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9. ABSTRACT

Mechanistic simulation models of ecological processes are becoming useful tools for the study and management of complex natural systems such as upwelling coastal regions. The goal of this cooperative project was to develop such a model, which would provide working experience with the methods and serve as a basis for future model development in Peru. This report discusses briefly the ecological assumptions and mathematical methods used in the model. The model currently runs satisfactoryily, although its simple nature precludes extensive comparison with detailed, observed data. A number of suggestions for future modifications are presented. In addition, tentative, short-term and long-term programs of economic research were drawn up. The short-term programs calls for a study of artisanal fishery in order to make recommendations for the development of the fishery. The long-term program still is not clear though probably will entail economic research required for the proper development and management of the exploitation of Peru's fishery resources.

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Ecosystems Modeling, Economic Considerations for a Peruvian Coastal Fishery

James N. Kremer and Jon G. Sutinen



International Center for Marine Resource Development

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Reports on a cooperative project off Ilo, Peru, between the Invernational Center for Marine Resource Development, University of Rhode Island, and the Division of Oceanography, Instituto del Mar, Callao, Peru. Support provided by the Agency for International Development. Additional copies can be obtained from the Marine Advisory Service, University of Rhode Island, Narragansett Bay Campus, Narragansett, Rhode Island 02882. <u>Kingston 1975</u>

Mechanistic simulation models of ecological processes are becoming useful tools for the study and management of complex natural systems such as upwelling coastal regions. The CUEA (Coastal Upwelling Ecosystem Analysis) Program in which Peruvian scientists are involved includes the development of such a simulation model as a major objective. Ine goal of the cooperative project between the Oceanography Division of Peru's Instituto del Mar and the Graduate School of Oceanography of the University of Rhode Island is to develop such a model, which may provide working experience with the methods and serve as a basis for future model development in Peru.

The short duration of this preliminary project meant that (1.) only a very simple model could be attempted, and (2.) it was desirable to draw heavily on the work of similar models. These included the comparate estuarine model of Narragansett Bay (Nixon and Nicola, 1975). Note that the upwelling model developed by John Walsh (1975). Note that of formulation and numerical methods for the ecological model were based on the Narragansett Bay model, while the ecological assumptions were in most cases very similar to those of Walsh. A physical circulation model also was developed and patterned after the general three-dimensional mixing scheme used by Walsh to simulate a simplified upwelling region.

In mechanistic modeling, the first step is to construct a simplified conceptual version of the system of interest, based on the much more detailed and elaborate understanding of the system. Thus, the extremely complex upwelling environment is characterized in reality by rapidly varying conditions, "patchiness" in chemical and biotic elements, and numerous inter- and intra-species interactions. Any of these may be included in a model if sufficient knowledge is available to allow hypothetical formulation of the basic interactions. Therefore, a certain amount of subjective judgment is required to specify what properties of the real system are to be treated in the model system. These decisions are usually based on the probable importance of the processes in the system as perceived by the researcher, although the personal interest of the investigator is, of course, another valid criterion. In addition, the decision is not final, and a simple preliminary model can and should be continuously modified--explaining areas proving to be of greater interest, and perhaps simplifying those where less detail seems needed. The present modeling effort is necessarily preliminary, and future work by Instituto del Mar personnel may mean that feedback from such theoretical analyses will affect practical and experimental research.

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The simplified conceptual model of the upwelling region is represented in figure 1. Radiation, temperature and physical circulation are external forces operating on the biological community. Phytoplankton, the primary producers, are of basic interest in this first version of the model, and a fair degree of physiological detail is represented in their compartment. But the response of the phytoplankton cannot be adequately represented apart from other biotic influences, such as grazing and excretion by herbivores. Thus these interactions are included in the model, albeit very crudely and simply. The logical path for future work to take is to expand the detail in these compartments and provide mechanistic formulations for such processes as ingestion, respiration, reproduction and growth.

The phytoplankton compartment of the present model includes mathematical formulations of growth and nutrient uptake as a function of temperature, ambient nutrient concentrations, and light. The approach taken here characterizes the complex, multi-factoral nature of the growth process by postulating a temperature-dependent maximum rate for otherwise optimum conditions. This maximum is then reduced by terms representing the extent to which nutrients (nitrogen, phosphorous and silica) and light are less than optimum. The predicted net rate of production is then the product of these three terms. The realized growth, or net community primary production, depends on the predicted growth rate as diminished by herbivore grazing. Nutrient uptake is commensurate with predicted growth, while herbivore excrction and physical upwelling supply nutrients to the system. The physical circulation is modeled in a simple way which is, however, fle.ible enough to allow various rates of transport and velocity gradients to be approximated in the model region.

The following report is a brief discussion of the ecological assumptions and the mathematical methods used in the model. Throughout this discussion, frequent reference is made to the model of John Walsh (1975). His section headings are used, and references to pages in his report (w_{D} -) are made to facilitate the comparison. All citations to the work of Walsh refer to this paper unless other-wise specified.

The model presently runs satisfactorily, although its simple nature precludes extensive comparison with detailed, observed data. A number of suggestions for future modifications are presented, and it is hoped that they will provide immediate direction for continued modeling efforts at the Instituto del Mar. Nevertheless, even this primitive model has begun to play a role in suggesting relevant questions, pointing out uncertainties in our present knowledge and assisting in the planning of future research.

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Figure 1.

FORMULATIONS

Physical Assumptions

<u>Circulation--general</u>. The physical circulation model "UPWELL" was written to be as similar as possible to the mixing model of Walsh. Basic differential equations including advective and turbulent mixing in three dimensions and biological changes are solved, using a simple forward finite-difference method. Two different velocity fields were chosen which are like Walsh's (Wp 209), with the slow upwelling simulated by velocities identical to his figure 7, and fast upwelling simulated by doubling these. While the general UFWELL model is for any three-dimensional region (10 x 10 x 10 grids) and any velocity field. biological applications were completed using a simple version with fewer computation cubes: a one-dimensional region, 100 kilometers offshore x 70 meters deep. Apparently Walsh used a spatial region of these same dimensions in his ecological model, rather than the more complete three-dimensional model.

<u>Diffusion</u>. The equations chosen by Walsh for advection (Wp 218) and for turbulent diffusion (Wp 218) were used in this model with the same parameter values for the constants. Some uncertainty exists about whether or not Walsh used both equations explicitly, or whether diffusion was ignored by assuming it was "implicitly" included in the advection equation.

<u>Boundary Conditions</u>. The assumption made by Walsh--that air and coast boundaries should use concentrations of zero--violates the assumption that no diffusion may occur through these boundaries. A different assumption was made in this model which assures no <u>gradient</u> of concentration at the boundaries. Therefore, all concentrations at the boundary are made equal to the value at first adjacent cube.

<u>Radient Energy</u>. A constant value of radiation of 300 langleys per day was used in all cases for the ecological model. The ability of the plants to acclimate the light of optimal photosynthesis makes the exact choice of the radiation less critical. Only the relative values of the light-to-optimum ratio is important in the model.

The incident light follows the common exponential decrease

$$I_{\tau} = I_{o} e^{-kz}$$

where I is radiation (ly/time) at surface (I₀) or at depth z (I_z).

The extinction coefficient (k) is calculated using the equation

$$k = k_0 + 0.2514 P + 0.5047 P^{0.66667}$$

where P = mg phytoplankton C/L. This is identical to the original equation of Riley (1956) for a ratio of phytoplankton C:Chl = 35. The same equation was used by Walsh but with P in units of nitrogen.

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A ratio equal to 35 was chosen based on a preliminary review of data in reports of *Anton Bruun* cruises, but should be verified or changed by more complete analysis by the staff at the Instituto.

The value of $k_{c} = 0.025$ (extinction coefficient of water due to nonchlorophyll materials also is based on *Antrn Bruun* data. A preliminary look at six samples by plotting observed k vs. Chl <u>a</u> suggested an intercept (Chl = 0) of about 0.025. This too should be verified or altered by future work.

Chemical Assumptions

Nutrient Regeneration. While the main source of nutrients is probably the rich upwelled water, the role of nutrients regenerated by zooplankton or anchovy excretion may also be important. In this model, as in Walsh's, excretion of ammonia, phosphorous and silicate by the herbivores takes place in the ratio N:Si:P of 11:0,5:1 for zooplankton and 5:0.22:1 for anchovy. Walsh assumed that excretion only occurs when the animals are feeding. While this may be an uncertain assumption, my results suggest that without it the contribution of nutrients becomes much more important than it should. It seems that the assumption is necessary, and the interpretation is useful, that it is an actempt to simulate the patchiness of the animals' distribution.

<u>Nutrient Uptake</u>. Within the PHYTO submodel, one nutrient is chosen as most limiting each hour of the simulation. The basis for this selection is different than Walsh's, since there appear to be difficulties with his method. Both methods have been tried in this model, and the simulation results are very interesting. Some valuable experiments are suggested by the disagreement of the methods, and perhaps in the future the Instituto staff will be able to complete the experiments thus providing some answers to the questions.

Walsh chooses to select the "most limiting" nutrient on the basis of a comparison of nutrient ratios in the water with the ratios required by the phytoplankton. This nutrient is then used in a kinetic calculation of growth (or uptake) using the familiar Michaelis-Menton equation:

Uptake = $U_{max} \cdot [N] / (K_s + [N])$ (Wp 212)

In this report's model, the kinetic calculation is made for all three nutrients, and the "most limiting" is chosen on the basis of this comparison; that is, the nutrient that is most difficult for the plant to take up from the water is selected as most limiting.

The two methods are very different primarily because the values of the half-saturation constants (K_s) have ratios that are very different from the requirements of the plants. The plants <u>require</u>

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nutrients in the ratio N:Si:P of 16:10:1, but the K constants used in the model have the ratio 6:3:1. Therefore it is possible--and in fact it frequently happens in the model simulations--that nitrogen or silica are in concentrations which are lower than the ratio in the phytoplankton, while the kinetic calculation indicates that phosphorous is much more difficult for the plant to take up, and therefore should probably be considered limiting.

The difference is a theoretical one. Walsh's is an assumption based on the standing-stock of nutrients. The alternative is a dynamic assumption, based on the physiological capabilities of the phytoplankton cells. However, the implications are important; Walsh predicts that nitrogen and/or silica are limiting while the kinetic calculations clearly show phosphorous to be limiting. Experiments that would be helpful to resolve this question include two.

1. Nutrient kinetic observations should be made to more accurately evaluate the value of the K constants that are appropriate for the species in the upwelling environment. For example, measurements of growth observed at different concentrations should show a hyperbola that agrees with the Michaelis-Menton theory.

2. Enrichment experiments should be conducted where natural populations are grown in freshly collected water to which different materials have been added. If nitrogen or silica is limiting, addition of these chemicals should stimulate growth, while the addition of phosphorous alone will not. Such experiments as these would add to a basic understanding of which controls are important in the growth of phytoplankton populations in upwelling areas.

<u>Inhibition</u>. In this model, the preference of phytoplankton for ammonia before nitrate is included in a very simple form. Rather than a gradual transition as in the equation used by Walsh (Wp 212) the phytoplankton use ammonia until there is no more. Any additional nitrogen is provided by the concentrations of nitrite and nitrate in the water.

Biological Assumptions

<u>Phytoplankton</u>. In the present preliminary form of the model, the rates of all processes are assumed to represent averages over 24 hours. No diel periodicity is now included, although the calculations are made on an hourly time-step (dt=one hour) for the circulation requirements. It would of course be desirable to include daily patterns for many processes, and this would be a good direction for future work on the model to take.

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The maximum possible growth of the phytoplankton (G_{max}) in the model is determined using the general equation of Eppley (1972). He has demonstrated that a strong upper limit appears to exist as a function of temperature. The equation has the familiar form of an exponential relation:

$$G_{max} = e^{(0.063 \text{ Temp}-0.16)}$$

The remaining parts of the FHYTO submodel estimate to what extent light and nutrients limit the ability of the plants to achieve this maximum growth. Two fractions are calculated which represent the limitations, LTLIM and MXLIM.

The limitation of less than optimum light levels (LTLIM) is based on a formulation by DiToro et al. (1971) which represents a double integration of the physiological response equation suggested by John Steele (1962): I

$$G = G_{\max} \frac{I_o}{I_{opt}} e^{\left(\frac{1}{2} - \frac{1}{1} \right)}$$

DiToro has shown that the integral of this equation, when considered over the total daylight period and the depth of the water (or mixed layer), is possible if the assumption is made that radiation is constant throughout the day. My work at the University of Rhode Island (Nixon and Kremer, in press) has suggested a correction factor of 0.85 should be applied when noonday inhibition at the surface is considered. This correction is used in this model only in the surface layers when the radiation exceeds the optimum for photosynthesis.

The exact value of I_0 , the optimum value for photosynthesis, is chosen to be 50 percent of the surface. While this choice is arbitrary, it is well supported by observations of Steele (1962) and others, and may be assumed to represent the acclimation of the plants to the changing seasonal light intensity. Walsh makes a similar acsumption, choosing a "half-saturation" constant for light of 10 percent of the surface value. Although Walsh includes diel variation, he ignores some aspects of the basic photosynthesis-light response which may affect production at the surface quite dramatically. In addition, he chooses to consider light and the nutrients together, so that only one of them is limiting growth at any depth and time. This model, however, considers a 24-hour average, and light effects are considered in addition to nutrient limitation. In the future, it would be desirable if diel variations could be added to the model, since the hourly time-step is already necessary for the circulation computations. In the phytoplankton formulation, this would mean that the original equation of Steele (for the instantaneous rate) should be used each hour and at every depth, with surface radiation following a curve from dawn to noon to dusk.

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The basis for the nutrient formulation has already been described, and only a brief mention of it is necessary here. Half-saturation constants (identical to those of Walsh) were chosen to be used in Michaelis-Menton equations. Three values are calculated, one for each of the nutrients, NLIM, PLIM and SILIM. The smallest of these numbers represents the most limiting nutrient, and is used in the growth prediction. A form of the equation is used which gives a unitless fraction:

$$NLIM = G/G_{max} = [N]/(K_{s}+[N])$$

MXLIM, the most limiting fraction, is simply the minimum of NLIM, PLIM, and SILIM.

The final predicted growth rate is:

 $GP = G_{max} \times LTLIM \times MXLIM$ (day ⁻¹)

The actual growth is determined in an exponential rate equation

 $P = P_{O,e}$ (GP-grazing) time biomass (mg C/L)

<u>Herbivores</u>. In this model, grazing and excretion by the herbivores are treated very simply. Rates for the two processes are chosen from the data presented by Walsh, and biomasses were selected, also airectly from Walsh. No diel vertical migration is considered for either the zooplankton or the fish. Instead, all rates are averaged over the total water column and over 24 hours.

Biomass Estimates--Zooplankton. Walsh suggests a biomass of 41 mg dw/m³ during the autumn characterized by relatively slow upwelling. During the winter when upwelling is faster, the biomass is more dispersed (12 mg dw/m³). Since the unit of volume in the biological model is liters, the estimates are converted:

Autumn: $41(\text{mg dw})/\text{m}^3 \ge 0.001 \text{ (m}^3/\text{L}) = 0.041 \text{ (mg dw/L})$ (24 hr. avg.)

Winter:
$$12 \times 0.001 = 0.012$$

Biomass Estimates--Anchovy. The same calculation m_y be made for the estimate of fish biomass used by Walsh:

Autumn: $(6 \text{ fish/m}^3) \ge 0.001 = (0.006 \text{ fish/L})$ (24 hr. avg.)

Winter: $0.04 \times 0.001 = 0.00004$

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<u>Grazing Stress</u>. Grazing Rate Estimates--Zooplankton. Walsh cites literature relating growth and assimilation to ingestion which allows the following budget:

Growth = assimilation - excretion (Wp 216)

 $0.268 I_{\pi} = 0.6 I_{\pi} - 0.03 (\mu g - at N)/(mg dw hr)$

where $I_z =$ ingestion of zooplankton. Solving this for ingestion:

 $I_{p} = 0.0903 (\mu g-at N)/(mg dw hr) ingested$

But the food that is ingested (phytoplankton) has the composition 106 carbon:16 nitrogen. Or

0.0795 mg C/µg-at N

Then the ingestion may be converted to carbon:

I₂ = 0.0903 x 0.0795 = 0.007 (mg Phyto. Carbon)/(mg dw Zoo hr)

ingested. This is the same as Walsh's rate of 0.02 hour⁻¹ (Wp 216)

Grazing Rate Estimates--Anchovy. The fish ingestion is estimated in the same way for two seasons:

Growth = assimilation - excretion

94.2 (μ g-at N)/(fish day) = 0.8 I_A - (486 μ g-at N)/(fish day) (Wp 215) I_A = 725.3 (μ g-at N)/(fish day) or 30.2 (μ g-at N)/(fish hr) (autumn biomass) I_A = 1180 or 49.2 (winter biomass)

Again, the ingested phytoplankton ratio is 106:16, and the carbon ration may be calculated:

 $I_A \approx 30.2 \times 0.0795 = 2.4$ (mg Phyto Carbon)/(fish hr) ingested (autumn) $I_A = 49.2 \times 0.0795 = 3.9$ (winter)

The rate of 2.4 is almost the same as Walsh's rate of 0.005 hour⁻¹ (Wp 215)

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Excretion Rate Estimates. Zooplankton. Walsh cites McCarthy (1971, see Wp 215) for the estimate of 0.03 ug-at NH₃/(mg dw hr). The ratio of nutrients excreted was chosen to be N:Si:P^{11:0.5:1} (Wp 211). Thus, the hourly excretion rates for zooplankton are:

Anchovy. The same estimates are made for the fish excretion using the ammonia excretion rate of 13.5 ug-at N/(g dw hr) and a ratio N:S1:P of 5:0.22:1 (Wp 215 and 211). To convert the estimates to ratesper-fish, a value for weight of each fish is required. Estimates of 1.5 grams (dry weight) per fish (autumn) and 2.5 grams (dry weight) per fish (winter) are based on the wet weight values of 6.5 and 10.5 given in Wp and the conversion of 25 percent of wet weight given in Wp 215. The excretion rates then are calculated:

$$EXCN_{A} = (\mu g - at N)/(g dw hr) \times 1.5 (g dw)/(fish) = 20. (\mu g - at N)/(fish hr)$$

 ${\rm EXCP}_{\rm A}$ and ${\rm EXCSi}_{\rm A}$ are calculated in the model similarly using the ratio of excreted nutrients and the appropriate biomass for the season.

<u>Grazing Threshold</u>. A more traditional grazing equation than that used by Walsh is used in this model to express food density dependence. Ivlev's equation (1945) has frequently been used for density dependence in grazing models:

Ration =
$$R_{max}$$
 (1. - $e^{-k(P-Po)}$)

or

XLIM =
$$R/R_{max} = 1. - e^{-k(P-Po)}$$

Walsh used the same rectangular hyperbola equation he uses for nutrients and light:

$$R/R_{max} = (P - Po)/[K_{s} + (P-Po)]$$

The two equations are similar at low food densities, but at the higher concentrations, Walsh's ration is significantly lower than Ivlev's. The distinction is arbitrary however, and either should be equally satisfactory (see Mullin et al. 1975).

Grazing thresholds are somewhat controversial (Frost 1975). While it seems reasonable that there should be such a threshold, for zooplankton at least, the experimental data do not always support the assumption. In simulation models, thresholds may avoid the mathematical artifact of over-

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grazing--when the predicted ration consumes more than the available phytoplankton, often forcing the concentration to go below zero. The mathematical method of solution used in this model, which will be discussed in more detail later, also avoids the artifact vroblem, so the threshold was not considered necessary. The additional interpretation given by Walsh is that the threshold simulates patchiness. Thus, when phytoplankton concentrations are below the threshold, he assumes that no herbivores will be in the region. This is perhaps more reasonable for the motile anchovy than the zooplankton, but it is nevertheless an interesting suggestion. Thresholds were used in the model, though not well established, so that their effect may be evaluated.

The calculation of excretion and grazing in this model results from the biomass estimate x (the rate/biomass):

excreted N = EXCN_z (Z)(XLIM_z) + EXCN_a (A)(XLIM_a)

 $NH_3 = NH_3 + excreted N - uptake$

where

Z, A = biomasses of zooplankton and anchovy

EXCN_{z,a} = the excretion rates of zoop] mkton and anchovy per unit biomass

XLIM, = the density-dependent food limitation terms (Ivlev)

Notice that this formulation assumes nutrient excretion only when grazing is taking place and in direct proportion to the rate of grazing. In my opinion, this is probably not a strong assumption since excretion certainly continues in a starving animal due to respiration, etc. The only basis for this assumption is the patchiness argument mentioned earlier concerning the possible feeding threshold. If the Ivlev equation is interpreted to mean that herbivores are only found in water where food is abundant, then the reduction of grazing is appropriate. This, however, is not the conventional interpretation. It is interesting to note that without this assumption, the excretion of nutrients is much too large, and the model gives less satisfactory results. The assumption is therefore included here, partly because Walsh used it, but more careful evaluation of the excretion formulation seems necessary before results may be interpreted in detail. Clearly, the uncertainty in these formulations is compounded by the uncertainty in the empirical data used in the rate estimates. Adequate formulations depend ultimately on increasing our data base in the area of herbivore excretion, as in others.

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Detritus. Because the role of the anchovy is only crudely represented here, detritus is not considered. Detritus in the form of zooplankton and anchovy fecal pellets (unassimilated ingested phytoplankton) is probably an additional food source for the herbivores, and may significantly contribute to the particulate nutrient levels in the water. For our purposes initially, no herbivore dynamic mechanisms are included in the model, and the role of detritus is ignored.

Computation Procedure

<u>Stability</u>. The grid scheme used in the circulation model UPWELL was selected to be the same as that of Walsh: 10 kilometers $x \ 10$ kilometers $x \ 10$ meters. For the velocity field used for both the autumn and winter system, numerical stability is not a problem for the same criteria mentioned by Walsh (Wp 217),

<u>Steady State</u>. The model is run to approximate a steady-state condition. Walsh mentions that for his model this is achieved in 1800 simulated hours (c 80 days), and my model also reached a stable condition in about 50 days. The criterion for stead state was a total change in phytoplarkton in all grids in the top two layers less than 0.1 percent.

Even though the model is used to represent only steady-state, it seems possible that time-variable simulations could be attempted too. In fact, the diel variation of Walsh seems to suggest that perhaps the interacting rates may track appropriate time-variations in a realistic way. This may prove a useful tool in investigating the spatial heterogeneity that may result from variations in the velocity fields along shore and offshore.

<u>Integration Method</u>. The solution of the differential equations of the model is accomplished using a method I developed for the Narragansett Bay (Rhode Island) model. The rates express basic growth processes, and thus suggest the use of exponential equations. Thus, instantaneous rates for growth and grazing may be evaluated simultaneously in the following form:

Let

GP = growth rate of Phyto (day⁻¹), i.e., mg C/(mg C·day)gz = grazing rate of herbivores (day⁻¹), i.e., L/(L·day) $P_o = initial condition of phytoplankton (mg C/L)$

Then

 $P = P_o e^{(GP-gz)t}$.

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This equation represents the <u>exact</u> integral of the differential equation for any time interval (t) during which the two rates (GP, gz) are constant. The differential equation is:

dp = (GP-gz)(P) dt

Conventionally, the differential equation is solved using a scheme of numerical integration--for each small time change, the value of dp is calculated and added to the present value of P for the next iteration. This is the method used by Walsh.

Thus

$$dp_{1} = (GP-gz)P_{0} dt$$
(1)

$$P_{1} = P_{0} + dr_{1}$$

$$dp_{2} = (GP-gz)P_{1} dt$$
(2)

$$P_{2} = P_{1} + dp_{2}$$

etc.

When the exact integral of simultaneous rates is used, the rates must all be of the same units. That is, both GP, the growth rate of the phytoplankton and the grazing rate of the herbivores, have only the units time⁻¹. The grazing of herbivores is originally calculated as a <u>ration</u> ingested (e.g., mg $C/_{hour}$). It is necessary to change this ration into a filtering rate of the correct units.

Ration =
$$I_z(Z)(XLIM_z) + I_a(A)(XLIM_a)$$
 (mg C/L·hr)
gz = [Ration (mg C/L·hr)]/[P (mg C/L)]

The filtering rate gz $(hour^{-1})$ represents the rate at which the combined grazing of the zooplankton and anchovy occurs. If the available food (P) is small, the herbivores will feed very fast, trying to get the desired ration. In some cases, gz may exceed 1.0--every liter of the water column is filtered more than one time. But because the rate is integrated in the exponential form, some phytoplankton will always remain. For example:

$$gz = 1.0 \text{ day}^{-1}$$
$$P_1 = P_0 e^{(-gz \cdot day^{-1})}$$

then

 $P_1 = 0.368 P_0$

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Thus, 36.8 percent of the original phytoplankton will remain after one day of intensive grazing, compared to 0.0 remaining if the conventional finite-difference integration method is used.

To correctly evaluate the uptake of nutrients and the actual amount of phytoplankton consumed by the herbivores, the exponential equation must be used again. The net result of one rate working in combination with other rates simultaneously may be specified.

If:

GP = growth rate

gz = filtering rate

Then:

$$PNET = -GP \times [P_0 (1 - e^{(GP - gz)})]/(GP - gz)$$

RTN = gz x [P_0 (1 - e^{(GP - gz)})]/(GP - gz)

Example. $GP = +0.698 \text{ day}^{-1}$ and $gz = 0.500 \text{ day}^{-1}$, growth is a little more than grazing.

After one day:

$$P = P_o e^{(0.693 - 0.500)} = 1.213 P_o$$

$$PNET = -0.693 \times P_o (1 - e^{0.193}) / (0.193) = 0.765 P_o$$

$$RTN = 0.500 \times P_o (1 - e^{0.193}) / (0.193) = -0.552 P_o$$

Final change = PNET + RTN = 0.765 $P_0 - 0.552 P_0 = 0.213 P_0$

$$P = P_0 + 0.213 P_0 = 1.213 P_0$$

In this example, the exact integral shows that the final change in the phytoplankton is $1.213 P_0$, and the other equations show that this change results from growth of 0.765 and grazing of 0.552. The uptake of nutrients must be computed from the PNET, since this is the total growth represented in the compartment, even though some is lost to grazing. Similarly, when calculations including the growth of the herbivores are added to the model, the actual ration consumed by them is the value RTN.

The method used in this model, in contrast to the finite difference schemes most frequently employed, evaluates the exact integral over " "dt" so that the only mathematical approximation is the assumption that the rates are constant over the time interval. For the simulations completed up to now, grazing (gz) is assumed to be constant at all times (a 24-hour average seasonally). The factors which determine

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GP are recalculated at short intervals. Nutrient limitation is usually the most rapidly changing condition, so that factor NUTLIM is computed every iteration in the simulation (hourly). Light limitation (LTLIM) only varies with changes in the extinction coefficient due to changes in phytoplankton biomass. These changes are relatively slow, and thus are computed only one time each day. The maximum growth rate as a function of temperature (G_{max}) depends only on the temperature field which is constant throughout the simulation. Thus this computation is done for every grid in the system only during the first iteration of the simulation.

During each iteration, the program computes the nutrient limitation terms (NLIM, PLIM, SiLIM) from which the most limiting nutrient term is chosen (MXLIM). GP is computed using the G_{max} and the daily value of LTLIM. The uptake of the nutrients is computed using the predicted growth of phytoplankton (PNET). The total changes of each of the nutrients and the phytoplankton for each of the grids are used in the UPWELL circulation program in combination with the diffusion and advection changes to compute the final change in each grid every hour.

RECOMMENDATIONS FOR CONTINUED RESEARCH

Model-Related Recommendations

To speed up the preliminary development of the ecological model, the three-dimensional circulation program (UPWELL) was modified to include only two dimensions. This form is well suited for many analyses because it is executed in a fraction of the time on the computer and is easier to deal with for the programmer. However, the questions of longshore gradients in horizontal and vertical velocities, which may be related to patchiness, cannot be evaluated in this simplified version. It would be desirable if a duplicate deck were prepared, on which the modifications were made to allow three-dimensional ecological simulations. The transition has been anticipated and only a few changes in dimension statements should be necessary.

Another question related to the circulation problem is the relative role of the advection and diffusion comporents of the mixing equation. In other words, what are the values of the eddy coefficients, especially in surface layers and regions of strong sheer associated with upwelling? The model UPWELL may be used alone to evaluate the effect of different values. For example, given only boundary values for nutrients, different eddy coefficients will result in different depth gradients which can be used to suggest the best choice of this parameter.

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Perhaps the most valuable improvement in the model which may be easily undertaken at this time is to include diurnal variations in some factors. For example, diel patterns of light may be represented by a simple sine equation, and the photosynthetic response of phytoplankton may then be evaluated, using the Steele equation, at each time and depth. The present assumptions calculate only a 24-hour average of the Steele equation integrated over each depth interval for the total day using the complicated equation of DiToro. Additional modifications appropriate here are vertical migration and grazing of herbivores.

Other Recommendations

The following are valuable experiments and field observations suggested by early work with the model.

The self-shading, extinction coefficient rela ionship, especially the appropriate value of k_0 , should be verified. A simple regression analysis of observed extinction coefficient (k) with Chl <u>a</u> (or plant carbon) will indicate both the intercept k_0 and an appropriate relationsnip.

The C:Chl ratio is used in the model to relate the state variable, carbon, to the more frequently measured biomass estimate, chlorophyll. Direct comparison of model results to field data requires as good an estimate as possible of this ratio and its range.

Similarly, the appropriate ranges of carbon:nutrient ratios for the phytoplaukton species are important values which need better estimation.

Nutrient kinetic experiments suggested by the model formulation of the most limiting nutrient would be very valuable. Additional surveys of appropriate half-saturation constants for growth (K_S) and enrichment experiments to evaluate nutrient limitation in the natural system would be important.

Grazing rates, excretion rates, feeding thresholds and biomass estimates of zooplankton as well as anchovy are essential before their role in the upwelling ecosystem may be even adequately approximated in the model. The formulation in the present model is certainly crude and serves primarily to point out areas where additional research would be extremely useful.

The potential role of switch-feeding of the anchovy between phytoplankton and zooplankton is another valuable line of research to pursue. The question of what factors may control this switching for juvenile and adult anchovy lends itself nicely to a combined experimental and modeling research effort.

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The potential role of detritus as a food source to zooplankton and anchovy also deserves future study. In many other marine and estuarine systems, detritus may be an important food or food supplement, and analysis of this possibility in the upwelling systems would be interesting.

THE ECOLOGICAL COMPUTER PROGRAM

The following is a short guide for the use of the physical circulation model UPWELL and the ecological oubroutines, ECOS and PHYTO. While a detailed, line-by-line description of the programs is not possible, this discussion of the critical variables and main input parameters is necessary for continued use of the model. The overall program-flow may be outlined as:

UPWELL

Initial specification, assignment of certain parameters; *READ velocities in y and z dimension for l plane (x=1); *READ temperature field; WRITE temp for reference; Convert input velocities (cm/sec) to program units (m/hr); Assign program control variables, and accumulators, etc.; Average pairs of input velocities to get center-cube flows; If CHECK = l, confirm continuity and grid-time stability; WRITE new velocity field - flows at center of grid cubes; *READ initial conditions for ecological variables, using Fortran NAMELIST input option. Begin iterations for ec_h time-step: Update timer and output-control variables;

Call ECOS: determine non-physical rates of change each hour;
Calculate boundary values using no-diffusion condition;
(At Air and Land, Cb=C so no concentration gradient exists at boundary b; at Deep and Offshore boundaries, constant initials kept);
Calculate advection and diffusion parts of change equation;
Integrate the physical and biological changes:
DCDT = Biol changes (R) - advection (DCZ, DCY) + diffusion (D2CZ, D2CY);
Transfer new concentrations into array VAL for ECOS and PHYTO accounting for uptake of NH4 first, then N02N03as required by assumption of inhibition of uptake by phytoplankton;
If IOUT=1 for output, WRITE arrays of state variables;
Subroutine STDYST checks for steady state of change if called.

End iterations for each time-step

Cal PROFIL subroutine to output final results



⁽Note additional output statements at end of program if desired to WRITE partial change values DCZ, D2CZ, etc.)

^{*}See sample data input example, figure 2.

SUBROUTINE ECOS (The derivation of the formulations are given in the official report; a brief outline of the program follows.)

Data assignments for grazing and excretion of herbivores, presently as DATA statements, my be changed to a READ; Call PHYTO;

Calculate excretion and grazing pressure by the zooplankton and anchovy (presently this is done only in the first iteration because constant rates are assumed--when a better formulation is used, this should be done every time);

Compute food-limitation term for herbivores; Compute excretion and grazing corrected for food limitation (this is done every iteration because phyto. is changing); Compute final rates of change (per hour) for all state variables; Return to UPWELL for integrations of rates.

SUBROUTINE PHYTO

Data assignments for important coefficients; Determine maximum potential growth rate for each grid based on the temperature, GMAX (this is only done once since temperature is constant); Determine degree of light limitation LTLLY (this is done once every day, since it depends on light which is assumed constant and the extinction coefficient which changes slowly due to selfshading by phytoplankton; Calculate kinetic nutrient limitation MXLIM based on the most limiting nutrient NLIM, PLIM, SiLIM; Assign correct symbol to array LSIGN to indicate the limiting nutrient for output. Calculate the estimate of net primary productivity GP as the product of GMAX x LTLIM x MXLIM; GP is the average rate of productivity (= C^{14}) during the 24-hour day, but it is an <u>hourly</u> rate, to agree with the time-step of UPWELL; Output useful variables for all grids at each output interval (IOUT = 1): Return to ECOS to complete rate-of-change calculations for the other state variables.

General Programming Considerations for UPWELL

Program used with ecological model was modified to only compute Y-Z plane, no X-direction (along shore) change is possible. This was done to save computer time and money. The basic model UPWELL is, however, three-dimensional, and conversion back to this form should be straightforward:

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a. Dimensions should be increased in UPWELL to allow a 10 x 10 x 10 array to replace the present $1 \times 10 \times 10$. Concentration and change arrays include extra boundary grids and also must be changed to $5 \times 12 \times 12 \times 12$ from present $5 \times 1 \times 12 \times 12$.

b. The X-direction changes must be replaced in the program, both in the DIMENSION statements (VX, DCX, D2CX) and in the body of the program where these are calculated. The original threedimensional version of UPWELL may be used for comparison, but <u>note</u> that the boundary conditions are different. The no-diffusion boundary conlition of the recent version is correct for the center-grid velocities.

The computational field of grids in the physical model is $1 \ge 10 \ge 10 \ge 10 \ge 10 \ge 10$ for three-dimensional) so velocities and rates arrays are dimensioned accordingly. Advection and diffusion calculations require an additional grid point at each boundary, so the array C is in dimensions of $3 \ge 12 \ge 12$. It is important in the program calculations to carefully keep track of which element of the C array agrees with elements of DCY, D2CZ, K, etc. For example, C(2,2,2) = R(1,1,1) for each substance that is mixed by the model.

The array VAL is used to transfer the concentrations of the state variables (PPL, NH_4 , etc.) between the subroutines and the main program UPWELL through the common block /VALUES/. VAL(500) is a one-dimension array because the many calculations in the subroutines ECOS and PHYTO may be completed independent of the physical location of the grid. (This is more efficient that the four-dimensional array C used in UPWELL because of the way the machine evaluates multi-dimensional subscripts every time they occur.) The concentrations of the state variables C are passed into array VAL for use in the subroutines, and any increase in the size of the grid field must have an increase in the dimension of VAL also.

Definitions of key variables in UPWELL include:

- VX, VY, VZ = velocities in the along shore, offshore, and down directions. Input as cm/sec across grid boundaries; converted in the program to center-grid averages as cubic meters per hour.
- D2CX, D2CY, D2CZ = partial change due to diffusion.
- DCX, DCY, DCZ = partial change due to advection.
- EDX, EDY, EDZ = eddy diffusion coefficients, assumed constant in x and y direction, variable with depth. Thus, EDZ(1) is coefficient for diffusion between Z_1 and Z_2 . A great deal of uncertainty about the correct values for EDZ with depth of zones of upwelling exists, and different values of this parameter should be tested in the model.

TEMP = temperature field assigned to grids. Constant with time.

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NX, NY, NZ = number of grids in each dimension.

- MXGRID = maximum number of grids = NX x NY x NZ. This number is used in sequential iterations of the concentrations stored in one-dimensional array VAL in ECOS and PHYTO.
- NC = number of state variables (concentrations) for UPWELL.
- BX, BY, BZ = array of size-limits including extra non-computational boundary grids. BX = NX + 2, etc.
- C = array containing the concentrations of the state variables in UPWELL, dimensioned C(NC, BX, BY, BZ).
- DX, DY, DZ = length in meters of the grids in the three-dimensions (presently 10km x 10km x 10m or 10000 x 10000 x 10m).
- DT = time-step of iterations, in hours (presently DT = 1 hr.).
- LAST = duration of the total simulation run, in hours. Steadystate seems to take 30-60 days, LAST=720 - 1440 hrs.
- CHECK = confirm continuity (conservation of water) and numerical stability of grid-size and time-step if CHECK = 1. This should be done once when new velocity fields are tried. Thereafter CHECK = 0 will avoid the checks. Note that the continuity check may indicate "water not conserved" for very insignificant volumes due to machine rounding error. This may be ignored after determining that the error is small.

General Programming Considerations for ECOS

The flow of this subroutine is not complicated. Calculations for all grids are done in the one-dimensional array of state variables P(MXGRID), NH4(MXGRID), etc. Rates of change DP, DNH4, etc., are calculated and returned to UPWELL by the common block /CHANGES/.

Note that all calculations for herbivore grazing and excretion are not done in every iteration, since average rates are not variable in time in the present model. This is one of the first changes that will be useful to make in the model.

Final rate-of-change estimates are based on the exponential evaluation of instantaneous growth and grazing. A discussion of this method is in the final report.

Definitions of key variables in ECOS:

- ANCH = biomass estimates of anchovy, no. of fish/L. Presently two values are specified, one for winter and one for autumn.
- ZOOP = biomass estimates of zooplankton, mg dry wt./L. two seasons.
- AEXCN, ZEXCN = rates of nitrogen excretion, ug-at N/L unit biomass. Two rates are specified for fish for different seasonal size.

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ARTN, ZRTN = rates of nitrogen excretion, ug-at N/L unit.biomass. Two rates are specified for fish for different seasonal size.

GRAZK = food-limitation coefficient for the Ivlev equation, L/mg C.

ZPO, APO = feeding threshold for no feeding of zooplankton and fish.

- SEASON = control of which season is selected for the herbivore biomass, etc. 1 = autumn, 2 = winter, 0 = no herbivores.
- ***Variables above presently must be specified for the simulation internally by Fortran DATA statements; in the future it might be desirable to read these values.
- XLIMA, XLIMZ = food-limitation fraction to adjust grazing pressure and excretion values (unitless fraction).
- RATION = total <u>preferred</u> ingested ration by herbivores; the final realized ration must be calculated from the instantaneous grazing rate FILT in the growth equations (mg C/hr).
- FILT = the predicted filtration rate of herbivores necessary to achieve the preferred ration based on available food.
- CHG = net phytoplankton rate of change = growth (GP) minus grazing (FILT).
- DP(grid) = predicted change in phytoplankton (mg C/hr) based on the exponential evaluation of the growth and filtration rates.
- DNN4(grid) = hourly change in ammonia, due to uptake by the phytoplankton (PNET) and e cretion by the herbivores.

DPO4(grid) and DSi(grid) = hourly change in phosphate and silicate.

General Programming Considerations for PHYTO

All calculations concerning the growth rate of phytoplankton are not done every iteration to save computer execution time. Since the temperature is constant for all computational grids, temperaturedependent maximum rates are only computed one time. Light limitation is calculated once every 24 hours, but nutrient limitation is determined every hourly time-step.

Although the final growth rate estimate (GP) represents an hourly instantaneous rate, it is based on the assumption of no diel variation; i.e., no day and night variations are represented during one day's 24-hour iterations. All conditions of nutrient availability change every hour, but the assumed light equation (LTLIM) corrects for the day-night effect.

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In this subroutine as well as ECOS, the one-dimensional arrays are used for the state variables. It is necessary to compute the level of light reaching each depth, however, so the actual location of each grid in the water column had to be determined. In the calculation of LTLIM, vertical columns of grids are computed in sequence from surface to bottom, with the light level continuously decreasing with depth through the series.

Definitions of Key Variables in PHYTO

- C\$NP, C\$PP, C\$SIP = carbon-to-nutrient ratios for the phytoplankton. Values for these important conversion factors are input to the program as ratios by atoms, C:N, C:P, C:Si. Because the units in the model are in milligrams of carbon, it is necessary to convert the ratios to µg-at/mg C in PHYTO.
- IOPT = the optimum light for photosynthesis. The actual value of this is not too important; only the ratio of the incident light to the optimum is used in the equations.
- RADN = radiation incident at the surface of the ocean. The units may be irradiance or insolation, but it is important that the choice of IOPT be based on the same units.
- F = photo-period, or day length, expressed as a fraction of a day.
- KO = extinction coefficient of water with no phytoplankton.
- KSN, KSP, KSSI = half-saturation constants for growth of phytoplankton for the three nutrients (Michaelis-Menton or Monod).
- DTT = conversion for the hourly rate of growth from the 24-hour value.
- GMAX(grid) = maximum growth rate of phytoplankton as a function of temperature (based on the work of Eppley, 1972).
- EXCOEF(grid) = estimated extinction coefficient due to absorption of light by the phytoplankton (converted from the equation of Riley, 1956).
- LTLIM(grid) = limitation to growth due to less than optimum light integrated over 24 hours for the depth interval of each grid.
- NTOT = total nitrogen available for phytoplankton growth, NH_L + NO₂ + NO₂.
- NLIM, PLIM, SILIM = kinetic factors for the limitation of growth due to nutrient concentrations (based on the hyperbolic Monod or Michaelis-Menton equation).
- MXLIM = the winimum of NLIM, PLIM, SILIM: the "most limiting" nutrient. (Notice that an alternative way to select the limiting nutrient is in the program as comments.)

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- $GP(grid) = \varepsilon$ stimated growth rate of the phytoplankton. The value is an instantaneous rate (hour⁻¹) representing net primary production, approximately the same as C^{14} data.
- OUTEX, OUTLT, OUTMX, OUTGP, LSOUT = arrays used for output of the part of the state variable arrays that are used in the calculations. (Now only 70 of the 100 grids are printed.)

Programming Considerations--Supplementary Subroutines

<u>BLOCK DATA</u>. This subroutine is necessary to assign initial values to variables and parameters placed in COMMON. It also would be possible to read these as inputs with Fortran READ statements.

<u>SUBROUTINE PROFIL</u>. This program is only to produce output of the simulated results in an easily readable format. Profiles of grid values for rates and concentrations offshore and with depth are produced for the following variables:

- CHL = estimated chlorophyll distribution assuming C:Chl = 35.
- PN = net phytoplankton primary productivity [mg C/ (m³.duy)]. This
 must be calculated from the growth rate and the biomass values.
- $AR = assimilation ratio, \mu g C/(\mu g Chl day).$
- DP = community production, i.e., net change of phytoplankton due to growth and grazing. This is the instantaneous hourly rate times a conversion factor to give a rate with units mg C mg C⁻¹ day⁻¹.

Profiles of the nutrient concentrations also may be easily obtained by adding additional WRITE statements.

SUBROUTINE STDYST. This program determines if the changes in the phytoplankton compartment are small enough to represent steadystate conditions. The criterion used is total change in the phytoplankton biomass in the top 20 grids (0-20m and 0-100km offshore) of less than 0.1 percent between successive calls of the subroutine STDYST. In the simulations up to now, STDYST is only called when output of the program variables occurs every five simulated days. Even with this rigorous test, the program converges to steady-state in 25 to 60 simulated days, depending on the exact conditions of the run.

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18. 0. <	17.	0.	0.	0.	0.	0.	0.	0.	0.	0.	ο.		
19.0.0.0.0.0.0.0.0.20.14.15.16.17.18.19.19.19.19.19.TC21.14.14.15.16.17.18.18.18.18.18.18.22.14.14.14.15.15.15.16.16.16.16.23.14.14.14.14.15.15.15.15.15.15.24.14.14.14.14.14.14.14.14.14.25.13.13.13.13.13.13.13.13.13.2728293031.& INITL VAL=3*.04,7*0.3*.04,7*0.3*.04,7*.01,3*.04,7*.01,3*.04,7*.01,3*.04,7*.01,33.100*0.5,34.10*0.5,10*10.,10*15.,10*20.,10*25.,50*33.,35.10*0.4,10*0.8,10*1.2,10*1.6,10*2.0,50*2.5,	18.	0.	Ο.	0.	ο.	0.	0.	0.	0.	0.	0.		
20. 14. 15. 16. 17. 18. 19. 19. 19. 19. 19. T C 21. 14. 14. 15. 16. 17. 18. 18. 18. 18. 18. 18. 22. 14. 14. 14. 15. 15. 15. 15. 16. 16. 16. 16. 23. 14. 14. 14. 14. 15. 15. 15. 15. 15. 15. 15. 24. 14. 14. 14. 14. 14. 14. 14. 14. 14. 1	19.	0.	0.	0.	0.	0.	0.	0.	0.	0.	0.		
21. 14. 14. 15. 16. 17. 18. 18. 18. 18. 18. 18. 22. 14. 14. 14. 15. 15. 15. 16. 16. 16. 16. 23. 14. 14. 14. 14. 15. 15. 15. 15. 15. 15. 15. 24. 14. 14. 14. 14. 14. 14. 14. 14. 14. 1	20.	14.	15.	16.	17.	18.	19.	19.	19.	19.	19.	Т	С
22. 14. 14. 14. 15. 15. 15. 16. 16. 16. 16. 23. 14. 14. 14. 14. 15. 15. 15. 15. 15. 15. 15. 24. 14. 14. 14. 14. 14. 14. 14. 14. 14. 1	21.	14.	14.	15.	16.	17.	18.	18.	18.	18.	18.		
23. 14. 14. 14. 14. 15. 15. 15. 15. 15. 15. 15. 24. 14. 14. 14. 14. 14. 14. 14. 14. 14. 1	22.	14.	14.	14.	15.	15.	15.	16.	16.	16.	16.		
24. 14. 14. 14. 14. 14. 14. 14. 14. 14. 1	23.	14.	14.	14.	14.	15.	15.	15.	15.	15.	15.		
25. 13. 13. 14. 14. 14. 14. 14. 14. 14. 14. 14. 14	24.	14.	14.	14.	14.	14.	14.	14.	14.	14.	14.		
26. 13. 13. 13. 13. 13. 13. 13. 13. 13. 13	25.	13.	13.	14.	14.	14.	14.	14.	14.	14.	14.		
27	26.	13.	13.	13.	13.	13.	13.	13.	13.	13.	13.		
28	27.	•	•	•	•	•	•	•	•	•	•		
<pre>29. 30. 31. &INITL VAL=3*.04,7*0.,3*.04,7*0.,3*.04,7*.01,3*.04,7*.01,3*.04,7*.01, 32. 3*.04,7*.01,3*.04,7*.01, 30*.01, 33. 100*0.5, 34. 10*0.5,10*10.,10*15.,10*20.,10*25.,50*33., 35. 10*0.4,10*0.8,10*1.2,10*1.6,10*2.0,50*2.5, 36. 10*3.0,10*4.0,10*7.0,10*10.,10*13.,50*25., &END</pre>	28.	•	•	•	•	•	•	•	•	•	•		
30. 31. &INITL VAL=3*.04,7*0.,3*.04,7*0.,3*.04,7*.01,3*.04,7*.01,3*.04,7*.01, 32. 3*.04,7*.01,3*.04,7*.01, 30*.01, 33. 100*0.5, 34. 10*0.5,10*10.,10*15.,10*20.,10*25.,50*33., 35. 10*0.4,10*0.8,10*1.2,10*1.6,10*2.0,50*2.5, 36. 10*3.0,10*4.0,10*7.0,10*10.,10*13.,50*25., &END	29.	•	•	•	•	•	•	•	•	•	•		
 \$\Lambda\$ INITL VAL=3*.04,7*0.,3*.04,7*0.,3*.04,7*.01,3*.04,7*.01,3*.04,7*.01, 3*.04,7*.01,3*.04,7*.01, 30*.01, 100*0.5, 10*0.5,10*10.,10*15.,10*20.,10*25.,50*33., 10*0.4,10*0.8,10*1.2,10*1.6,10*2.0,50*2.5, 10*3.0,10*4.0,10*7.0,10*10.,10*13.,50*25., 	30.	•	•	•	•	•	•	•	•	•	•		
32. 3*.04,7*.01,3*.04,7*.01, 30*.01, 33. 100*0.5, 34. 10*0.5,10*10.,10*15.,10*20.,10*25.,50*33., 35. 10*0.4,10*0.8,10*1.2,10*1.6,10*2.0,50*2.5, 36. 10*3.0,10*4.0,10*7.0,10*10.,10*13.,50*25., & END	31.	&INITL	VAL=3*	.04,7*0	.,3*.0)4,7*0.,	3*.04,7	*.01,3*	.04,7*.	.01,3*.0	04,7*.01,		
 33. 100*0.5, 34. 10*0.5,10*10.,10*15.,10*20.,10*25.,50*33., 35. 10*0.4,10*0.8,10*1.2,10*1.6,10*2.0,50*2.5, 36. 10*3.0,10*4.0,10*7.0,10*10.,10*13.,50*25., & END 	32.	3*.04	,7*.01,	3*.04,7	*.01,	30*.01	•						
34. 10*0.5,10*10.,10*15.,10*20.,10*25.,50*33., 35. 10*0.4,10*0.8,10*1.2,10*1.6,10*2.0,50*2,5, 36. 10*3.0,10*4.0,10*7.0,10*10.,10*13.,50*25., &END	33.	100*0.	5,										
35. 10*0.4,10*0.8,10*1.2,10*1.6,10*2.0,50*2.5, 36. 10*3.0,10*4.0,10*7.0,10*10.,10*13.,50*25., & END	34.	10*0.5	,10*10.	,10*15.	,10*20).,10*25	.,50*33	3.,					
36. 10*3.0,10*4.0,10*7.0,10*10.,10*13.,50*25., & END	35.	10*0.4	,10*0.8	,10*1.2	,10*1.	6,10*2.	0,50*2	5,					
	36.	10*3.0	,10*4.0	,10*7.0	,10*10).,10*13	.,50*25	j.,	&END				

Figure 2. Sample Data Input. Presently the ecological program reads the following data cards. Lines 1 through 10 read Y-velocities; 11-20, Z-velocities; 21-30, temperatures; 31 and 32, initial phytoplankton; 33, initial ammonia: 34, nitrate plus nitrite; 35, phosphate; 36, silicate. -24-

```
PROGRAM UPWELL AND ECOS AND PHYTO FOR PERU PROJECT APRIL 1975
C
     DECK MODIFIED FUR ONLY 1 Y-Z PLANE, NX = 1
ALL VX, DCX, D2CX OMITTED
DIMENSION VY( 1 , 10, 10
С
С
                                              VY( 1 , 10, 10), VZ( 1 , 10, 10),
                                                                                                     NO VX
                    E02(10).
       1
                   C( 5, 3, 12, 12), R(1 ,10 ,10 , 5), TEMP( 1 , 10, 10),
DCY( 5, 1 , 10, 10),
       ž
       3
                                                                                                     NO DOX
       NU D2CX
        CONMON /CNTRL/MXGRID, IOUT, NX, NY, NZ, DX, DY, DZ, DT, DAY, HOUR, TEMP
3 /VALUES/VAL(500) /CHANGE/ R
       3 /YALUES/VAL(500 )
        NAMELIST /INITL/ VAL
        DATA C/2160+0./
     DATA C//IDUFUS/
OIMENSIONS ARE ( C, X, Y, Z) -- INDICES ( N, I, J, K)
====>> INITIALIZE EDDY OIFFUSION PARAMETERS
HORIZONTAL PLANE EDX & EDY = 1.0E6 CM2/SEC
VERTICAL PLANE EDZ = 5.0 CM2/SEC ABOVE 20 M, 1.0E-3 BELOW
С
C
C
C
C
        VERTICAL PLANE EDZ = 5.0 CM2/SEC ABOVE 20 M, LOUE-3 DELLA
COEFFICIENTS MAY HAVE TO BE DIMENSIONED IN A NON UNIFORM FIELD...
M2/HR
        EOY=EDX
                                                                                                     M2/HR
        DATA EDZ/10* 0.36E-3/
EDZ(1)=1.8
        EDZ(2)=1.8
        READ IN VY AND VZ -- VX INITIALLY ALL 0.
READ(5,10)((VY(1,J,K),J=1,10),K=1,10),
C
C
                     ((VZ(1,J,K),J=1,10),K=1,10)
       1
        FURMAT(10F7.3)
10
     READ IN TEMPERATURE FIELD
С
        READ(5,10)((TEMP(1,J,K),J=1,10),K=1,10)
        I = I
        WRITE(6,910)ID(15),1,((TEMP(I,J,K),J=1,NY),K=1,NZ)
CUNVERT FROM CM/SEC INPUT TO PROGRAM UNITS M/HR; AND FILL ARRAYS
С
С
         (36 M/HR PER CM/SEC )
        DO 20 J=1+NY
DO 20 K=1+NZ
                                                                                                     M/HR
        VY/1,J,K}=VY{1,J,K}*36.0
       1 * 2.0
                                                                                                     RAPIDO
        VZ(1, J,K)=VZ(1, J,K)*36.0
                                                                                                     M/HR
                                                                                                     RAPIDO
       1 # 2.0
        CONTINUE
20
                                    PREPARE FOR COMPUTATION ITERATIONS
C
        essas)
        NC=5
```

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```

```
LAST=24
LAST=LAST*51
CHECK=1.
                                                                                                                                                                     HOUPS
              CHECK=0.
              IPG=0
              DX2=Dx+9X
             UY2=DY*DY
DZ2=DZ*DZ
              T=0.
DAY=1
C RESET COUNTERS FOR SPECIFIC ROUNDARY LIMITS FOR THIS PROBLEM
              NX=1
              NY=10
             NZ=7
EX=NX+2
             BY = NY + 2
              RZ=NZ+2
              8X1=8X-1
              BY1=BY-1
              BZ1≠BZ-1
С
              AXY=DX*DY
              AXZ=DX+DZ
              AYZ=DY+DZ
             VXMAX=0.
VYMAX=0.
              VZMAX=J.
              1=1
              (NDTE: IF MAX ARRAY SIZE IS INCHEASED, CHANGE '10' IN IF STMTS BELDW)
С
             DO 30 J=1+NY
J1=J+1
             IF(J1.GT.10)J1=10
PO 30 K=1,NZ
              K1=K+1
            K1=K+1

IF(K1.GT.10)K1=10

AVERAGE VELOCITIES AT GRID BOUNDARIES TO GET CENTER POINT VALUES

VY(1,J,K)=(VY(1,J,K)+VY(1,J1,K))/2.0

VZ(1,J,K)=(VZ(1,J,K)+VZ(1,J1,K))/2.0

IF(CHECK .NE. 1.) GO TO 29

CUNFIRM CONTINUITY (CONSERVATION OF WATER) FUR ALL GRIDS

DATA UMASSX/0./ .ARSVX/0./ VAMAX/0./

DMASSY=(VY(1,J,K) - VY(1,J1,K))*AXZ

DMASSY=(VY(1,J,K) - VZ(1,J,K))*AXZ

DMASS2=(VZ(1,J,K) - VZ(1,J,K1))*AXY

DMASS.NE.U.JWRITE(6,20)1,J;K,DMASSX,DMASSY,DMASSZ,DMASS

FORMAT(* WATEP NOT CONSERVED IN GPID (*,313,*) DMASS X,Y,Z=*,

1 3E15.4,1E20.0)
С
 С
 26
            1 3615.4.1620.8)
```

```
1 3E15.4, LE20.8)
C CHECK STABILITY CRITERIA
```

```
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```

```
495VY=A35(VY(I,J,K))
          ABSVZ=ABS(VZ(1,J,K))
IF(ABSVY.GT.VYMAX)VYMAX=ABSVY
          IF FAB SVZ.GT.VZMAXIVZMAX=ABSVZ
          CONTINUE
29
30
          CONTINUE
         I=1

WRITE(6,913)ID(7),I,(( VY(I,J,K),J=1,WY),K=1,NZ)

WRITE(6,910)IJ(3),I,(( V2(I,J,K),J=1,WY),K=1,NZ)

IF(CHECK,NE,L)GO TO 32

ADVFCTION CRITERIA:

DATA XSTAB,STABX/2+0./

YSTAB=VYMAX=DT/DY

JCTAU JUNATEDT/07
          1=1
C
          ZSTAH=VZ'AAX+DT/DZ
          TURBULENT DIFFUSION CRITERIA:
STABY=EDY +DT/DY2
С
          STABZ=EDZ(1)*97/072
          HRI FE (6, 35) XSTAR, YSTAB, ZSTAR, STADX, STABY, STABZ
        CONTINUE
FORMAT('DADVECTION STABLLITY CRITERIA X,Y,Z:',3E12.4,
I ' (MAXIMUM FOR EACH DIMENSION)'/
I ' TURBULENT STABLLITY CRITERIA X,Y,Z:',3E12.4]
DATA INSET,IDSET/5,0/
IF(INSET.ED.5) READ(5, NITL)
IF(INSET.ED.5) READ(INSET.915)DAY,VAL
HEITE(6.0(7)INSET.DAY
          CONTINUE
32
35
       WRITE (6,917) INSET, DAY
INPUT INITIAL CONDITIONS FUR PPL, NH4, NO2NO3, PO4, & SI
C
č
           ====> REGIN ITERATIONS
          07 1000 1T=1+LAST
           T=T+DT
          HOUR=AMOD(T+24+3)
      IF(HOUR-EO.J.)DAY=DAY+1
IOUT DETERMINES FREQUENCY OF OUTPUT
С
                                                                                                                            10 DAYS
          INUT=AMOD(T, 240.)
           CALL SUBROUTINES HERE TO DETERMINE NON-PHYSICAL KATES OF CHANGE IN C
С
          CALL ECOS
       TRANSFER VALUES INTO C-ARRAY FOR MIXING
С
           1=2
           IGRID=0
          DO 40 K=2,87
DO 40 J=2,871
IGRID=IGRID+1
           C(1+[+J+K)=VAL(IGRID )
C(2+1+J+K)=VAL(IGRID+100)
           C(3+1+J+K)=VAL(1GRID+200)
           C(4+1+J+K)=VAL(IGRID+300)
C(5+1+J+K)=VAL(IGRID+400)
```

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```

```
40 ·
C
                   CONTINUE
             DETERMINE ROUNDARY VALUES FOR CONCENTRATIONS
I.E. AT COAST AND SURFACE, NO DIFFUSION: C(B) = C(B-1)
LATERALLY AND OFFSHORE, C(B) = C(B-1) (or unchanging)
  C
C
C
                   DD 100 J=1+BY
DD 100 K=1+BZ
                    DU 100 N=1.NC
                  IF (J.EQ. 1) C(N, I, J, K) = C(N, I, 2, K)

IF (J.EQ. RY) C(N, I, J, K) = C(N, I, RYI, K)

IF (K.EQ. 1) C(N, I, J, K) = C(N, I, J, 2)

NO CHANGE IN DEEP BOUNDARY
                                                                                                                                                                                                                             COAST Y-
                                                                                                                                                                                                                             OF SHR Y+
SFC Z-
 C
                                                                                                                                                                                                                              BOTTOM
 100
                   CUNTINUE
                   SPECIAL CALCULATIONS NEEDED FOR NON-COMPUTATIONAL GRIDS WITHIN BOUNDARIES.
 С
                  I=2
DO 110 N=1,NC
([+2+8] IS A PHYSICAL BCUNDARY.
C(N,1,2+8)=C(N,1+2+9)
CONTINUE
C
110
                 CONTINUE
С
č
                  CALCULATE PARTIAL CHANGES IN X, Y, & Z DIRECTIONS DUE TO ADVECTION AND TURBULENT (EDDY) DIFFUSION
                  [=]
С
                  MATPIX C INDICES ARE ONE LARGER THAN LOOP INDEX; I => C(I-1)
                  11=1+1
                                                                                                                                                                                                                            =C(1)
                  12=1+2
                                                                                                                                                                                                                            =C(I+1)
                 DO 150 J=1.44
                  J1 = J+1
                  J2= J+2
                 00 150 K=1,NZ
                 K1=K+1
                 K2=K+2
                 4¥=1.
                 IF(VY(1, J, K). GT. 0. ) AY=0.
                 AY1=1.-AY
AY2=AY1-AY
                 AZ=1.
                 IF (VZ(1, J,K).GT.0.)AZ=0.
                AZ1=1.-AZ
AZ2=AZ1-AZ
                 00 150 N=1.AC

      9D
      150
      N=1,KC

      DCY(N,I,J,K)=
      VY(I,J,K)+(AY+C(N,I1,J2,K1)+AY2*C(N,I1+J1+K1)

      -
      AY1*C(N,I1,J,K2)+AZ2*C(N,I1+J1+K1)

      0CZ(N,I,J,K)=
      VZ(I,J,K)*(AZ*C(N,I1+J1+K2)+AZ2*C(N,I1+J1+K1)

      -
      AZ1*C(N,I1,J1,K1)+(AZ*C(N,I1+J1+K2)+AZ2*C(N,I1+J1+K1))

      -
      -
      AZ1*C(N,I1+J1,K1)+(AZ*C(N,I1+J1+K2)+AZ2*C(N,I1+J1+K1))

      1
      -
      -

      -
      AZ1*C(N,I1+J1,K1)+(AZ*C(N,I1+J1+K2)+AZ2*C(N,I1+J1+K1))
      -

      1
      -
      -
      AZ1*C(N,I1+J1+K1)

      1
      -
      -
      -
      AZ1*C(N,I1+J1+K1)

      1
      -
      -
      -
      -
      -
      -
      -

      1
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -
      -

             1
             1
                D2CYIN, I, J, K) = EDY
                                                                                      *(C(N, [1, J2, K1)+C(N, [1, J, K1)-TWOC)/DY2
```

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```
D2CZ(N,I,J,K)=EDZ(
Continu's
                                          K) + (C(N, I1, J1, K2) + C(N, I1, J1, K) - TWUC) / DZ2
150
C
C
          SUM PARTIAL CHANGES AND INTEGRATE
         F=1
11=1+1
         DO 200 J=1.NY
         J1=J+1
         DO 200 K=1.NZ
         K1=K+1
         DU 200 N=1+NC
DCDT=R(I+J+K+N)
                                                      \begin{array}{l} - DCY(N_{1}I_{1}J_{1}K) - DCZ(N_{1}I_{1}J_{1}K) \\ + D2CY(N_{1}I_{1}J_{1}K) + D2CZ(N_{1}I_{1}J_{1}K) \end{array}
                                                                                                            NO DCX
        1
                                                                                                            NO D2CX
         C(N, I1, J1, K1)=C(N, I1, J1, K1) + DT*DCDT
200 CONTINUE
      TRANSFER NEWLY VIXED CONCENTRATIONS BACK INTO ARRAY VAL
С
         IGRID=0
         I=2
        I=2

DD 160 K=2,11

DD 160 J=2,6Y1

IGRID=IGRID+1

VAL(IGRID )=C(1,I,J,K)

IF(VAL(IGRID )=C(4,I,J,K)

VAL(IGRID+300)=C(4,I,J,K)

VAL(IGRID+300)=C(4,I,J,K)
         VAL(IGRI0+400)=C(5,I,J,K)
С
    USE NH4 UNTIL DEPLETED, THEN USE NO2NO3:
         DEBT=C(2,1,J,K)
IF(DEBT.GE.O.)G0 TO 15A
DEBT=DEBT+C(3,1,J,K)
         IF(DEHT.LT.O.JDEBT=0.
VAL(IGRID+100)=J.
VAL(IGRID+200)=DERT
         C(2,I+J+K)=0.
C(3+I+J+K)=DEBT
GO TO 160
158
         VAL(IGRID+100)=DEBT
        VAL(IGRID+200)=C(3,1,J,K)
CONTINUE
160
ĉ
         IF(100T.NE.1)00 TO 1000
C
         WRITE Y-2 PLANE
         1=2
         DO 300 N=1,NC
         1PG=1PG+1
         IF(MOD(IPG,5).EQ.1)WRITE(6,900)
         WRITE(6,913)DAY, HOUR. ID(N), I, (( C(N, I, J; K), J=2, BY1), K=2, BZ1)
        CONTINUE
300
C
     CHECK TO SEE IF STEADY STATE HAS BEEN ACHIEVED
```

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```

.

	CALL STUYST
1000	CONTINUE
	IF(IOSET.NE.0)WRITE(IUSET.915)DAY.VAL
	LF(105FT.NE.0)WRITE(6.916) 105FT
	CALL PROFIL
	STOP
С	WRITE(6+910)10(4)+((DCX(1+1+1+K)+1=1+NY1+k=1+N7)
С	WRITE (6, 910) $[D(5), 1, (1, DCY(1, 1,, k, 1,, k,, k,$
С	WRITE (6,910) IU(6) + I + (1,0) CZ (1 + I + J + K + J + I + N) + K + N + NZ
С	WRITE $(6, 910) 10(7) \cdot (102Cx(1, 1, 1, 4, K)) = 1 \cdot Ny 1 \cdot K = 1 \cdot Ny 1$
С	WRITE $\{0, y_1, 0\}$ ID $\{0\}$, $\{1, 0, 2, C, y_1\}$, $1 = 1 = 1 = 1 = 1, y_1 = 1 = 1, y_2 = 1, y_1 = 1, y_2 = 1, y_2 = 1, y_1 = 1, y_2 = 1, y_2 = 1, y_1 = 1, y_2 = 1, y_1 = 1, y_2 = 1,$
С	WRITE (6+510) 10(9) + (102C7(1+1+1+1+K)+1+1+NY)+K=1+NY)
900	FORMAT(191)
910	FORMAT(101,5X,A4,1 X-GRID1,14//(1 1,10X,10F10 3,1)
C	WKITE(6,912)DAY+HUK+ID(N)+I+((C(N+1+J+K)+J=1+BY)+K=1+B7)
912	FORMAT('ODAY', 15.' HOUR', EG. 2.' CONCENTRATION 1. AG. Y COND. 16
	1/(* * 12F10-3))
913	FORMAT('UVAY', 15, ' HOUR', F6.2. CUNCENTRATION LAGELY CONDUCT
	1/(**,10F10,3))
915	FORMAT(15/(10F8.5))
916	FURMAT() ===> SUIPUT INITIAL CONDITIONS TO SCIE. 1621
917	FORMATING INPUT DATA READ FROM SETUIDS TO SECURITY START-UP ON DA
	1Y', (5)
	END

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```
      BLUCK DAIA

      COMMON /CNTRL/MXGP[D,10LT,NX,NY,NZ,DX,DY,DZ,DT,UAY,HDUR,T(100)]

      1 /PHYT/GMAX(100),LTLIN(100),MXLIM(100),LSIGN(100),GP(100)]

      2 /CNVRT/ C$NP,C$SPP.C$SIP

      1 /CHANGF/R(500)

      INTEGER DAY

      REAL GMAX,LTLIM,MXLIM,GP

      0ATA GMAX,LTLIM,MXLIM,GP

      0ATA C$NP,C$SIP,C$PP(16., 10., 1)

      C:N:SI:P GY ATCMS

      DATA C$NP,C$SIP,C$PP(16., 10., 1./

      0ATA

      MXGRID,NX,NY,NZ,DX,DY,DZ,DT,DAY,HDUR

      1 / 100, 3,10,10,2*1.E4.10.,1.0,0.

      ATA R/500 *0./

      .LSIGN/ 100***/
```

С

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```

```
SUBROUTINE FCOS
С
Ĉ
          HERBIVORF GRAZING AND EXCRETION INCLUDED HERE:
           AIDMASS ESTIMATES FOR HERBIVORES: FISHAL, MG DW ZOO/L
REAL AMCH(2)/.003, .00002/, ZUNP(2)/.02, .006/
PATES OF NITROGEN EXCRETION PER BIOMASS UNIT:
Ĉ
                                                                                                                                    PER L
С
          I , AEXCN(2)/20., 34. /, ZEXCN /.03/
INGESTED RATIUN PER BIGMASS UNIT:
                                                                                                                                    UGATN/L
С
           INCESTED RATION MER BLIMASS UNIT:

, ARTN[2]/2.4 , 3.9/, ZRTN /.007/ MG C/HR

GRAZK IS THE EXPONENT IN THE IVLEY GRAZING EQUATION: 1 - EXP(K*PHYTO)

3 , GRAZK/ -5.776/ L/MG C

"SEASON" SPECIFIES AUTUMN (1), WINTER (2), GR ZERO (0) BIOMASSES AND RATES
          ,
C
          3
С
           INTEGER SEASUN /2/
DATA 11/0/
С
           REAL LILIM, MALIM
           INTEGER LAB(1,6)/'PRDN', 'MGC/','L/DA','Y ','
                                                                                                    ...
                                                                                                                11
           COMMON /CNTPL/MXGRID, IOUT, MX, NY, NZ, DX, DY, DZ, DT, DAY-HOUR, TEMP(100 )
         1 /PHY1/GMAX(100 ),LTLIM(100 ),MXLIM(100 ),LSIGN(200 ),GF(100 )
2 /CNVKT/ C1NP,C1PP,C1SIP

        2
        //dlub/
        NH4(100 ), NU2N03(100 ), P04(100 ), SI(100)

        3
        //dlub/
        NH4(100 ), NU2N03(100 ), P04(100 ), SI(100)

        4
        /CHANGE/DP(100 ), DNH4(100 ), DNDX(100 ), DP04(100 ), DSI(100 )

с
с
           MXGRID=NX*NY*NZ
                                                               FRUM MAIN
           CALL PHYTO
           IF(11.NE.0)G0 TO 15
           [1=1
           TELSEASON.GT.0100 TO 10
           SFASON=1
           2009(1)=0.
           ANCH(1)=0.
          ANCH(1)=0.

ZFXCN = ZEXCN * ZOUP(SEASGN)

ZRTN = ZRTN * ZOUP(SEASGN)

ANEXCN=4EXCN(SEASUN) * ANCH(SEASON)

ANRTN = ARTN (SEASON) * ANCH(SEASON)

FXCRETION OF OTHER NUTRIENTS ACCORDING TO RATIO VS. N

FXCRETION OF OTHER NUTRIENTS ACCORDING TO RATIO VS. N
10
                                                                                                                                    UGAT/L/H
                                                                                                                                    MGC/L/HR
С
С
           ZUPPL EXCN= 11:0.5:1 AND ANCHVY EXCN= 5:0.22:1
ZFXCP=ZEXCN/11.0
                                                                                                              (N:SI:P)
                                                                                                                                    N:P=11
           ZEXCSI=ZEXCP+0.5
                                                                                                                                    SI:P=0.5
           ANEXCP=ANEXCN/5.0
                                                                                                                                    N:P=5.0
           ANEXS1=ANEXCP+0.22
                                                                                                                                   SI:P=.22
15
           CONTINUE
          COMPUTE RATES OF CHANGE PER HOUR FOR STATE VARIABLES
D(1 20 GRID=1,MXGRID
IF(P(GRID).GT.0.)G) TO 16
С
```

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	XLIMZ=0.	
	XLIMA=O.	
	RATION=0.	
	i IL T=0.	
14	9P(GR(9)=0.	
	PNET=0.	
	GN TO LA	
16	XLIMZ=1.0-EXP(GRAZK*(P(GR10)-ZP0))	
Ċ	FEEDING THRESHOLDS FUP ZUDPL (ZPU) A. J ANCHUVY (APU)	
	DATA 200.AP0/0.04. 0.24/	
	IF(XLIMZ,LT.O.O) XLIMZ=0.	
	XLIMA=1.0-EXP(GRAZK*(P(GRID)-APO))	
	IF(XLIMA.LT.O.O) XLIMA=0.	
	RATION=ZRTN+XLIMZ + ANRTN+XLIMA	
	FILT=RATION/P(GR1D)	1/HR
	CHG=GP(GRID)-FILT	
	IF (CHG. EQ. 0.) GO TO 14	
	TERM=EXP(C(1G)+1.0)	
	OP(GQ1D) = P(GR1D) * TERM	
	PNFT=0-	
	TEICHG-NF-Q-) PNET=GP(GRID)*P(GRID)*TERM/CHG	
1.8	DNH4(G91))= -PNET/C\$N9 + 7EXCN#XLIMZ + ANEXCN#XLIMA	UGAT/L
••	DPD4(GRID) = -PNET/CSPP + 2EX(P+XLIM2 + ANIXCP+XLIMA	UGAT/L
	DSI(GRID) = -PNET/CSSIP+ 7EXCSI#XLIM7+ ANEXSI#XLIMA	
20		
	RETIRA	
	END	
	6.1977	

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SUBROUTINE PHYTO IMPLICIT REAL (I,K,N) INTEGER GRID, I.K, DAY, TCDAY/0/, LIM(3)/'N', 'P', 'S'/ 1,NX, NY, NZ, NXNY, IOUT REAL EXCREF(100),LTLIM, MALIM T=TEMP COMMON /CNTRL/MXGRID,IOUT,NX,NY,NZ,OX,DY,DZ,DT,DAY,HOUR,T(100) 1 /PHYT/GMAX(100),LTLIM(100),MXLIM(100),LSIGN(100),GP(100) 2 /CNVRT/ C\$NP,C\$PP,C\$SIP 3 /VALUES/ P(100), NH4(100),NO2NO3(100),PO4(100),SI(100) DATA 10PT/150./, F/.5/, RADN/300./, K0/.025/, KSN/1.5/, 1 KSP/.25/, KSSI/.75/ NOT ALL STEPS NECESSARY IN EACH ITERATION --TEMP - DEPENDENT GMAX DETERMINED ONE TIME FOR CUNSTANT TEMP FIELD 15 (100) NO 2000 C C IF(TODAY.NE.01GO TO 20 NXNV#NX#NV IOPT SHOULD BE DETERMINED FOR THE WATER COLUMN AND SEASON, ETC. C DATA DTT/4.166667E-2/ DAYS/HR DO 10 GRID=1,MXGRID EPPLEY GMAX(GRID)=EXP(0.063+T(GRID)-0.16) GMAX(GFID)=GMAX(GFID) + UTT COMPUTE NUTRIENT CONVERSIJN FACTORS FROM RATIO BY ATOMS: / HR 10 C:N:S1:P С (UG-AT NUTR)/(1.272 MGC PER 106 UG-AT C) = UG-AT/MG C С C\$NP =1.272/C\$NP C\$PP =1.272/C\$PP MGC/UGAT C\$SIP=1.272/C\$SIP 20 CONTINUE č IF(DAY.FQ.TODAY)GO TO 50 TODAY => DAY ONLY DURING FIRST TIME-STEP OF EACH DAY TODAY=DAY DETERMINE LILIM ONE TIME DAILY С TERM=2.72*F LTLIM IS COMPUTED IN SEQUENCE FOR GRIDS DOWN THRU THE WATER COLUMN С 00 30 I=1,NX DO 30 J=1,NY J1=(J-1)*NX I1=RADN DU 30 K=1,NZ CONVERT FROM 3-DIM SUBSCRIPTS TO LINEAR LOCATIONS С GRID=I+JI+(K-1)+NXNYEXCDEF(GRID)=K0+0.2514+P(GRID)+0.5047+(P(GRID)++0.606667) RILEY PILEY EON. DRIG.: .008BICHL UG/L) + .054(CHL**.67) ASSUMING C:CHL * 35. KO BASED UN 6 SAMPLES FRCM PERU, = 0.025 1/M С С

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```
KZ=EXCCEF(GR10)+DZ
          TERM1=11/IOPT
          EKZ=EXP(-KZ)
          TERM2=TERM1+EKZ
          LTLIM(GRID)=TERM/KZ*(FXP(-TEPM2)-EXP(-TERM1))
IF(I1.GE.ICPT)LTLIM(GRID)=LTLIM(GRID)*0.85
                                                                                                                         KREMER
          11=11+EKZ
          CONTINUE
30
50
          CONTINUE
C
C
         WUTRIENT CALCULATIONS MAY NOT NEED TO BE MADE HOURLY, ...
NUTRIENT LIMITATION -- 3 NUTRIENTS
NH4 PREFERENCE MAY BE INCLUDED WITH UPTAKE SECTION
с
с
          DO 100 GRID=1.MXGRID
          NTUT=NH4(GRIU)+NU2NU3(GRID)
         NLIM \pi_N TOT /(KSN +NTOT )

PLIM = PO4(GRID)/(KSP + PO4(GRID))

SILIM = SI(GRID)/(KSSI+ SI(CRID))

MXLIM IS THE LIMITATION FACTOR OF THE MOST LIMITING NUTRIENT
¢
      MXLIMIGRID)=AMINI(NLIM,PLIM,SILIM)
WALSH METHOD OF IDENTIFYING MXLIM: VS. 16:10:1 N:SI:P
IF(P04(GRID).GT.O.)NRATIC=(NTOT /P04(GRID))/16.0
C
r:
          IF(PU4(GRID).GT.O.)SIRATO=( SI(GPID)/PU4(GRID)//10.0
С
ċ
          MXLIM(GRID)=PLIM
         IF(NRATIO.LT.1.0) MXLIM(GRID)=NLIM
IF(SIRATO.LT.NRATIO)MXLIM(GRID)=SILIM
LSIGN IDENTIFIES WHICH NUTRIENT IS NOW LIMITING
С
Ċ
С
         LSIGN(GRID)=LIM(L)
          IF(MXLIM(GRID).EQ.PLIM
                                                         HISIGN(GRID)=LIM(2)
          IF(MXLIN(GRID).EQ.SILIM
                                                         ILSIGN(GRID)=LIM(3)
         GP(GRID)=GMAX(GRID)+LTLIM(GRID)+MXLIM(GRID)
100
         CONTINUE
         IF([OUT.EQ.1)
        IWRITE(6,105) OUTEX, OUTLT, OUTMX, OUTGP, LSUUT

FORMAT(*1 K*/ 7(10X,10F7.4/), *0 LT*/ 7(10X,10F7.4/),

1 *0 MAX*/ 7(10X,10F7.4/), *0 GP*/ 7(10X,10F7.4/),

1 *0NUTR*/ 7(10X,10A4 /1)
105
         RETURN
         END
```

```
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```

SUBROUTINE PROFIL SUBROUTINE PROFIL INTEGER LASPC(7)/'FITO','DPLA','NCTO','N ','MG C','/L ',' '/ L & ABCHL(7)/'CHLO','ROFI','LA ',' UG ','CHL/','L ',' '/ L ABC14(7)/'PROD','UCC1','ON ',' MG ','C/M3','/DIA',' '/ L ABNH4(7)/'AMMO','NIAC','O ','UG-A','T/L ',' ',' '/ L ABNOX(7)/'NITR','ATOS',' Y N','ITRI','TOS ',' UG-','AT/L'/ L ABSOX(7)/'FOSF','ATOS',' ','UG-A','T/L ',' ',' '/ L ABSOX(7)/'FOSF','ATOS',' ','UG-A','T/L ',' ',' '/ L ABSOX(7)/'SILI','CATO','S ','UG-A','T/L ',' ',' '/ L ABSOX(7)/'ASSI','MILA','TION',' MG ','C/MG',' CHL','/DIA'/ L ABOPP(7)/'PROD','UCCI','ON C','OMMU','NIOA','D C/','M3/D'/ INTEGEO DAY 1 2 3 4 5 6 7 8 INTEGER DAY REAL LILIM, MXLIM, NH4, NO2NU3 1, CHL(100 }, AR(100)/100 *0./, PN(100) COMMON /CNTRL/MXGRID,IOUT,NX,NY,NZ,DX,DY,DZ,DT,DAY,HOUR,TEMP(100) 1 /PHYT/GMAX(100),LTLIM(100),MXLIM(100),LSIGN(100),GP(100) 2 /CNVRT/ C\$NP,C\$PP,C\$SIP 3 /VALUES/ P(100), NH4(100),NO2NO3(100),PO4(100),S1(100) 4 /CHANGE/DP(100), DNH4(100),DNOX(100),DPO4(100),DS1(100) С С MXGRED=70 C С PRINT OUT FINAL VALUES AND RATES 00 100 I=1, MXGRID С CHLOROPHYLL DISTRIBUTION DATA CSCHL/35.0/ DATA C3CHL735.07 CHL(1)=P(1)/C\$CHL *1000. NET PHYTG. PRODUCTIVITY (MG C/M3.DAY; =C-14) PN(1)=P(1)*1000.*(EXP(24.*GP(1))-1.0) ASSIMILATION RATIO -- UG C/UG CHL/DAY 20 C 30 L/H-M3/D Ċ 40 [F(CHL(1).GT.O.)AR(1)=PN(1)/CHL(1) #24. /HR-/DAY C NET COMMUNITY PRODUCTION -- PRODUCTIVITY + GRAZING DP(I)= 1000. * EXP(DP(I)=24.0) 50 L/H~M3/D 100 CONTINUE WRITE (6,200)LABCHL, (CHL(I),I=1,MXGRID) WRITE(6,200)LABC14, (PN(I),I=1,MXGRID) WRITE(6,200)LABCL4, (AR(I), (=1,MXGRID) WRITE(6,200)LABDPP,(DP(1),1=1,MXGRID) RETURN 200 FORMAT(1H1,T41,744///'0',8X,'_',1X,99('_')/7X,' 0 [',T111,']'/

 FORMAT(1H1,T41,744///0',0',0X,"_',1X,99('_')/7X,* 0 |*,T111,*[*/

 10X,*[*,10(F3.3,2X),T111,*[*/

 10X,*[*,T111,*[*/* P 10_[*,T111,*]*/10X,*[*,T111,*]*/

 10X,*[*,T111,*[*/* P 10_[*,T111,*]*/10X,*[*,T111,*]*/* 0 20_[*,

 * R*,7X,*[*,10(F8.3,2X),T111,*[*/10X,*[*,T111,*]*/* 0 20_[*,

 T111,*[*/*00,*]*,T111,*[*/* F*,7X,*]*,10(F3.3,2X),T111,*[*/10X,

 [,T111,*]*/* U 30_[*,T111,*]*/10X,*[*,T111,*]*/* N*,7X,*]*,

 10(F8.3,2X),T111,*[*/10X,*]*,T111,*]*/* U 40_[*,T111,*]*/10X,

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T X-GRID 1

	14.000 14.000 14.000 14.000 13.000 13.000 0.0 0.0	15.030 14.030 14.030 14.030 14.033 13.030 13.030 0.0 0.0 0.0	16.000 15.000 14.000 14.000 14.000 14.000 13.007 0.0 0.0	17.000 16.000 15.000 14.000 14.000 14.000 13.009 0.0 0.0	15.000 17.000 15.000 14.000 14.000 14.000 13.000 0.0 0.0	19.000 18.000 15.000 15.000 14.000 14.000 13.000 0.0 0.0	19.000 18.000 15.000 14.000 14.000 13.000 0.0 0.0 0.0	19.000 18.000 15.000 14.000 14.000 14.000 13.000 0.0 0.0	19.000 13.000 15.000 15.000 14.000 14.000 13.000 0.0 0.0 0.0	19.000 18.000 15.000 14.000 14.000 14.000 13.000 0.0 0.0
٧Y	X-GRID 1									
	549.090 144.000 -144.000 -144.000 -144.000 -252.090 9.0	1224.000 324.000 -324.000 -324.000 -324.000 -324.000 -324.000 -326.000 -396.000	1+04-000 396-000 -360-000 -360-000 -360-000 -360-000 -324-000	1440.003 432.000 -360.000 -360.000 -360.000 -360.000 -360.000 -432.000	1449.000 432.000 -360.000 -360.000 -360.000 -360.000 -360.000 -432.000	1440.00 432.00 -360.00 -360.00 -360.00 -360.00 -360.00 -432.00	1440.000 +32.000 -360.000 -360.000 -360.000 -360.000 -360.000 -432.000	1440.000 432.000 -360.000 -360.000 -360.000 -360.000 -437.000	1440.000 452.000 -360.000 -360.000 -366.000 -360.000 -360.000 -432.000	1440.000 432.000 -360.000 -360.000 -360.000 -360.000 -432.000
vz	K-GRID 1									

- ∪. 5∻0	-3.144	-0.035	0. 0	0.0	0.0	U. 0	0.0	7. 0	0.0
-1.224	-0.324	-0.193	0.0	0.0	0.)	0.0	0.0	() . ()	3.0
-1.224	-0.324	-0.144	0.J	0.0	J _J	0.0	3.0		0.0
-0.936	-0.252	-0.14+	0.0	0.0	0.0	0.0	0.0	0.0	0.0
-0.648	-0.140	-0.144	ن.(ა.ა	J.)	0.0	9.2	0.0	1.1
-0.252	-0.252	-0.103	0.0	ປ.0	3.0	0.0	0.0	ບູ້ບ	3.0
0.0	-0.130	-0.336	0.0	0.0	0.0	0.0	0.0	3.7	4.9

INPUT DATA READ FROM SET 5, EFFECTIVE START-UP ON DAY 1

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```
SUBROUTINE STOYST
Integer Day
Real Chg(6)/60+0./
           INTEGER GRID
COMMON /VALUES/VAL(500 )
1 /CNTRL/MXGRID,IOUT,NX,NY,NZ,DX,DY,CZ,DT,DAY,HOUR,T(100 )
DATA SUM20/10000./
DATA STABLE/0.001 /
DATA 1/0/
              1=1+1
             SUM=0.
NO 10 GRID=1.20
SUM=SUM+VAL(GRID)
 10
              CONTINUE
             CHANGE=(SUM2O-SUM)/SUM
SUM2O=SUM
CHG(I)=CHANGE
IF/DAY.E4.51)WRITE(6,30)CHG
              IF (ABS(CHANGE).GT.STABLEIRETURN
           WRITE(6,20)CHANGE, DAY, HOUK
FORMAT(*0 STEADY-STATE CRITERION MET, RUN TERMINATED */
1 *0 TOTAL CHANGE IN TOP 20 GRIDS =*,E15.5,* OF PREVIOUS TOTAL'/
1 *0 TIME AT TERMINATION: *, 15,F5.1)
20
           WRITE(6.30)CHG
FORMAT('O SEQUENCE OF CHANGE VALUES LEADING TO STEADY STATE:'/
1 (1X,10E12.4))
DATA 10SET/6/
IF(10SET.NE.6)WRITE(10SET,915)DAY,VAL
30
            WRITE(10,916)IOSET

FORMAT(15/(10FR.5))

FORMAT(*0 ===> OUTPUT INITIAL CONDITIONS TO SET*,15/)

CALL PROFIL

STOP 1
915
916
```

- END

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.*!',TI11 !'/' I',TX,*!',10(Fd.3,2X),TI11,*!'/10X,*!',TI11,*!'/
.* 0 50_!',TI11,*!'/10X,*!',TI11,*!'/' A',TX,*!',10(F8.3,2X),
.TI11,*!'/10X,*!',TI11,*!'/' D 60_!',TI11,*!'/10X,*!',TI11,*!'/
.10X,*!',10&F8.3,2X),TI11,*!'/10Y,*!',TI11,*!'/' (M) 70_!',TI11,
.*!'/ 3(10X,*!',TI11,*!'/3,TX,*80_!',TI11,*!'/' (M) 70_!',TI11,*!'/')
..TX,*90_!',TI11,*!'/3(10X,*!',TI11,*!'/),6X,*100_!',99(*_'),*!'/
.11X,10(9X,*!')/10X,*10',3X,*20',3X,*30',8X,*100',8X,*50',8X,*60',6X
.**T0',8X,*80',8X,*70',8X,*100'//T44,*DISTANCIA JE LA COSTA (KM)*)
END

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DAY	1	HOUR 1.00	CONCENTRA	TTON PPL	X GRID 2	2					
	0.041	0.041	0.040	0.006	Ū.0	0.0	0.0	0 0	• •		
	0.041	0.041	0.041	0.002	9.000	0.000	0.000	0.000	0.0	0.0	
	0.041	0.041	0.039	0.010	J.010	0.010	0.010	0.010	0.000	0.000	
	0.040	0.040	0.039	0.010	0.010	0.010	0.010	0.010	0.010	0.010	
	0.040	0.040	0.039	0.010	0.010	0.010	0.010	0-010	0.010	0.010	
	0.039	0.040	0.039	0.010	0.010	0.010	0.010	0.010	0.010	0.010	
	0.010	0.039	9.039	0.010	0.010	0.010	0.010	0.019	0.010	0.010	
					01010	0.010	0.010	0.010	0.010	0.010	
DAY	1	HUUR 1.00	CONCENTRA	TION NH4	X GRID 2						
	0.493	0.493	0.492	0.500	0.500	0.500	0 500	0 500			
	0.487	0.487	0.486	0.500	0.500	0 500	0.500	0.500	0.500	0.500	
	0.492	0.492	0.492	0.446	0 496	0.404	0.500	0.500	J. 500	0.500	
	0.496	0.496	0.456	0.496	0.496	0.490	0.496	0.496	0.496	0.496	
	0.498	0.498	0.498	0.497	0.407	0.495	0.496	0.496	0.496	0.496	
	0.499	0.499	0.499	0 497	0.471	0.497	0.497	0.497	Ů.497	0.497	
	0.500	0.500	0 500	0 400	0.498	0.498	0.498	0.498	0.498	0.498	
		00000	0.000	0.499	0.444	0.499	0.499	0.499	0.499	0.499	
DAY	1	HOUR 1.00	CUNCENTRA	TION NOR	X 6310 2						
	1.184	0.808	0.705	0-671	0.671	0 671					
	10.531	10.081	9.973	9.917	9,919	9 919	0.011	0.671	0.671	0.671	4
	15.612	15.162	15.072	15.000	15.000	15 000	7.717	9.919	9.919	9.919	P
	20.468	20.126	20.07?	20.000	20.300	20.000	19.000	15.000	15.000	15.000	•
	25.518	25.144	25.115	25.000	25.000	20.000	20.000	20.000	23.000	20.000	
	33.000	33.000	33.000	33.000	23.000	23.000	25.000	25.000	25.000	25.000	
	33.000	33.000	33,000	33-000	33.000	33.000	33.000	33.000	33.000	33.000	
				23.000	33.000	33.000	33.000	33.000	33.000	33.000	
DAY	1	HOUR 1.00	CONCENTRA	TION PO4	X GRID 2						
	0.428	0.413	0.408	0.407	0.407	0.407	0.4.37	0 407			
	0.648	0.812	0.803	0.800	0.800	0.300	0.800	0.900	0.407	0.407	
	1.248	1.212	1.205	1.200	1.200	1.200	1 200	1 200	0.800	0.800	
	1.637	1.610	1.606	1.600	1.600	1.600	1 6 30	1.200	1.200	1.200	
	2.032	2.009	2.007	2.000	2.000	2 000	1.000	1.600	1.600	1.600	
	2.500	2.500	2.500	2.500	2.50.0	2.000	2.000	2.000	2.000	2.000	
	2.500	2.500	2.500	2.500	2.500	2.500	2.500	2.500	2.500	2.500	
	,			20203	2.00	2.500	2.500	2.500	2.500	2.500	
)A Y	1	HOUR 1.00	CONCENTRA	TION SI	X GRID 2						
	3.068	3.028	3.017	3.018	3.018	3.618	3.018	3 016	2 010		
	4.395	4.125	4.059	4.036	4-036	4.036	2.010	3.018	3.018	3.018	
	7.362	7.092	7.038	6.997	6.997	6.997	4.030	*•036	4.036	4.036	
	10.279	10.073	10.041	9.997	9.497	9.007	0.77/	0.331	6.997	6.997	
	13.777	13.215	13.172	12.998	12.998	17 000	12 000	9.997	9.997	7.997	
	25.000	25.000	25.000	24.999	74.000	74 000	12.998	12.998	12.998	12.998	
;	25.000	25.000	25.000	24,999	24.900	24 000	24.999	24.999	24.999	24.999	
					270777	24833A	24.999	24.999	24.999	74.999	

CHLOROFILA	UG	CHL.	/ι
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				DISTANCES	DE LA COS	TA (244)				
100_1	 10	1 20	 30	1 40	 50	 1 60	1 70	ا رو	 1 90	اا ا 10
100	_									
90_										
30_ 	E 									
70_										
	 0.296	0.345	:)+405	0.419	0.419	0.419	0.41 9	0.419	0.419	0.419
60_	C_389	0.453	0.513	0.549	C.550	0.550	J .5 49	0.549	0.549	3.549
5J_	0.461 	0.626	J.710	0.711	0.783	0.731	0.779	0.778	0.778	0.778
40_										_
J.J	0.579	0.975	1.196	1.360	1.357	1.346	1.338	1.334	1.332	1.331
30	1 0.808	1.781	2.293	2.545	2.446	2.368	2.323	2.298	2.285	2.280
20_	1.163	2.137	2.960	3.698	4.258	4.747	5.211	5.690	6.196	6.701
10_										
0	2.082	2.64-	3.386	4.317	5.431	6.712	8.035	9.370	16.697	11.973

Objectives of the economic study were three:

1. planning with the staff of the Instituto del Mar, Peru (IMARPE) for an economic analysis of issues of interest, using available data

2. reviewing fishery economic issues in the Ilo, Peru, area with special reference to (a.) artisanal fisheries, and (b.) the production, distribution, and marketing of fish for human consumption

3. exploring the possibility of constructing an economic model capable of interfacing with the ecosystem model to be developed by Scott Nixon and James Kremer.

These general objectives were achieved, though each to a different extent. As should become clear in the next section, my visit focused on objectives 1. and 2. Constructing the model in 3. probably is not feasible in the near future due to the short time allocated and lack of appropriate data on the food fishery.

Results

Following discussions with persons in the Ministry of Fisheries and related entities (such as EPSEP and Pesca Peru), with artisan fishermen and the IMARPE laboratory staff in Ilo, Dr. W. Macedo and I drew up a tentative research program for his office.

The tentative program consists of a short-term and a long-term program of economic research. The short-term program was the major focus of concern, and although developed in some detail it still needs refinement as well as an indication of support by IMARPE. It calls for a study of the artisanal fishery around the port of Ilo (including Villa Villa and Meca) with the objective of making recommendations for the development of the fishery. (A tentative outline of such a study is presented in table 1.) Much of the information necessary for the above study is contained in an existing paper by A. Pastor and N. Gelarza, <u>Estudio Preliminar Sobre La Pesca Artesanal En El Puerto De Ilo</u>, IMARPE, 1974. Additional information was gathered during our recent visit to the Ilo area.

However, information on the costs and earnings of capital, and on the earnings and characteristics of the fishermen has not yet been collected. Since this information is a critical part of the study, we believe its collection should begin immediately. The first step necessary for collecting such data is to develop an appropriate form for the field staff to use. Examples of such forms were left at IMARPE. The appropriate form will be developed by the IMARPE economist in consultation with the Ilo field staff, who will be responsible for collecting the data. Development of these forms can begin at once without adding much of a burden on IMARPE's staff.

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Table 1. A tentative outline of an economic study of the Ilo artisanal fishery.

- I. Introduction
- II. Description of the fishery
 - A. The resource (magnitude, location, species, etc.)
 - B. The harvesting sector
 - 1. Output and prices (past and present)
 - 2. Capital inputs
 - a. description of gear, techniques, and the enterprises' operations
 - b. costs and earnings of the enterprises
 - c. supporting infrastructure (repair and construction,
 - 3. Labor inputs
 - a. description of skills, education, ethnic and social status, and alternative economic opportunities
 - b. earnings
 - c. supporting infrastructure (schools, housing, hospitals,
 - etc.)
 - C. Processing
 - D. Distribution
 - E. Marketing
- III. Prospects for Development
 - A. Resource potential
 - B. Harvesting sector
 - 1. New vessels, gear and techniques

 - Fishermen training
 Supporting infrastructure development
 - C. Processing
 - D. Distribution E. Marketing
- IV. Recommendations for development and future research

Data collection will be for an appropriate sample size and composition For example, 25 vessels and their crews may suffice; however, other considerations may call for this number to be reduced or expanded. The vessels in the sample should be chosen to capture a variety of vessel sizes, gear types, major species caught, locations of operation, etc. The costs and earnings of these enterprises will be monitored for a full fishing season and may be collected at the end of every fishing trip along with the current collections of landings quantity, composition, and value data.

The parts of the study concerning processing, distribution and marketing probably will be of a cursory nature for two reasons: 1. processing, distribution, and marketing are more naturally the concern of EPSEP, the processing, distribution, and marketing arm of the Fisheries Ministry, and 2. whereas IMARPE routinely collects catch statistics and therefore has established close links with the harvesting sector, it would require a substantial increase in personnel to monitor the other activities. Of course, IMARPE personnel would be remiss if they did not attempt to obtain as much of this information as possible from EPSEP. However, any major study of processing, distribution and marketing should be carried out by EPSEP or another branch of the Fisheries Ministry, or at least postponed until IMARPE develops the capability to thoroughly study these issues.

A logical question at this point is how will this study meet its objectives, i.e., what are some recommendations for the fishery's development likely to emerge from the study? From the analysis of costs and earnings data on existing and new forms of capital, the study will likely prescribe any beneficial changes in the types of vessels, gear and techniques to be used. The study will propose actions to resolve any problems of financing vessels and gear, of repairing and constructing vessels, of supplying gear and other equipment, etc. Similarly with labor, the study will likely assess the manpower needs for developing the fishery, i.e., the number of fishermen and the types of skills requires, and propose means of providing this manpower, i.e., the wage level, training, housing, schools and medical facilities needed. In the areas of processing, distribution and marketing, the study will at least identify major problem areas, and suggest the necessary research to resolve these problems.

The second, or long-term, part of the research program is only speculative. It is not clear at this time what IMARPE's role will be in fishery economic research in the distant future. What is clear, however--or at least becoming more evident to a number of Peruvian researchers and administrators--is that much more economic research is required for the proper development and management of the exploitation of Peru's fishery resources. Inter alia, the monitoring and analysis of costs and earnings in the industry will be an important part of such research. Therefore, in the long run IMARPE may wish to develop the capability to routinely collect and analyze costs and earnings data. If so, the Ilo research program can be regarded as a first step in developing such a capability.

Role of AID/ICMRD in Peruvian Fishery Economic Research

While there appears to be a substantial interest in expanding Peru's fishery economic research capabilities, no single entity at this time is engaged in developing a comprehensive economic research program. Just how, when and where such a program will be developed is not clear. Given that such a program will be developed in the future, however, some thought and commitment should be made soon regarding training Peruvian economists and outlining a comprehensive research program. AID could play a meaningful role by funding the training of Peruvian fishery economists at U.S. universities. Although FAO has played a major role in economic research in the past, AID, it seems, can fill a valuable niche by providing skilled advisors to develop the research program and to conduct some of the research.

Should the above scenario be realized, the implications for ICMRD and the Department of Resource Economics are obvious. However, in the very near future there exists a need for continued resource economics involvement. If the above study is to be carried out, more refinement of its exact contents is necessary. I am sure further analytical assistance will be desired by IMARPE's economist. Therefore, a request for additional resource economics involvement is likely to be forthcoming in the near future.

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Jon G. Suti	nen		
<u>Time</u> (Days)	<u>Dates</u> (1975)	<u>Place</u>	
18	20 Jan-7 Feb	Peru	Site visit to assist Instituto del Mar personnel in planning economic analysis; review fishery economic issues in the Ilo area with reference to artisanal fisheries and to production, distribution and marketing of fish.
James N. Kr	emer		
<u>Time</u> (Days)	<u>Dates</u> (1975)	Place_	
17	1-17 Feb	R. I.	Preparation phase: review physical characteristics of study area and available biological data supplied by Instituto del Mar personnel; write computer program for physical circulation model.
22	18 Feb-11 Mar	Peru	Discuss circulation model; formulate and analyze basic conceptual phyto- plankton model. Survey available data with preliminary hand calcula- tions comparing theoretical formula- tions with observed data. Preliminary estimates of some parameters appro- priate to the Peru system.
49	12 May-29 Apr	R. I.	Development of phytoplankton sub- model as discussed in Peru; develop- ment of ecological model framework to complement PHYTO submodel, including herbivore (anchovy and zooplankton) grazing and excretion influences. Preliminary computer simulation runs and evaluation.
20	30 Apr-19 May	Peru	Analyze and evaluate final model formulations. Discuss suggested modifications in the model and research and field observations. Discuss with Instituto del Mar personnel the technical details of the model methods. Present lecture on ecosystem model.
4	20-23 May	R. I.	Preparation of final report

University of Rhode Island - Instituto del Mar Project Schedule

Scott W. Nixon, Assistant Professor at URI. Ten days, 7-16 May, in Peru, consultation and discussion with Dr. Guillen of Instituto del Mar on modeling and ecosystem analysis. Present lecture on role of modeling in ecosystem research. Seven days, 17-23 May, in R. I. assisting in preparation of final report.

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