the regression period. This differs from Murphy's (1966) method of using an anchovy: sardine ratio for each anchovy estimate. It also differs from Ahlstrom's (1968) method of tying each estimate to the biomass of sardine adults in 1958. Since regressions between two variables with time trends are influenced by the degree of temporal coherence of each variable, caution is warranted in their use. Confidence intervals may be calculated but their meaning is not clear due to the violation of the assumption of independence among the values used to calculate the regression. The regression estimates are used here as a simple shorthand method of reporting an apparent relationship.

The spawner biomass estimates of anchovy and sardine which result from this and previous


Figure 12.-A time-series comparison of the annual total regional census estimates of anchovy larvae as grouped by season of spawning. The highest line (solid circle) is comparable to that found in Figure 10 and to the height of "A" in Figure 9.
studies are found in Table 12. The primary objective of this study concerns the period of increase from 1951 to 1966 . Despite various technical deficiencies in the basic data, the years 1940, 1941, 1950, and 1969 are also estimated in the section to follow.

The first column of Table 12, Murphy's (1966) sardine spawner biomass estimates, is reproduced without change. If those estimates are changed by later studies, all other values in the table must be adjusted. The important features of Table 12 are 1) a regression estimate to derive sardine spawner biomass estimates from the regional census estimates of total sardine larvae; 2) a ratio estimate of anchovy spawner biomass which is half the product of the anchovy: sardine ratio and the sardine biomass; and 3 ) a regression estimate of anchovy spawner biomass derived from the regional census estimate of total anchovy larvae.

The equation for a least squares estimate of sardine spawner biomass and sardine total larvae derived for this study is:

$$
B_{s}=0.230 L_{s}-0.057
$$

where $B_{s}$ is the annual estimate of sardine spawner biomass in thousands of short tons and $L_{s}$ is the regional census estimate annual total of sardine larvae in numbers times $10^{12}$ in the years 1951-58, excluding 1953. The years 1953 and 1959 were eliminated as outlyers on a scatter diagram when it was found that the assumption of constant size distribution of larvae had been violated. The equation was later simplified by assuming a zero intercept to:

$$
B_{s}=0.206 L_{s}
$$

The anchovy spawner biomass estimates were derived from the ratio of anchovy: sardine larvae and the Murphy sardine biomass estimate in the following way:

$$
B_{a}=c\left(\frac{L_{a}}{L_{s}} B_{s}\right)
$$

where $B_{a}$ is the ratio estimate of anchovy spawner biomass, $L_{a}$ is the regional census estimate

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Table 11.-Anchovy larvae per positive station.

| Year | No. of <br> positive <br> stations | No. of <br> anchovy larvae | No. of <br> larvae per <br> positive station |
| :---: | :---: | :---: | :---: |
| 1951 | 344 | 26,951 | 78 |
| 1952 | 399 | 53,457 | 134 |
| 1953 | 539 | 85,178 | 158 |
| 1954 | 626 | 126,191 | 202 |
| 1955 | 558 | 134,017 | 240 |
| 1956 | 482 | 104,192 | 216 |
| 1957 | 518 | 135,966 | 262 |
| 1958 | 667 | 197,082 | 295 |
| 1959 | 729 | 182,176 | 250 |
| 1960 | 628 | 254,263 | 405 |
| 1961 | 420 | 88,925 | 258 |
| 1962 | 374 | 190,735 | 510 |
| 1963 | 370 | 172,611 | 466 |
| 1964 | 372 | 141,498 | 380 |
| 1965 | 392 | 234,850 | 599 |
| 1966 | 738 | 364,689 | 494 |

of anchovy total larvae, and $c$ is a constant representing the relative fecundity of sardine relative to anchovy, here assumed to be approximately 0.5. Lenarz (1972) suggests a constant of escapement of 1.1 to counteract the tendency for anchovy larvae to pass through the mesh openings of the standard silk net to a greater extent than sardine larvae. This experimentally derived estimate from paired tows which captured both species in standard and fine mesh
nets is considered too small to be appreciable on the scale of variability encountered in plankton tows. Murphy's (1966) estimate of the escapement factor, i.e., 2, is rejected.

The equation for a least squares estimate of the anchovy spawner biomass and anchovy total larvae is:

$$
B_{a}=0.094 L_{a}+0.072
$$

defined in the same way as the sardine case above. Similarly, the zero intercept was forced, giving:

$$
B_{a}=0.098 L_{a}
$$

which is the equation used for the regression estimate of anchovy spawner biomass.

The aberrant years 1953 and 1959 were recalculated assuming the slope and the Murphy sardine spawner biomass estimate are correct and that the sardine larva estimates are biased or imprecise. The recalculated estimates (parentheses, Table 12) appear to conform better to the trends of anchovy biomass and anchovysardine ratio.

The data from 1940, 1941, 1950, and 1969 have been manipulated to extend the biomass estimates. The 1940 and 1941 cruises were

Table 12.-Sardine and anchovy spawner biomass estimates by ratio and regression methods.

|  | Murphy sardine spawner ( $\times 10^{35}$ ) | Regression sardine spowner biomass ( $\times 10^{3}{ }^{3}$ ) | Sardine larval estimate $\left(\times 10^{12}\right)$ | Anchovy larval $\underset{\left(\times 10^{12}\right)}{\text { estimate }}$ | Anchovy sardine ratio | Ratio anchovy spowner ( $\times 10^{3 \mathrm{~T}}$ ) | Regression anchovy spawner biomass ( $\times 10^{3} \mathrm{~T}$ ) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1940 | 1,296 |  | 1,634* | 5,943* | 3.64 | 2,359 |  |
| 1941 | 2,001 |  | 2,476* | 7,104* | 2.87 | 2,871 |  |
| $\cdots$ | 716 |  | 3,343 | 2,602 | 0.78 | 279 |  |
| 1951 | 570 | 553 | 2,685 | 6,504 | 2.42 | 690 | 637 |
| 1952 | 554 | 542 | 2,633 | 8.132 | 3.09 | 856 | 797 |
| 1953 | 709 | 450 | 2,189 (3,442)** | 13,632 | 6.23 (3.96)** | 2,209 (1,404)** | 1,335 |
| 1954 | 668 | 658 | 3,193 | 18,533 | 5.80 | 1,937 | 1,816 |
| 1955 | 425 | 404 | 1,959 | 17,100 | 8.73 | 1,855 | 1,676 |
| 1956 | 293 | 351 | 1,706 | 15,215 | 8.92 | 1,307 | 1,491 |
| 1957 | 212 | 234 | 1.137 | 20,040 | 17.63 | 1,869 | 1.964 |
| 1958 | 281 | 299 | 1,453 | 28,272 | 19.46 | 2,875 | 2,771 |
| 1959 | 190 | 117 | 570 (922)** | 23,463 | 41.16 (25.45)** | 3,910 (2,418)** | 2,299 |
| 1960 |  | 201 | 975 | 31,414 | 32.22 |  | 3.079 |
| 1961 |  | 132 | 642 | 32,538 | 50.68 |  | 3,189 |
| 1962 |  | 151 | 731 | 63,758 | 87.22 |  | 6,248 |
| 1963 |  | 78 | 379 | 61.533 | 162.36 |  | 6,030 |
| 1964 |  | 104 | 505 | 52,253 | 1103.47 |  | 5,121 |
| 1965 |  | 226 | 1,098 | 79,292 | 72.21 |  | 7,771 |
| 1966 |  | 151 | 735 |  |  |  |  |
|  |  |  |  | 52,200 | 71.02 |  | 5,116 |
| 1969 |  | $27 \dagger$ | 132 $\ddagger$ | 33,623 $\dagger$ | $254.72 \dagger$ |  | 3,293 $\dagger$ |

* 1940, 1941-larval estimates seasonally adiusted.
+ 1969-larval counts $75 \%$ complete; adiusted for extra retention of small larvae.
conducted during the sardine spawning season, but excluded an important portion of the anchovy spawning season. Similarly, the cruises only sampled $20 \%$ of the area we now consider routinely surveyed. The ratio of sardine and anchovy larvae was used for 1950 and will not be discussed further. The 1969 samples systematically violated the assumption of stable size composition of larvae.

The cruises of 1940 and 1941 were conducted over the southern California inshore and offshore regions in the spring and summer. The total for both species in both regions and in both years was derived by simulation of analogous cruises within the 1951-60 survey period and the pro-duct-moment correlation coefficient is listed for each species, for each region, and year.

| Year | Species | Region Original |  | Seasonally adjusted | Correlation coefficient |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1940 | Anchovy | SCI | 1,545 | 3,229 | 0.794 |
|  |  | SCO | 820 | 2,714 | 0.945 |
|  |  | Total | 2,365 | 5,943 |  |
| 1940 | Sardine | SCI | 502 | 726 | 0.794 |
|  |  | SCO | 891 | 908 | 0.955 |
|  |  | Total | 1,393 | 1,634 |  |
| 1941 | Anchovy | SCI | 1,494 | 3,257 | 0.845 |
|  |  | SCO | 2,104 | 3,847 | 0.962 |
|  |  | Total | 3,598 | 7,104 |  |
| 1941 | Sardine | SCI | 987 | 1,305 | 0.807 |
|  |  | SCO | 1,161 | 1,171 | 0.981 |
|  |  | Total | 2,148 | 2,476 |  |

The ratio of this group of samples, within the same boundaries as the other regional census estimates used here, change from 1.70 to 3.64 in 1940 and change from 1.68 to 2.87 in 1941. Accordingly, these ratios are used in Table 12.

The 1969 data point for anchovy is modified in a crude attempt to adjust for the extra retention of anchovy larvae in a new sampling net (see Figure 4) which retains approximately $50 \%$ more larvae. The 1969 surveys were conducted with a $1-\mathrm{m}$ net with $50 \%$ more open area, more regular mesh apertures, and an average mesh width of 0.505 mm rather than the 0.55 mm aperture silk net (used and wet). The effect of additional mesh on reducing the extrusion effect of filtration pressure is discussed in Tranter and Smith (1968).


Figure 13.-A time-series comparison of sardine and anchovy biomass estimates from 1940 through 1969. The solid circle represents sardine biomass as calculated from the fishery by Murphy (1966) and extends from 1940 to 1959 . The solid triangle represents sardine biomass derived from a regression estimate of the relationship between the Murphy biomass estimate and the annual total regional census estimate of sardine larvae during the reference years and is extended from 1951 through 1969. The open circle represents the estimates of biomass derived from the ratio of anchovy larvae to sardine larvae and the Murphy sardine biomass estimate from 1940 to 1959. The open triangle represents anchovy biomass derived from a regression estimate of the relationship between the anchovy tonnage calculated from the anchovy:sardine larvae ratio, and the annual total regional census estimate of anchovy larvae and extends from 1951 through 1969. The open squares represent the Murphy estimate of anchovy spawning biomass by 3 -year averages (Murphy, 1966, Figure 17, p. 65). Dashed lines represent interpolations between nonadjacent years.

Figure 13 contains all the estimates resulting from this study. Murphy's sardine biomass estimates are plotted from 1940 to 1959 . The anchovy biomass estimates derived from the anchovy sardine ratio and the anchovy larva regional census in 1940, 1941, and 1950 through 1959. The regression estimates are reproduced from 1951 to 1959 for comparison with the Murphy sardine biomass and ratio-derived anchovy biomass. The regression estimates of both sardine and anchovy spawner biomass are extended through 1966 and a tentative estimate for 1969 is placed for comparison.

## DISCUSSION

Important changes in the size of the anchovy population have occurred in the California Current area over the past two decades. The increase of anchovy has coincided with the continuing decrease of sardine in the same area. All available evidence indicates that the anchovy and sardine populations declined between 1941 and 1951, and thereafter the anchovy underwent a sustained increase reaching a plateau of $5-8$ million metric tons between 1962 and 1966. This population size may be between 2 and 3 times the anchovy spawning population of 1940-41 and between 5 and 10 times the anchovy population in 1950-51. Changes of this magnitude in the absence of a fishery underscore the importance of natural fluctuations in the population size of a fish species. Fishery management of such a species must be responsive to changes of this magnitude and rapidity.

Murphy $(1966,1967)$ speculated on the effect of the anchovy population on the recovery of the sardine population. Neither the feeding of anchovy and sardine nor the population dynamics of the food organisms is well enough understood in the California Current. Since the size frequency curves of larvae are relatively invariable (Ahlstrom, 1965; Lenarz, in press) one would expect competition, if any, to occur in the juveniles and pre-recruits of either species.

One interesting fact may be the decline of sardine spawning in the spring and summer has coincided with an increase of spawning in spring
by the anchovy. In the estimates of spawner biomass of sardine and anchovy, no attention has been given the possibility that fecundity in numbers of eggs per batch and number of batches per unit time is plastic.

The numbers of anchovy and sardine larvae have been used here exclusively to describe changes in the adult biomass which spawned them. Since this process seems so effective, one might wonder what is required to refine techniques so that spawning, and larval and juvenile survival, may be used to predict the recruitment of year classes to the fishery. I believe the two major barriers are that 1) the size composition of the larvae of both species is so dependent on the sampling gear that mortality rates will remain too crude to project survival and 2) the effect of transport of larvae, particularly offshore to "unfavorable" areas, is neither well enough measured nor understood to effect predictive sampling systems.

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[^0]:    ${ }^{1}$ National Marine Fisheries Service, Southwest Fisheries Center, La Jolla, CA 92037.

[^1]:    A review of estimates and procedures was conducted for the California Marine Research Committee (MRC) in 1964 and the texts of the presentations are to be found there. Statements and calculations were neither edited nor formally derived. Quotations from the Appendix to these minutes will be regarded as informal and cited (MRC, date).

[^2]:    ${ }^{4}$ Smith, P. E. Lunar periodicity in the spawning of the northern anchovy (Engraulis mordax). Unpublished manuscript filed at National Marine Fisheries Service, Southwest Fisheries Center, P.O. Box 271, La Jolla, CA 92037.

[^3]:    ${ }^{5}$ Vrooman, A. M. Anchovy subpopulations. Unpublished manuscript filed at National Marine Fisheries Service, Southwest Fisheries Center, P.O. Box 271, La Jolla, CA 92037.

