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COMMERCIAL FISHING WITH PREDATOR-PREY INTERACTION

Koji OKUGUCHI*

Abstract: The Smith model of commercial fishing of a single species is generalized, taking into consideration of natural interaction between a prey and a predator. The stability of the non-extinct steady state populations of the two species is proved, and the effects of changes in parameters upon the steady state are examined. Finally, the steady state populations under harvesting and no harvesting are compared.

1. INTRODUCTION

A dynamic model of commercial fishing incorporating economic as well as biological aspects of a single species has been formulated and analysed by Smith [4], and the stability of the steady state or bionomic equilibrium has been investigated in great detail by Leung and Wang [3] and Wang and Cheng [6] under the assumptions that the price of the fish is a strictly decreasing function of the total harvest and that the total cost excluding the opportunity cost per boat or investment is proportional to the square of the harvest per boat or investment and inversely proportional to the stock or population of the fish.

Solow [5], on the other hand, has attempted a comparative dynamical analysis of fishing within a framework of an optimal control theoretical model involving natural predator-prey interaction à la Volterra. Clark [1] also has studied a similar model involving a Gause-type interspecific competition between two species of fishes which feed on a common food supply, each species growing according to a logistic law in the absence of its competing species. Moreover, he has analysed a model with predator-prey interaction à la Larkin. Both Solow and Clark have assumed the constancy of prices of fishes harvested over time. More recently, May, Beddington, Clark, Holt and Laws [4] have analyzed multispecies fisheries assuming intertemporal constancy of fishing efforts. In addition to a prey-predator (krill-baleen whale) model, they have considered one prey-two predator (krill-whale and seal) model and a three trophic level (krill, cephalopod and sperm whale) model. They have made some observations on economic aspects of multispecies fisheries, but have not analyzed the optimal determination of fishing efforts from the viewpoint of economic rationality of fishing firms.

The purpose of this paper is to extend the Smith model of commercial fishing of a single species taking into account natural interaction between a predator and a

* Thanks are due to a referee who has suggested a simple proof of our stability result.

prey. In Section 2 we present a commercial fishing model involving predator-prey interaction, and analyze the stability of the steady-state with non-extinct populations of the two species. In Section 3 a comparative statical analysis of the steady state with respect to changes in parameters is given. Further, the populations of the two species in the steady are compared for the case of harvesting and that for no harvesting. Section 4 concludes.

2. THE MODEL AND A DYNAMIC ANALYSIS

Let there be two species of fishes in the same territory, one as predator and the other as prey, and let X_1 and X_2 denote the populations of the prey and predator, respectively. If uninterrupted by fishing, the two species grow according to

$$(1) \quad \dot{X}_1 = X_1(a_1 - b_1 X_1 - c_1 X_2),$$

$$(2) \quad \dot{X}_2 = -X_2(a_2 - b_2 X_1 + c_2 X_2),$$

where the parameters are assumed to satisfy:

$$(3) \quad \begin{aligned} a_1 &> 0, & b_1 &> 0, & c_1 &> 0 \\ a_2 &\geq 0, & b_2 &> 0, & c_2 &\geq 0. \end{aligned}$$

The meanings of these equations and restrictions are as follows. The prey species grows in accordance with a logistic law in the absence of the predator, and the number of prey predated by the predator is proportional to the number of the predator for given number of the prey and to the number of the prey for given number of the predator. The predator, on the other hand, grows according to a logistic law also in the absence of the prey if $a_2 < 0$ and $c_2 > 0$ are satisfied simultaneously. The additional increase in the predator population in the presence of predation is proportional to the number of the prey for given number of the predator and to the number of the predator for given number of the prey.

Solow has analysed the Volterra case where $a_2 > 0$, $b_2 > 0$ and $c_2 = 0$. In this case the predator decreases exponentially without predation. The case where $a_2 < 0$, $b_2 > 0$ and $c_2 > 0$ has been considered by Larkin (see Clark [1, Chap. 9]), and the case where $a_2 > 0$, $b_2 > 0$ and $c_2 > 0$ has been treated by Hirsch and Smale [2, Chap. 12]. May, Beddington, Clark, Holt and Laws [4] have assumed the following differential equation for the growth of the predator, instead of (2):

$$(2)' \quad \dot{X}_2 = r_2 X_2 (1 - X_2 / \alpha X_1),$$

where $r_2 > 0$ is the intrinsic growth rate of the predator and $\alpha > 0$ is a proportionality constant relating the carrying capacity of the predator to the stock of the prey when the predator is not harvested.

Let us now introduce selective harvesting of the two species of fishes. In contrast to Smith [5], Leung and Wang [3], and Wang and Cheng [7], we assume à la Solow [6] and Clark [1] that both species of fishes are sold at constant prices, p_1 and p_2 ,

over time. Let Y_1 and Y_2 be the amounts of harvest of the prey and predator, respectively. The selective harvesting cost per boat or investment are given by $\gamma_1 Y_1^2/X_1$ and $\gamma_2 Y_2^2/X_2$, respectively, for the prey and predator. Profit per boat or investment for fishing the i -th species (the prey for $i=1$ and the predator for $i=2$) is given by

$$(4) \quad \pi_i = p_i Y_i - \gamma_i Y_i^2/X_i - \pi_i^0, \quad i=1, 2,$$

where π_i^0 is the opportunity cost per unit of investment in fishing the i -th species.

Maximising π_i with respect to Y_i , we get¹

$$(5) \quad Y_i = p_i X_i / 2\gamma_i, \quad i=1, 2,$$

which substituted into (4) yields

$$(6) \quad \pi_i = p_i^2 X_i / 4\gamma_i - \pi_i^0, \quad i=1, 2.^2$$

Let K_i be the number of boats engaged in fishing the i -th species. In the presence of fishing the natural growth equations (1) and (2) have to be modified as:

$$(7) \quad dX_1/dt = X_1(a_1 - b_1 X_1 - c_1 X_2) - K_1 Y_1$$

$$(8) \quad dX_2/dt = -X_2(a_2 - b_2 X_1 + c_2 X_2) - K_2 Y_2,$$

or in view of (5), as

$$(9) \quad dX_1/dt = X_1(a_1 - b_1 X_1 - c_1 X_2 - p_1 K_1 / 2\gamma_1)$$

$$(10) \quad dX_2/dt = -X_2(a_2 - b_2 X_1 + c_2 X_2 + p_2 K_2 / 2\gamma_2).$$

We assume that the number of boats engaged in fishing increases or decreases in

¹ It might be more faithful to current practice to consider maximization of the integral of discounted profit than to consider current profit maximization. Technical difficulties associated with the former maximization, however, forces us to be satisfied with the latter maximization as a first approximation. Moreover, this approximation is valid if the discount factor is sufficiently large, or if future is very uncertain.

² Let the demand function and the harvesting cost per boat be given by $p_i = f^i(Y_i)$ and $C^i = C^i(Y_i, X_i)$, respectively, where $f_{Y_i}^i < 0$, $C_{Y_i}^i > 0$ and $C_{X_i}^i < 0$, $i=1, 2$. The profit per boat for fishing the i -th species is

$$(a) \quad \pi_i = X_i f^i(Y_i) - C^i(Y_i, X_i) - \pi_i^0, \quad i=1, 2.$$

Maximizing π_i with respect to Y_i , we get

$$(b) \quad Y_i \equiv h^i(X_i), \quad i=1, 2,$$

where

$$(c) \quad dh^i/dX_i \geq 0 \quad \text{according as} \quad C_{Y_i X_i}^i \leq 0, \quad i=1, 2.$$

Substituting (b) into (a),

$$(d) \quad \pi_i = h^i(X_i) f^i(h^i(X_i)) - C^i(h^i(X_i), X_i) - \pi_i^0, \quad i=1, 2.$$

More generally, the demand function should be formulated as

$$(e) \quad p_i = f^i(Y_i, Y_j), \quad i \neq 1, 2.$$

proportion to profit or loss per boat. Hence

$$(11) \quad dK_i/dt = \delta_i(p_i^2 X_i/4\gamma_i - \pi_i^0), \quad i=1, 2,$$

where δ_i 's are positive constants.

Equations (9)–(11) constitute our fundamental system of differential equations, the stability of which steady state being our main concern in this section. If dK_i/dt approaches zero independently of (9) and (10), the stability may be proved by appealing to the results already obtained by others. However, (9), (10) and (11) are interdependent, making our stability analysis rather complex.

Let X_i^* and K_i^* be the steady state values of X_i and K_i , respectively, where we assume that X_i^* and K_i^* are both positive. Expand (9)–(11) in the neighbourhood of X_i^* and K_i^* to derive the following system of linear differential equations in four variables.

$$(12) \quad \begin{pmatrix} dZ_1/dt \\ dZ_2/dt \\ dN_1/dt \\ dN_2/dt \end{pmatrix} = \begin{pmatrix} -b_1 X_1^* & -c_1 X_1^* & -p_1 X_1^*/2\gamma_1 & 0 \\ b_2 X_2^* & -c_2 X_2^* & 0 & -p_2 X_2^*/2\gamma_2 \\ \delta_1 p_1^2/4\gamma_1 & 0 & 0 & 0 \\ 0 & \delta_2 p_2^2/4\gamma_2 & 0 & 0 \end{pmatrix} \begin{pmatrix} Z_1 \\ Z_2 \\ N_1 \\ N_2 \end{pmatrix} \\ \equiv \begin{pmatrix} -a_{11} & -a_{12} & -a_{13} & 0 \\ a_{21} & -a_{22} & 0 & -a_{23} \\ a_{31} & 0 & 0 & 0 \\ 0 & a_{42} & 0 & 0 \end{pmatrix} \begin{pmatrix} Z_1 \\ Z_2 \\ N_1 \\ N_2 \end{pmatrix},$$

where

$$(13) \quad Z_i \equiv X_i - X_i^*, \quad N_i \equiv K_i - K_i^*, \quad i=1, 2,$$

and

$$(14) \quad a_{11}, a_{12}, a_{13}, a_{21}, a_{23}, a_{31}, a_{42} > 0, \quad a_{22} \geq 0.$$

If the steady value of either X_1 or X_2 is zero, then π_1^0 or π_2^0 has to be zero because of (11). We therefore assume away the possibility of extinction of both species of fishes, as the opportunity cost of investment in fishing can, in general, be considered to be positive. Note also that in deriving (12), use has been made of the following two equations relevant to the steady state.

$$(15) \quad a_1 - b_1 X_1^* - c_1 X_2^* - p_1 K_1^*/2\gamma_1 = 0,$$

$$(16) \quad a_2 - b_2 X_1^* + c_2 X_2^* + p_2 K_2^*/2\gamma_2 = 0.$$

The characteristic equation for the coefficient matrix of (12) is shown to be

$$\begin{aligned}
(17) \quad & \lambda^4 + (a_{11} + a_{22})\lambda^3 + (a_{11}a_{22} + a_{12}a_{21} + a_{13}a_{31} + a_{24}a_{42})\lambda^2 \\
& + (a_{11}a_{24}a_{42} + a_{13}a_{22}a_{31})\lambda + a_{13}a_{24}a_{31}a_{42} \\
& \equiv \lambda^4 + \alpha_1\lambda^3 + \alpha_2\lambda^2 + \alpha_3\lambda + \alpha_4 \\
& = 0 .
\end{aligned}$$

In view of the definition of α_1 , α_2 , α_3 and α_4 , a simple calculation shows that all of the upper left-hand corner principal minors of the matrix

$$\begin{pmatrix}
\alpha_1 & \alpha_2 & 0 & 0 \\
1 & \alpha_2 & \alpha_4 & 0 \\
0 & \alpha_1 & \alpha_3 & 0 \\
0 & 1 & \alpha_2 & \alpha_4
\end{pmatrix}$$

are positive. Hence by the Routh-Hurwitz stability criterion, the real parts of all characteristic roots of (17) are negative, establishing the local stability of the steady state for the system of differential equations (9)–(11). We should note here that we have been able to establish the stability independently of the sign of a_2 which may be of any sign.

3. COMPARATIVE STATICS

We now conduct a comparative statical analysis of the steady state with respect to changes in the values of the parameters. From (11) we have

$$(18) \quad X_i^* = 4\gamma_i \pi_i^0 / p_i^2, \quad i = 1, 2 .$$

This coupled with (15) and (16) yields:

$$(19) \quad K_1^* = (a_1 - 4\gamma_1 b_1 \pi_1^0 / p_1^2 - 4\gamma_2 c_1 \pi_2^0 / p_2^2) 2\gamma_1 / p_1 ,$$

$$(20) \quad K_2^* = (-a_2 + 4\gamma_1 b_2 \pi_1^0 / p_1^2 - 4\gamma_2 c_2 \pi_2^0 / p_2^2) 2\gamma_2 / p_2 .$$

By simple calculations we get the following Table 1.³

³ If the demand function and the harvesting cost are given as in the footnote 2, the steady state value of X_i is determined by

$$(f) \quad h^i(X_i^*) f^i(h^i(X_i^*)) - C^i(h^i(X_i^*), X_i^*) - \pi_i^0 = 0, \quad i = 1, 2 .$$

Totally differentiating (f), and taking into account the first order condition for maximization of (d), we have

$$(g) \quad dX_i^* / d\pi_i^0 = -1 / C_{X_i}^i > 0, \quad i = 1, 2 .$$

However, comparative statics for other variables and parameters are not as straightforward as for the table given above.

TABLE 1. COMPARATIVE STATICS

	X_1^*	X_2^*	K_1^*	K_2^*
p_1	—	0	?	—
p_2	0	—	+	?
π_1^0	+	0	—	+
π_2^0	0	+	—	—
c_1	0	0	—	0
b_2	0	0	0	+

Table 1 is not exhaustive, as we are interested only in the effects of changes in the values of some parameters. From Table 1 we can read, for examples, $\partial X_1^*/\partial p_1 < 0$, $\partial X_2^*/\partial p_1 = 0$, $\partial K_1^*/\partial p_1 \cong 0$, $\partial K_1^*/\partial p_2 > 0$, etc. We should note especially the asymmetrical sign patterns pertaining to changes in K_1^* and K_2^* . These asymmetries reflect the effects of the predator-prey interaction.

The populations in the steady state with and without harvesting can be compared on the basis of Figs. 1–5 as follows.

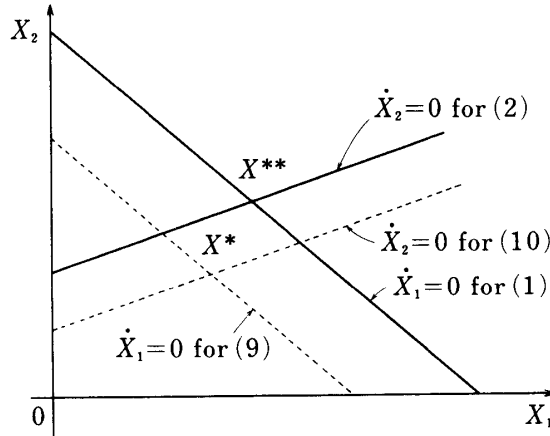


Fig. 1. Harvesting of both species.

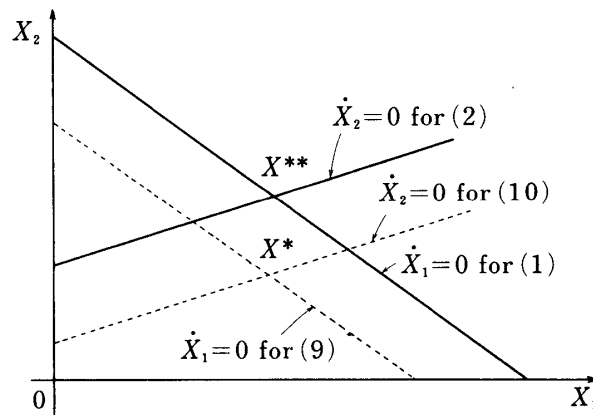


Fig. 2. Harvesting of both species.

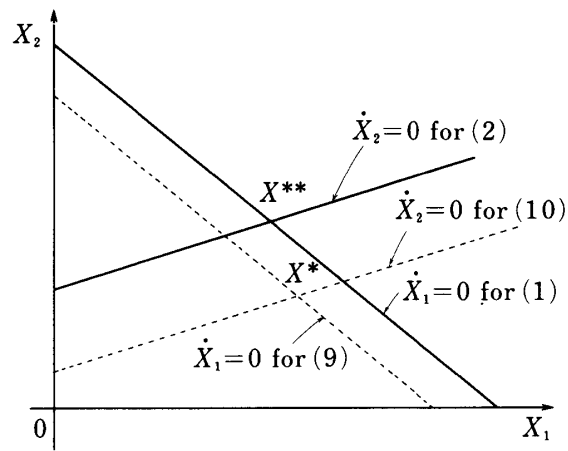


Fig. 3. Harvesting of both species.

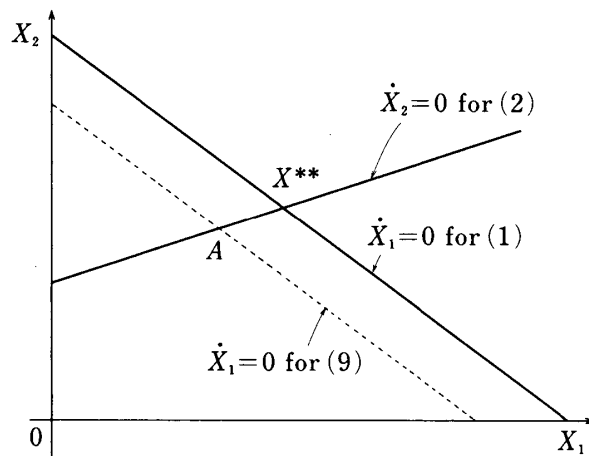


Fig. 4. Only prey is harvested.

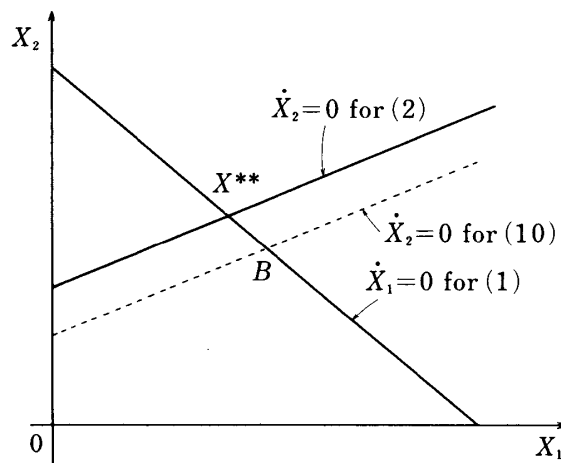


Fig. 5. Only predator is harvested.

Consider first the case where two species are both harvested. Let $a_2 < 0$,⁴ and let $X^{**} = (X_1^{**}, X_2^{**})$ be the pair of the steady state values of X_1 and X_2 in the absence of harvesting, and $X^* = (X_1^*, X_2^*)$ be that in the presence of harvesting. Three cases are possible (see Figs. 1–3). In these cases, we have unambiguously $X_2^* < X_2^{**}$. However, for the prey we may have $X_1^* < X_1^{**}$ (Fig. 1), or $X_1^* = X_1^{**}$ (Fig. 2), or $X_1^* > X_1^{**}$ (Fig. 3). If only the prey is harvested (Fig. 4), the steady state shifts from X^{**} to A , and the steady state populations of both species are decreased. If, on the other hand, only the predator is harvested (Fig. 5), the steady shifts from X^{**} to B . In this case, the prey population increases and the predator population decreases.

4. CONCLUSION

We have formulated a dynamic model of commercial fishing, as distinct from recreational fishing, allowing for existence of natural interaction between the predator and prey, and assuming that the prices of both species will remain constant over time during which fishes are harvested. Our model is different from that of May et al. [4] mainly in that economic rationality of fishing firms is taken into account. It has been found that the steady state or bionomic equilibrium with non-extinct populations of both species is locally stable, regardless of the signs of a parameter pertaining to the intrinsic growth rate of the predator. We have also given a comparative statical analysis of the steady state with respect to changes in exogeneous parameters. The steady state population of the predator in the presence of harvesting is smaller than that in the absence of harvesting, while the prey population in the presence of harvesting may be of any size compared with that in the absence of harvesting. The above comparison holds when both species are simultaneously harvested. Figures 4 and 5 are concerned with similar comparison when only one species is harvested.

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⁴ The case where $a_2 > 0$ may be similarly analyzed.

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