Dynamical Analysis of a Harvesting Model of Phytoplankton-Zooplankton Interaction

Anuj K. Sharma, Amit Sharma, Kulbhushan Agnihotri

Abstract—In this work, we propose and analyze a model of Phytoplankton-Zooplankton interaction with harvesting considering that some species are exploited commercially for food. Criteria for local stability, instability and global stability are derived and some threshold harvesting levels are explored to maintain the population at an appropriate equilibrium level even if the species are exploited continuously. Further, biological and bionomic equilibria of the system are obtained and an optimal harvesting policy is also analysed using the Pontryagin’s Maximum Principle. Finally, analytical findings are also supported by some numerical simulations.

Keywords—Phytoplankton-Zooplankton, Global stability, Bio nomic Equilibrium, Pontryagin-Maximum Principle.

I. INTRODUCTION

PLANKTON refer to all the plants and animals in marine environment that drift with the oceanic currents as inhabitants of the ocean water. Zooplankton, the planktonic animals, are all weak swimmers, whereas phytoplankton, planktonic plants, do not swim at all. They are the staple item for the food web and are producers and recyclers of most of the energy that flow through the oceanic ecosystem. The phytoplankton species in the pelagic zone are excessive small, microscopic and single-celled, buoyantly supported by the density of the surrounding water which include: Cyanophyta, Bacillariophyta and Xanthophyta. Plankton, especially the phytoplankton play an important role not only in aquaculture but they also stabilize the environment by consuming half of the universe carbon dioxide and releasing huge oxygen for the living organisms. Aquatic ecologist have long been fascinated by the non-equilibrium dynamics of explosive phytoplankton bloom explosions (i.e., the rapid explosions and declines in their population). Frequent outcome of a planktonic bloom formation, leads to massive cell lysis and rapid disintegration of large planktonic population. This is closely followed by an equally rapid increase in bacterial number, and in turn, by fast deoxygenation of water, which could be detrimental to aquatic plants and animals. Plankton remained fascinated area of the research for the last three decades but now these days some species are exploited for the food such as: Nori, Kelp and Eucheuma are phytoplankton and Jellyfish, Krill, and Acetes are zooplankton species. During recent years, many research models were applied to plankton system in the presence of nutrients and role of different functional forms in phytoplankton-zooplankton interactions were studied [1-7]. In [4] phytoplankton-zooplankton system was studied and it was concluded that the toxin producing phytoplankton may be used as controlling agents for the termination of plankton blooms. So far, not much attention is given on the impact of harvesting on the plankton system. To the best of author’s knowledge, recently Lv et al. [8] were the first to propose a harvesting model of toxin producing phytoplankton-zooplankton system in which local and global stability of the various equilibria were studied and concluded that over-exploitation of the system lead to the extinction of the species. Again Lv et al. [9] studied two zooplankton one phytoplankton model with harvesting. They concluded, in the absence of harvesting, the type of zooplankton with the higher biomass conversion ratio and the lower natural death rate persists only whereas harvesting may lead to the persistence of the type of zooplankton with the lower biomass conversion ratio and the higher natural death rate. Following along the lines of Lv et al. [8], in this paper, we assume both the populations grows logistically and the interaction of species are of Holling type-I follows law of mass action [10]. Our paper is organized as follows: In the first section, formation of model, positivity and boundedness is discussed. Second section continues with local and global stability analysis of the boundary and planner equilibrium, in the next section existence of bionomic equilibrium and optimal policy is determined by using Pontryagin’s maximal principle. Numerical simulation is carried out in the final section followed by conclusion.

A. Formation of Model

In this section, phytoplankton-zooplankton interaction is modeled with the help of system of simultaneous differential equations with the following assumptions:

(i) The variable \( P(t) \) is the density of the phytoplankton population and \( Z \) is the density of zooplankton population at any instant of time \( t \) subject to the non-negative initial condition \( P(0) = P_0 > 0 \) and \( Z(0) = Z_0 > 0 \).

(ii) The parameter \( r_1 \), \( r_2 \) are the intrinsic growth rates and \( K_1 \), \( K_2 \) are the environmental carrying capacities of phytoplankton and zooplankton population respectively.

(iii) The constant \( \rho_1 > 0 \) is the maximum uptake rate for zooplankton species, \( \rho_2 > 0 \) denotes the ratio of biomass conversion (satisfying the obvious restriction \( 0 < \rho_2 < \rho_1 \)).

(iv) \( \mu_1 \), \( \mu_2 \) are the natural death rates of populations and \( \alpha \) denotes the rate of toxic substances produced by per unit biomass of phytoplankton.

(v) The term \( \alpha PZ \) describes the distribution of toxic substances produced by phytoplankton.
substances which ultimately contributes to the death of zooplankton populations where \( r_2 > \alpha \), i.e., the ratio of biomass consumed by zooplankton is greater than the rate of toxic substance liberation by phytoplankton species.

(vii) Both populations are subjected to constant harvesting with harvesting terms \( c_1EP, c_2EZ \), where \( c_1, c_2 \) are catchability coefficients and constant \( E \) is the harvesting effort.

Thus the phytoplankton-zooplankton interaction with above assumptions are represented by

\[
\begin{align*}
\frac{dP}{dt} &= r_1P(1 - \frac{P}{K_1}) - (\mu_1 + c_1E)P - \rho_1PZ \\
\frac{dZ}{dt} &= r_2Z(1 - \frac{Z}{K_2}) - (\mu_2 + c_2E)Z + (\rho_2 - \alpha)PZ
\end{align*}
\]

with the initial conditions \( P(0) = P_0 > 0 \), \( Z(0) = Z_0 > 0 \).

1) Positivity and Boundedness of Solution: In this section we discuss the positivity and boundedness of the system (1) under the given initial conditions for all \( t \geq 0 \). The system equations (1) yields

\[
P(t) = P(0) \exp \int_0^t (r_1(1 - \frac{P}{K_1}) - (\mu_1 + c_1E) - \rho_1Z)ds \geq 0 \quad \text{and} \quad Z(t) = Z(0) \exp \int_0^t (r_2(1 - \frac{Z}{K_2}) - (\mu_2 + c_2E) + (\rho_2 - \alpha)P)ds \geq 0
\]

Further, \( \frac{dP}{dt} \leq r_1P(1 - \frac{P}{K_1}) - c_1EP \) and \( \frac{dZ}{dt} \leq r_2Z(1 - \frac{Z}{K_2}) - c_2EZ \)

By using a standard comparison theorem [11]

\[
\limsup_{t \to +\infty} P(t) \leq \xi_1 \quad \text{and} \quad \limsup_{t \to +\infty} Z(t) \leq \xi_2
\]

where \( \xi_1 = \max\{P(0), \frac{K_1(1 - c_1E)}{r_1}\} \) and \( \xi_2 = \max\{Z(0), \frac{K_2(1 - c_2E)}{r_2}\} \).

Thus all solution curves of the system (1) with given initial conditions enter the region, \( \Gamma = \{(P(t), Z(t)) | 0 \leq P(t) \leq \xi_1, 0 \leq Z(t) \leq \xi_2\} \).

Remark: In biological context boundedness may be interpreted as natural restriction to growth of species due to limited resources.

2) Stability of equilibria: The possible steady states of the system (1) are

(i) \( R_0 = (0, 0) \), the extinction equilibrium which always exist

(ii) \( R_1 = \left(\frac{(1 - c_1E)r_1}{\mu_1}, 0\right) \), zooplankton free equilibrium exist if \( E < \frac{r_1c_1E}{\mu_1} \).

(iii) \( R_2 = \left(0, \frac{(1 - c_2E)r_2}{\mu_2}\right) \), zooplankton dominance equilibria exist if \( E < \frac{r_2c_2E}{\mu_2} \), and

(iv) The interior equilibrium \( R^* = (P^*, Z^*) \)

\[
P^* = \frac{K_1(1 - c_1E)}{r_1} \quad \text{and} \quad Z^* = \frac{K_2(1 - c_2E)}{r_2}
\]

where \( P^* = \left(\frac{(r_1 - \mu_1 - c_1E)r_1}{\rho_1} - \rho_1Z^*\right) \), and \( Z^* = \left(\frac{(r_2 - \mu_2 - c_2E)r_2}{\rho_2} - \rho_2\right) \)

The dynamical behaviour of the system around various equilibria is determined by the nature of the eigen values of community matrix,

\[
J = \begin{bmatrix} r_1 - \frac{2\alpha c_1E}{K_1} - \mu_1 - c_1E & -\rho_1 P^* - c_1 E \\ (\rho_2 - \alpha)Z^* & r_2 - \frac{2\alpha c_2E}{K_2} + (\rho_2 - \alpha)P^* - \mu_2 - c_2E \end{bmatrix}
\]

At \( R_0 \), the characteristic equation is

\[
J(R_0) = \begin{vmatrix} r_1 - \mu_1 - c_1E - \lambda & 0 \\ 0 & r_2 - \mu_2 - c_2E - \lambda \end{vmatrix} = 0.
\]

The roots of this equation are \( \lambda_1 = r_1 - \mu_1 - c_1E \) and \( \lambda_2 = r_2 - \mu_2 - c_2E \).

Thus \( R_0 \) is a stable node if \( r_1 - \mu_1 - c_1E < 0 \) and \( r_2 - \mu_2 - c_2E < 0 \), i.e., \( E > \max\left[\frac{\mu_1}{1 - c_1E}, \frac{\mu_2}{1 - c_2E}\right] \).

Remark: The biological significance of the above theorem is that, if \( E \) crosses a threshold level, the extinction equilibria \( R_0 \) becomes stable which further reaffirms the ecologically well known fact that over exploitation would result in population extinction.

Proposition 1. For the system (1), \( R_0 = (0, 0) \) always exist and when \( E > \max\left[\frac{\mu_1}{1 - c_1E}, \frac{\mu_2}{1 - c_2E}\right] \), it is asymptotically stable.

Proposition 2. For the system (1), if \( r_1c_1E < r_2c_2E < r_1(\mu_1 - c_1E)K_1(1 - c_1E) \), then \( R_0, R_1 \) and \( R_2 \) exist, and \( R_0, R_1 \) become unstable, \( R_2 \) is locally asymptotically stable if \( E > \frac{r_2(1 - c_2E)r_2}{\mu_2} \).

Next, if the interior equilibrium \( R^* \) exists, then the characteristic equation at \( R^* \) is,

\[
\lambda^2 - \text{Trace} J\lambda + \text{det} J = 0
\]

where, \( \text{Trace} J = (r_1 - \mu_1 - c_1E) - \frac{2\alpha c_1E}{K_1} + (\rho_2 - \alpha)P^* \) and \( \text{det} J = [(r_1 - \mu_1 - c_1E) - \frac{2\alpha c_1E}{K_1} - \rho_1(\rho_2 - \alpha)P^* + \rho_1(\rho_2 - \alpha)P^*Z^*] [r_2 - \mu_2 - c_2E - \frac{2\alpha c_2E}{K_2} + (\rho_2 - \alpha)P^* + \rho_1(\rho_2 - \alpha)P^*Z^*].

Theorem 1. For the system (1), if \( \frac{r_2}{\mu_2} < E < \frac{r_1}{\mu_1} \), then \( R^* \) exists and is locally asymptotically stable if \( \text{Trace} J < 0 \) and \( \text{det} J > 0 \).

For the global stability of the equilibrium, we have the following theorem’s.

Theorem 2. The extinction equilibrium \( R_0 \) is globally asymptotically stable (GAS) if \( E > \max\left[\frac{\mu_1}{1 - c_1E}, \frac{\mu_2}{1 - c_2E}\right] \).

Proof: Consider the following Lyapunov function
\[ V(P, Z) = \frac{1}{m} P(t) + \frac{1}{p_2 - \alpha} Z(t) \]

Its time derivative along the solutions of system (1) yields

\[ \frac{dV}{dt} = (r_1 - \mu_1 - c_1 E) P + \frac{r_1 P^2}{\rho_1 K_1} + \frac{r_2 Z}{\rho_2 - \alpha} \frac{r_2 Z^2}{(p_2 - \alpha) K_2} \]

If \( r_1 - \mu_1 - c_1 E \leq 0 \) and \( r_2 - \mu_2 - c_2 E < 0 \), then we obtain \( \frac{dV}{dt} < 0 \).

Further, the Lyapunov theorem on stability [12] implies that all solutions ultimately approach the equilibrium \( R_0 \). This establishes our global result of the equilibrium \( R_0 \).

**Theorem 3.** The interior equilibrium \( R^* \) is globally asymptotically stable (GAS) in the positive quadrant of \( \mathbb{R}^2 \).

**Proof of theorem:** Let us define a lyaponuv function

\[ V(P, Z) = \int_{P_0}^{P} \frac{1}{x} - \frac{1}{P^*} dx + \frac{\rho_1}{(p_2 - \alpha)} \int_{Z_0}^{Z} \frac{1}{x} - \frac{1}{Z^*} dx \]

Then \( V(P, Z) = 0 \) if and only if \( P = P^* \) and \( Z = Z^* \) and \( V(P, Z) \geq 0 \) in \( p_2 \)-plane.

The time derivative of \( V \) along the trajectories of system is

\[ \frac{dV}{dt} = \frac{P - P^*}{P^*} \frac{dP}{dt} + \frac{\rho_1}{(p_2 - \alpha)} \frac{Z - Z^*}{Z^*} \frac{dZ}{dt} \]

\[ = (P - P^*) [r_1 \frac{1}{K_1} - (\mu_1 + c_1 E) - \mu_1 Z^*] \]

\[ + \frac{\rho_1}{(p_2 - \alpha)} (Z - Z^*) [r_2 \frac{1}{K_2} - (\mu_2 + c_2 E) + \mu_2 Z^*] \]

After some algebraic calculations, we can obtain

\[ \frac{dV}{dt} = \frac{r_1 (P - P^*)^2}{K_1} - \frac{\rho_1}{(p_2 - \alpha)} \frac{r_2 (Z - Z^*)^2}{K_2} < 0 \]

Thus \( \frac{dV}{dt} \leq 0 \) and \( \frac{dV}{dt} = 0 \) iff \( P = P^* \) and \( Z = Z^* \). Thus by lasalle’s theorem [12], \( R^* \) is globally asymptotically stable (GAS) in some neighborhood of \( p_2 \)-plane.

**B. Non Existence of Periodic Solutions**

**Theorem 4.** System (1) does not have any limit cycle in the positive quadrant of \( p_2 \)-plane.

**Proof:** For the proof of the above theorem, consider a continuous and differentiable function \( D(P, Z) = \frac{1}{p_2 - \alpha} \) in the simple connected domain \( \Omega \) of the region \( \Gamma \) in the positive quadrant of \( p_2 \)-plane.

Let \( H(P, Z) = r_1 P(1 - \frac{P}{K_1}) - (\mu_1 + c_1 E) P - \rho_1 P Z \)

and \( G(P, Z) = r_2 Z(1 - \frac{Z}{K_2}) - (\mu_2 + c_2 E) Z + (p_2 - \alpha) P Z \)

Then

\[ \Delta(P(t), Z(t)) = \frac{\partial(DH)}{\partial P} + \frac{\partial(DG)}{\partial Z} \]

\[ = -\frac{r_1}{K_1 Z} - \frac{r_2}{K_2 P} < 0 \]

which is negative. Thus \( \Delta(P(t), Z(t)) \) neither change sign nor identically zero in the positive quadrant of \( p_2 \)-plane. Therefore Bendixon-Dulac criteria confirm the non-existence of any limit cycles or closed trajectory in the positive quadrant of \( p_2 \)-plane.

**C. Bionomic Equilibrium**

The bionomic equilibrium is said to be achieved when the total revenue obtained by selling the harvested biomass equals the total cost of harvesting it. Let \( C \) be the harvesting cost per unit effort; \( p_1, p_2 \) are the prices per unit biomass of the phytoplankton and zooplankton respectively. Then net economic revenue or economic rent at any time \( t \) is given by, \( \pi(P, Z, E, t) = (p_1 c_1 P + p_2 c_2 Z - C) E \).

The bionomic equilibrium \( (P_\infty, Z_\infty, E_\infty) \), of the phytoplankton-zooplankton system is the solution of the biological equilibrium given by \( \frac{dP}{dt} = 0 \), \( \frac{dZ}{dt} = 0 \), which yields

\[ E = \frac{r_1}{c_1} (1 - \frac{P}{K_1}) - \mu_1 - \mu_1 Z \]

\[ = \frac{r_2}{c_2} (1 - \frac{Z}{K_2}) - (\mu_2 + c_2 E) \quad (3) \]

or

\[ \frac{(r_1 + \frac{c_1 (p_2 - \alpha)}{c_2}) P}{c_2} + (p_1 - \frac{c_1 (r_2)}{c_2 K_2}) Z \]

\[ + (c_1 (r_2 - \mu_2) - (r_2 - \mu_1)) = 0 \quad (4) \]

and the economic equilibrium which is said to be achieved when the economic rent is completely dissipated, i.e.

\[ \pi(P, Z, E, t) = (p_1 c_1 P + p_2 c_2 Z - C) E = 0 \quad (5) \]

On solving (4), (5) and using (3), we can find the bionomic equilibrium

\[ P_\infty = \frac{(r_1 - \mu_1) c_2 - c_1 (r_2 - \mu_2) + \frac{C (c_1 r_2 - \rho_1 c_2 K_2)}{c_2 p_2 K_2}}{r_1 c_2 + c_1 (p_2 - \alpha)} + \frac{c_2 p_2 K_2}{c_2 p_2 K_2} \]

which is positive if \( \frac{c_1}{r_2} < \min (\frac{r_1 - \mu_1}{r_2 - \mu_2}, \frac{p_1 K_2}{r_2}) \),

\[ Z_\infty = \frac{C - c_1 P_\infty}{c_1 p_1} \quad \text{exist if} \quad P_\infty < \frac{C}{c_1 p_1} \]

and \( E_\infty = \frac{C}{c_1} [(1 - \frac{r_1}{K_1}) - \mu_1 - \mu_1 Z_\infty] \quad \text{exist if} \quad P_\infty < \frac{C}{c_1 p_1} \).
D. Optimal Harvesting Policy

In this section, the aim is to find an optimal harvesting policy for maximum sustainable yield by assurance to regulatory agency to achieve its objective. We consider the present value \( R \) of a continuous time-stream of revenues given by:

\[
R = \int_0^\infty e^{-\delta t} \left( p_1 \lambda_1 P(t) + p_2 \mathcal{C}(t) - C \right) E(t) dt
\]

where \( \delta \) is the instantaneous rate of annual discount. Thus, our objective is to maximize \( R \) subject to (1) and to the control constraints 0 \( \leq \) \( E \) \( \leq \) \( E_{\text{max}} \), here \( E_{\text{max}} \) is the upper limit for the harvesting effort.

By using the Pontryagin Maximum Principle [13], the associated hamiltonian function is given by:

\[
H = e^{-\delta t} (p_1 \lambda_1 P + p_2 \mathcal{C}(Z - C)) E + \lambda_1 (r_1 P(1 - \frac{P}{K_1}) - (\mu_1 + c_1 E)P - \mu_1 P - \rho_1 PZ) - \mu_2 \mathcal{C}(Z - \frac{Z}{K_2}) E + \lambda_2 (r_2 (1 - \frac{P}{K_1}) - (\mu_1 + c_1 E)P - \mu_1 P - \rho_1 PZ) + \lambda_2 (\rho_2 - \alpha PZ)
\]

where \( \lambda_1, \lambda_2 \) are the adjoint operators which satisfy the equations

\[
\frac{d\lambda_1}{dt} = -\frac{\partial H}{\partial E}, \quad (6)
\]

\[
\frac{d\lambda_2}{dt} = -\frac{\partial H}{\partial Z}, \quad (7)
\]

and \( \sigma(t) = (e^{-\delta t} (p_1 \lambda_1 P + p_2 \mathcal{C}(Z - C)) - \lambda_1 \lambda_1 P - \lambda_2 \mathcal{C}(Z - C)) \]

represents the switching function [15]. Since the Hamiltonian \( H \) is linear in control variable \( E(t) \) so in this case only singular control for optimisation problem will be obtained. Thus the necessary condition to maximize the hamiltonian \( H \) under the singular control variable \( E(t) \) is

\[
\frac{\partial H}{\partial E} = 0
\]

So (8) gives:

\[
e^{-\delta t} (p_1 \lambda_1 P + p_2 \mathcal{C}(Z - C)) - \lambda_1 \lambda_1 P - \lambda_2 \mathcal{C}(Z - C) = 0. \quad (9)
\]

or it can be written as \( (\lambda_1 \lambda_1 P + \lambda_2 \mathcal{C}(Z - C)) = e^{-\delta t} \frac{\partial \pi}{\partial \mathcal{E}} \), which implies that the total user’s cost of harvesting per unit effort is equal to the discounted values of the future price at the steady state effort level.

Now from (6) and (7), we have

\[
\frac{d\lambda_1}{dt} = -c_1 p_1 e^{-\delta t} - \lambda_1 (r_1 - \frac{2r_1 P}{K_1} - (\mu_1 + c_1 E) - \rho_1 Z) - (\rho_2 - \alpha) Z \lambda_2
\]

\[
\frac{d\lambda_2}{dt} = -c_2 p_2 e^{-\delta t} + \rho_1 \lambda_1 - \lambda_2 (r_2 - \frac{2r_2 Z}{K_2}) - (\mu_2 + c_2 E) + (\rho_2 - \alpha) P
\]

Now in order to find the optimal equilibrium we consider the equilibrium value of \( E \) from (3) at \( R^* \) as

\[
E = \frac{r_1}{c_1 \left(1 - \frac{P^*}{K_1} \right) - \mu_1 - \rho_1 Z^*} - \frac{r_2}{c_2 \left(1 - \frac{Z^*}{K_2} \right) - \mu_2 + (\rho_2 - \alpha) P^*}
\]

and using it in (10) and (11), the following system of simultaneous linear equations can be obtained,

\[
\frac{d\lambda_1}{dt} = -c_1 p_1 e^{-\delta t} + \frac{r_1 P \lambda_1}{K_1} - (\rho_2 - \alpha) Z \lambda_2
\]

\[
\frac{d\lambda_2}{dt} = -c_2 p_2 e^{-\delta t} + \rho_1 \lambda_1 + \frac{r_2 Z \lambda_2}{K_2}
\]

Eliminating \( \lambda_2 \) from (13) and (14), we have

\[
\frac{d\lambda_1}{dt} - M_1 \lambda_1 = -M_2 e^{-\delta t},
\]

where, \( M_1 = \frac{r_1}{K_1} + \frac{(\rho_2 - \alpha) c_1}{c_2} P^* \) and \( M_2 = c_1 r_1 E + \frac{(\rho_2 - \alpha) c_2}{c_2} (c_1 P^* + c_2 P^* Z - C) \)

On solving (15), we get \( \lambda_1 = \frac{M_2}{M_1 + \delta} e^{-\delta t} \) and using this value in (14), we obtain

\[
\frac{d\lambda_2}{dt} - N_1 \lambda_1 = -N_2 e^{-\delta t}
\]

which results into \( \lambda_2 = \frac{N_2}{N_1 + \delta} e^{-\delta t} \)

where \( N_1 = \frac{r_2 Z^*}{K_2} \) and \( N_2 = c_2 P^* - \mu_1 \frac{M_2}{M_1 + \delta} P^* \)

Here \( \lambda_2 e^{\delta t} \) represents shadow prices [16] along the singular path and from the solution of (15) and (16), it is clear that shadow prices remains constant over time interval in optimal equilibrium when they strictly satisfy the transversality condition at \( t \to \infty \) [14-15]. It implies they remain bounded as \( t \to \infty \). Using the values of \( \lambda_1 \) and \( \lambda_2 \) in (9), the equation of the path of singular control is:

\[
c_1 (p_1 - \frac{M_2}{M_1 + \delta}) P + c_2 (p_2 - \frac{N_2}{N_1 + \delta}) Z = C
\]

Considering the values of \( M_1, M_2, N_1 \) and \( N_2 \) at the equilibrium value of \( E \); (5) and (17) gives the optimal equilibrium of the population of phytoplankton and zooplankton i.e.(\( P_1, Z_1 \)). Moreover from (3), the corresponding optimal harvesting effort \( E_E \) can be estimated. Now from (17), it can be concluded that

\[
\pi(P, Z, E) = c_1 p_1 P + c_2 p_2 Z - C = \frac{c_1 M_2}{M_1 + \delta} P + \frac{c_2 N_2}{N_1 + \delta} Z \to 0
\]

\[
as \delta \to \infty
\]

Therefore, the net economic revenue \( \pi(P_\infty, Z_\infty, E, t) = 0 \). This implies that in case of infinite discount rate, the net economic revenue becomes zero and harvesting will no longer take place [16]. Moreover (18) shows that, the net economic rent only be maximised in the optimal equilibrium if zero discount rate being offered.
Again equation (18) shows that, \( c_1 M_2 (N_1 + \delta) P + c_2 N_2 (M_1 + \delta) \) is of \( O(\delta) \) and \( (M_1 + \delta) (N_1 + \delta) Z \) of \( O(\delta^2) \), thus \( \pi \) is of \( O(\delta^{-1}) \) and this implies, \( \pi \) (the economic rent function) is a decreasing function of the instantaneous annual rate of discount \( \delta \). Hence, \( \delta = 0 \) leads to maximization of \( \pi \).

E. Numerical Simulation

(i) Consider the following set of parametric values of the given system,
\[
\begin{align*}
r_1 &= 6, \quad r_2 = 0.1, \quad K_1 = 50, \quad K_2 = 30, \quad \rho_1 = 0.3, \quad \rho_2 = 0.25, \\
\alpha &= 0.04, \quad \mu_1 = 0.05, \quad \mu_2 = 0.08, \quad c_1 = 0.22, \quad c_2 = 0.1, \\
E &= 0.5.
\end{align*}
\]
It is easy to find \( \mu_{\text{max}} = 27.0455, \quad \mu_{\text{min}} = 0.2, \quad Z_{\text{max}} = 26.8005, \quad \text{Trace} \, J = -0.1182 < 0 \) and \( \text{Det} \, J > 0.5490 > 0 \).
So, the condition of Theorem 1 holds, then the interior equilibrium \( R_* = (0.4490, 19.2871) \) is globally asymptotically stable, which is shown in fig. 1. In the absence of harvesting effort \( (E=0) \), the globally asymptotically stable equilibrium \( R_* \) exists at \((0.2184, 19.7419)\) (see fig. 2).

(ii) Choosing \( E = 30 \), keeping the other parametric values as in (i)
It is easy to verify that \( E > \max \left( \frac{r_1 - \mu_1}{c_1}, \frac{r_2 - \mu_2}{c_2} \right) = \max(27.0455, 0.2) = 27.0455 \).
Then the conditions of Theorem 2 are satisfied. Hence the extinction equilibrium \( R_0 \) is globally asymptotically stable, which is shown in fig. 3.

(iii) Taking the same set of parametric values in (i) with \( p_1 = 2, \quad p_2 = 3, \quad C = 5.9, \quad \delta = 0.03 \) and
\[
E = \frac{1}{\mu_2} \left[ 1 - \frac{c_2}{\mu_2} \right] - \mu_1 - \rho_1 Z^* \]
instead taking \( E = 0.5 \) or 1, we obtain the bionomic equilibrium is \( (P_\infty, Z_\infty) = (0.9834, 18.2243) \) and effort value to reach this equilibrium is \( E_\infty = 1.6577 \). The optimal equilibrium solution is \( (P_0, Z_0) = (0.5137, 18.9132) \) and the effort value to reach this equilibrium is \( E_0 = 0.9745 \).

II. Conclusion

In this paper, a mathematical model of phytoplankton-zooplankton interactions with harvesting (i.e. some species are exploited commercially for food supply) is analysed. It is assumed that, both populations grow logistically and some species of phytoplankton releases toxic substances which reduces the predator’s grazing pressure on their prey (phytoplankton) species. Using stability theory of ordinary differential equations the dynamical properties (local and global) of the system along with the effects of harvesting efforts are discussed. It is shown, in case of excessive harvesting the system never recovered and entire populations goes to extinction. Further, it has been proved that the interior equilibrium exists under certain conditions and it is globally asymptotically stable. Moreover, numerical results shows that, in the absence of harvesting \( (E=0) \) interior equilibrium exists at lower population level \( R_* = (0.0575, 19.8102) \) compared to \( R_0 = (0.2815, 19.3555) \) (in the presence of harvesting) for the phytoplankton but at a higher population level for the zooplankton. It has also been shown that the system under consideration does not have any limit cycle by using dulac’s criterion. Next, the existence of bionomic equilibria (intersection of the zero profit line and the biological equilibriums) and optimal harvesting policy are discussed. The present value of revenues is maximized by using Pontryagin’s maximum principle subject to the state equations and the control constraints. It is found that the shadow prices remain constant over time in optimal equilibrium when they satisfy the transversality condition. It is established that the zero discounting leads to the maximization of economic revenue and that an infinite discount rate leads to complete dissipation of economic rent.

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