Simulation of Optimal Harvesting of Three Species Ecological Model with Closed Interval of Biological Parameter using by using First Order Nonlinear Ordinary Differential Equations

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Abstract
The paper presents the study of three species ecological model with Prey N 1, predator N 2 and competitor to the Predator N 3 and neutral with the predator N 2 with imprecise biological parameters. The model is characterized by a set of first order nonlinear ordinary differential equations. Due to the lack of precise numerical information of the biological parameters such as prey population growth rate, predator population decay rate and predation coefficients, we consider the model with imprecise data as form of an interval in nature. Many authors have studied prey–predator harvesting model in different form, here we consider a simple prey–predator model under impreciseness and introduce parametric functional form of an interval and then study the model. Equilibrium points of the model are identified, the local stability is discussed using Routh - Hurwitz criteria and global stability by Liapunov function. The existence of bionomic equilibrium of the system has been discussed and optimal harvesting policy is given using Pontryagin’s maximum principle. The stability analysis is supported by Numerical simulation using Mat lab.

Keywords: Prey; Predator; Competitor to the predator; Equilibrium points; interval number, Stability of the equilibrium points; Bionomic Equilibrium; Optimal harvesting policy; Pontryagin’s maximum principle; Numerical simulation using mat lab.

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INTRODUCTION

1. Background
Mathematical modeling of ecosystems is a field of study which helps us to understand the interactions between different species and the mechanisms that influence the growth of species and their existence and stability. Mathematical models have been used to study the dynamics of prey-predator systems since Lotka (1925) and Volterra (1927). They proposed the simple mathematical model which describes the interaction between prey and the predator. Since then, many mathematical models have been constructed based on more realistic explicit and implicit biological assumptions.

Mathematical modeling is a frequently evolving process, to gain a deep understanding of the mathematical aspects of the problem and to yield non trivial biological insights; one must carefully construct biologically meaningful and mathematically tractable population models. Some of the aspects that need to be critically considered in a realistic and plausible mathematical model include; carrying capacity which is the maximum number of prey that the ecosystem can sustain in the absence of predator, competition among prey and predators which can be intraspecific or inter specific, harvesting of prey or predators and functional responses of the predators.

In this research work, a mathematical model to study the ecological dynamics of prey and predator system is proposed and analyzed. And also as an example some of the prey and predator system in some areas be studied.

1.1 Pre-Requisite Mathematics

1.1.1 Interval Number
Interval arithmetic, interval mathematics, interval analysis, or interval computation, is a method developed by mathematicians since the 1950s and 1960s as an approach to putting bounds on rounding errors and measurement errors in mathematical computation and thus developing numerical methods that yield reliable results. Very simply, it represents each value as a range of possibilities. For example, instead of estimating the height of someone using standard arithmetic as 2.0 meters, using interval arithmetic we might be certain that person is somewhere between 1.97 and 2.03 meters.

An interval number $A$ is represented by closed interval $[a_l, a_r]$ and defined by:

$$ A = [a_l, a_r] = \{x: a_l \leq x \leq a_r, x \in \mathbb{R}\} $$

Where $R$ is the set of real numbers and $a_l, a_r$ are the left and right limit of the interval number respectively. Also every real number $a$ can be represented by the interval number as $[a, a]$ for all
$a \in \mathbb{R}$

Classical arithmetic defines operations on individual numbers; interval arithmetic defines a set of operations on intervals.

$T \cdot S = \{ x \mid \text{there is some } y \text{ in } T, \text{ and some } z \text{ in } S, \text{ such that } x = y \cdot z \}$. 

The basic operations of interval arithmetic are, for two intervals $[a, b]$ and $[c, d]$ that are subsets of the real line $(-\infty, \infty)$,

i. $[a, b] + [c, d] = [a + c, b + d]$

ii. $[a, b] - [c, d] = [a - c, b - d]$

iii. $[a, b] \times [c, d] = [\min(a \times c, a \times d, b \times c, b \times d), \max(a \times c, a \times d, b \times c, b \times d)]$

iv. $[a, b] + [c, d] = [\min(a + c, a + d, b + c, b + d), \max(a + c, a + d, b + c, b + d)]$, when 0 is not in $[c, d]$

Division by an interval containing zero is not defined under the basic interval arithmetic. Instead of working with an uncertain real $x$ we work with the two ends of the interval $[a, b]$ which contains $x$ such that $x$ lies between $a$ and $b$, or could be one of them. Similarly a function $f$ when applied to $x$ is also uncertain. Instead, in interval arithmetic $f$ produces an interval $[a, b]$ which is all the possible values for $f(x)$ for all $x \in [a, b]$.

### 1.1.2 Interval-Valued Function

Let $a > 0$, $b > 0$ and consider the interval $[a, b]$. From a mathematical point of view, any real number can be represented on a line. Similarly, we can represent an interval by a function. If the interval is of the form $[a, b]$, the interval-valued function is taken as $h(p) = a^{(1-p)}b^p$ for $p \in [0,1]$.

Here we present some arithmetic operations on interval valued functions as follows: Let $A = [a_l, a_u]$ and $B = [b_l, b_u]$ be two interval numbers.

**Addition:** $A + B = [a_l + b_l, a_u + b_u]$. Provided $A_l + B_l > 0$. The interval-valued function for the interval $A + B$ is given by $h(p) = a_l^{(1-p)}a_u^p$ where $A_l + B_l > 0$ and $A_u + B_u = a_u + b_u$.

**Subtraction:** $A - B = [a_l - b_u, a_u - b_l]$. Provided $A_l - B_l > 0$. The interval-valued function for the interval $A - B$ is given by $h(p) = b_l^{(1-p)}b_u^p$ where $b_l = a_l - b_l$ and $b_u = a_u - b_u$.

**Scalar multiplication:** $\alpha A = \alpha[a_l, a_u] = \{[(\alpha a_l, \alpha a_u)\} \text{ if } \alpha \geq 0 \text{ and } \{[(\alpha a_l, \alpha a_u)\} \text{ if } \alpha < 0 \text{ provided } A_l > 0 \text{ and } A_u > 0$. The interval valued function for interval $\alpha A$ is given by $h(p) = c_l^{(1-p)}c_u^p$ if $\alpha \geq 0$ and $h(p) = -d_l^{(1-p)}d_u^p$ if $\alpha < 0$ where $C_l = \alpha a_l, C_u = \alpha a_u, d_l = |a|a_l$ and $d_u = |a|a_u$.

### 1.1.3 Pontryagin’s Maximum Principle

Pontryagin’s maximum principle is a powerful method for the computation of optimal controls, which has the crucial advantage that it does not require prior evaluation of the informal cost function. Let $b, c$ and $C$ are differentiable function in $t$ and $x$ with continuous derivatives, and that the stopping set D is a hyperplane, thus

$D = \{y + \sum \Sigma \in \mathbb{R}^n \text{ for some } y \in \mathbb{R}^n \text{ and some vector subspace } \Sigma \text{ of } \mathbb{R}^n \}$

Define for $\lambda \in \mathbb{R}^n$ the Hamiltonian function as:

$H(t, x, u, \lambda) = \lambda^T b(t, x, u) - c(t, x, u)$

Pontryagin’s maximum principle states that if $(x_t, u_t) \leq \tau$ is optimal, then there exist adjoint paths $\left(\lambda_t^\tau, \mu_t^\tau\right) \leq \tau$ in $\mathbb{R}^n$ and $\mathbb{R}$ with the following properties for all $t \leq \tau$

$H(t, x_t, u_t, \lambda_t^\tau, \mu_t^\tau)$ has maximum value 0, at the point $u_t = u_t$

$\dot{\lambda}_t^\tau = -\lambda_t^\tau \nabla b(t, x_t, u_t) - \nabla c(t, x_t, u_t)$

$\dot{\mu}_t^\tau = \lambda_t^\tau \nabla b(t, x_t, u_t) - c(t, x_t, u_t)$

Moreover the following transversality conditions hold: $(x_t, u_t) \leq \tau$

$\left(\lambda_{\tau}^\tau + \nabla C(t, x_t)\right)\sigma = 0$ for all $\sigma \in \Sigma$ and, in the time-unconstrained case
\[ ii) \quad \mu_t + \dot{C}(r, X_r) = 0 \]

Note that, in the time-unconstrained case, if \(b, c\) and \(C\) are time-independent functions, then \(\mu_t = 0\) for all \(t\).

The Hamiltonian serves as a way of remembering the first four statements, which could be expressed alternatively as:

i. \[ \frac{\partial H}{\partial u} = 0 \]

ii. \[ -\frac{\partial H}{\partial x} = \dot{\lambda} \]

iii. \[ -\frac{\partial H}{\partial t} = \dot{\mu} \]

iv. \[ \frac{\partial H}{\partial \lambda} = \dot{x} \]

The condition \(\frac{\partial H}{\partial u} = 0\) is not always correct. For example in cases where the set of actions is an interval and where the maximum is achieved at an endpoint.

2. MODEL FORMULATION AND ANALYSIS

In section deals with the mathematical modeling of the prey-predator dynamics where there are two predators which compete for the same limited resources. In addition, the section deals with the stability analysis of the equilibrium points and the numerical simulation of the model.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Parameter Definition</th>
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<tbody>
<tr>
<td>(R)</td>
<td>net economic rent</td>
</tr>
<tr>
<td>(E_i, i = 1,2,3)</td>
<td>harvesting efforts</td>
</tr>
<tr>
<td>(\delta)</td>
<td>instantaneous annual rate of discount</td>
</tr>
<tr>
<td>(q_i, i = 1,2,3)</td>
<td>catch ability coefficients</td>
</tr>
<tr>
<td>(c_i, i = 1,2,3)</td>
<td>harvesting cost per unit effort</td>
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Table 3.1 Definition of some parameters

2.1 Prey–Predator Model

The ecological model is as follows. There is one prey and two predators, where the two predators are compete with each other for the use of common recourse i.e. food. But the two predators cannot eat each other (one is not eaten by the other). By assuming that the predator and competitor to the predator have alternative food in addition to prey population (but the competitor to the predator can’t eat the prey population), then the model for one Prey and two Predator and harvesting on the both species is given by the following system of first order ordinary differential equations employing the following notation:

Let \(N_1\) denotes the size of the prey population, \(N_2\) denotes the size of the predator population and \(N_3\) denotes the size of the competitor to the predator population, lets assuming that there is demand for all species in the market so the harvesting of both species are carried out. Let prey, predator and competitor to the predator species are subjected to harvesting efforts (effort applied to the harvest the prey, predator and competitor species) \(E_1, E_2\) and \(E_3\) respectively. Then the dynamics of the prey-predator is described by:

\[
\frac{dN_1}{dt} = rN_1 - \alpha_1(N_2)^2 - \beta_1N_1N_2 - q_1E_1N_1 \quad (3.1)
\]

\[
\frac{dN_2}{dt} = sN_2 - \alpha_2(N_2)^2 + \beta_2N_1N_2 - \delta_1N_2N_3 - q_2E_2N_2 \quad (3.2)
\]

\[
\frac{dN_3}{dt} = \ell N_3 - \alpha_3(N_3)^2 - \beta_3N_2N_3 - q_3E_3N_3 \quad (3.3)
\]

where \(r, s\) and \(\ell\) are natural growth rate of prey, predator and competitor to the predator species respectively. Whereas:

- \(\alpha_1\) is rate of decrease of the prey population due to inter species competition
- \(\beta_1\) is rate of decrease of the prey population due to inhibition by the predator population,
- \(\alpha_2\) is rate of decrease of the predator population due to inter species competition,
- \(\beta_2\) is rate of increase of the predator population due to successful attacks on the prey population,
- \(\delta_1\) is rate of decrease of the predator population due to the competition with the third species (competitor),
- \(\alpha_3\) is rate of decrease of the competitor population to the predator population due to inter species competition,
\( \beta_s \) is rate of decrease of the competitor population due to the competition with the third species (predator population). Where all the parameter values \( \alpha_1, \alpha_2, \alpha_3, \beta_1, \beta_2, \beta_3 \) and \( \delta_s \) are positive in nature and are considered precise. Intuitively if any of the parameters are imprecise, furthermore when any parameter of the right hand side of equations (3.1) - (3.3) are interval number rather than a single value, then it is not so straightforward to convert equations to the standard form like (3.1), (3.2) and (3.3). For an imprecise Coefficient we present the problem with an interval coefficient.

2.2.2 Prey – Predator Model with Interval Coefficient

Let \( \hat{r}, \hat{s}, \hat{\ell}, \hat{\alpha}_1, \hat{\alpha}_2, \hat{\alpha}_3, \hat{\beta}_1, \hat{\beta}_2, \hat{\beta}_3 \) and \( \hat{\delta}_s \) be the interval counterparts of \( r, s, \ell, \alpha_1, \alpha_2, \alpha_3, \beta_1, \beta_2, \beta_3 \) and \( \delta_s \) respectively, then the Prey – Predator model with combined harvesting \( E_1, E_2 \) and \( E_3 \) can be written in the following form:

\[
\frac{dN_1}{dt} = \hat{r}N_1 - \hat{\alpha}_1(N_1)^2 - \hat{\beta}_1N_1N_2 - q_1E_1N_1 - q_1E_1N_1
\]

\[
\frac{dN_2}{dt} = \hat{S}N_2 - \hat{\alpha}_2(N_2)^2 + \hat{\beta}_2N_1N_2 - \hat{\delta}_sN_2N_3 - q_2E_2N_2
\]

\[
\frac{dN_3}{dt} = \hat{\ell}N_3 - \hat{\alpha}_3(N_3)^2 - \hat{\beta}_3N_2N_3 - q_3E_3N_3
\]

Where \( \hat{r} \in [r_1, r_2], \hat{s} \in [s_1, s_2], \hat{\ell} \in [\ell_1, \ell_2], \hat{\alpha}_1 \in [(\alpha_1)_1, (\alpha_1)_2], \hat{\alpha}_2 \in [(\alpha_2)_1, (\alpha_2)_2], \hat{\alpha}_3 \in [(\alpha_3)_1, (\alpha_3)_2], \hat{\beta}_1 \in [(\beta_1)_1, (\beta_1)_2], \hat{\beta}_2 \in [(\beta_2)_1, (\beta_2)_2], \hat{\beta}_3 \in [(\beta_3)_1, (\beta_3)_2] \) and \( \hat{\delta}_s \in [(\delta_s)_1, (\delta_s)_2] \) With \( r_1, s_1, \ell_1, (\alpha_1)_1, (\alpha_2)_1, (\alpha_3)_1, (\beta_1)_1, (\beta_2)_1, (\beta_3)_1 \) and \( (\delta_s)_1 \) are all positive.

2.2.2 Prey – Predator Model with Parametric Interval Valued Function Coefficient

The parametric form of the equations (3.4), (3.5) and (3.6) are:

\[
\frac{dN_1(t; p)}{dt} = (r_1)^{1-p}(r_2)^{p}N_1 - ((\alpha_1)_u)^{1-p}(\alpha_1)_uN_1 + ((\beta_1)_u)^{1-p}(\beta_1)_uN_1 - q_1E_1N_1
\]

\[
\frac{dN_2(t; p)}{dt} = (s_1)^{1-p}(s_2)^{p}N_2 - ((\alpha_2)_u)^{1-p}(\alpha_2)_uN_2 + ((\beta_2)_u)^{1-p}(\beta_2)_uN_2 - q_2E_2N_2
\]

\[
\frac{dN_3(t; p)}{dt} = (\ell_1)^{1-p}(\ell_2)^{p}N_3 - ((\alpha_3)_u)^{1-p}(\alpha_3)_uN_3 + ((\beta_3)_u)^{1-p}(\beta_3)_uN_3 - q_3E_3N_3
\]

Theorem 1: The differential equations with interval valued coefficient

\[
\frac{dN_1}{dt} = \hat{r}_0N_1 - \hat{\alpha}_0(N_1)^2 - \hat{\beta}_0N_1N_2 - q_1E_1N_1
\]

\[
\frac{dN_2}{dt} = \hat{s}_0N_2 - \hat{\alpha}_0(N_2)^2 - \hat{\beta}_0N_1N_2 - q_2E_2N_2
\]

\[
\frac{dN_3}{dt} = \hat{\ell}_0N_3 - \hat{\beta}(N_3)^2 - \hat{\beta}_0N_2N_3 - q_3E_3N_3
\]

Where \( \hat{r}_0 \in [r_1, r_2], \hat{s}_0 \in [s_1, s_2], \hat{\ell}_0 \in [\ell_1, \ell_2], \hat{\alpha}_0 \in [\alpha_1, \alpha_2], \hat{\beta}_0 \in [\beta_1, \beta_2], \hat{\beta}_0 \in [\beta_1, \beta_2], \hat{\alpha}_0 \in [\alpha_1, \alpha_2], \hat{\beta}_0 \in [\beta_1, \beta_2], \hat{\beta}_0 \in [\beta_1, \beta_2] \) and \( \hat{\beta}_0 \in [\beta_1, \beta_2] \) are based on CPUE (catch-per unit-effort). But the competitor to the predator can’t change in the absence of the prey population. Where \( q_1, q_2, q_3 \) are the catch ability coefficients of three species and strictly positive. The catch-rate function: \( q_1E_1N_1, q_2E_2N_2, q_3E_3N_3 \) are based on CPUE (catch-per unit-effort).

Proof: Replacing in place of \( \hat{r}_0, \hat{s}_0, \hat{\ell}_0, \hat{\alpha}_0, \hat{\beta}_0, \hat{\gamma}_0, \hat{\delta}_0, \hat{\rho}_0 \) and \( \hat{\gamma}_0 \) by \( [r_1, r_2], [s_1, s_2], [\ell_1, \ell_2], [\alpha_1, \alpha_2], [\beta_1, \beta_2], [\gamma_1, \gamma_2], [\delta_1, \delta_2], [\rho_1, \rho_2] \) and \( [\gamma_1, \gamma_2] \) respectively, then equation (3.10), (3.11) and (3.12) will be come:

\[
\frac{dN_1}{dt} = [r_1, r_2]N_1 - [\alpha_1, \alpha_2](N_1)^2 - [\beta_1, \beta_2]N_1N_2 - q_1E_1N_1
\]
\[
\frac{dn_1}{dt} = [s_1, s_n]N_2 - [r_1, r_n](N_3)^2 + [\gamma_1, \gamma_n]N_1N_2 - [\delta_1, \delta_n]N_2N_3 - q_2E_2N_2 \tag{3.17}
\]
\[
\frac{dn_3}{dt} = [\ell_1, \ell_n]N_3 - [\rho_1, \rho_n](N_3)^2 - [\psi_1, \psi_n]N_2N_3 - q_3E_3N_3 \tag{3.18}
\]

Let \( r_1' \in [r_1, r_n], s_1' \in [s_1, s_n], \ell_1' \in [\ell_1, \ell_n], \alpha_1' \in [\alpha_1, \alpha_n], \gamma_1' \in [\gamma_1, \gamma_n], \delta_1' \in [\delta_1, \delta_n] \) and \( \psi_1' \in [\psi_1, \psi_n] \) respectively. Following the interval arithmetic operations and properties, equations (3.16), (3.17) and (3.18) reduces to:
\[
\frac{dn_1}{dt} = r_1'N_1 - \alpha_1'N_1^2 - \beta_1'N_1N_2 - q_1E_1N_1 \tag{3.19}
\]
\[
\frac{dn_2}{dt} = s_1'N_2 - \tau_1'N_2^2 + \gamma_1'N_1N_2 - \delta_1'N_2N_3 - q_2E_2N_2 \tag{3.20}
\]
\[
\frac{dn_3}{dt} = \ell_1'N_3 - \rho_1'N_3^2 - \psi_1'N_2N_3 - q_3E_3N_3 \tag{3.21}
\]

For fixed \( n \), let us consider the interval-valued function \( h_n^{(p)} = a_n^{(1-p)}b_n^{(p)} \) for \( p \in [0, 1] \) and interval \( a_n \in [a_n, b_n] \). Since \( h_n^{(p)} \) is a strictly increasing and continuous functions, then the above equation reduces to:
\[
\frac{dn_1}{dt} = r_1''N_1 - \alpha''(N_1)^2 - \beta''N_1N_2 - q_1E_1N_1 \tag{3.22}
\]
\[
\frac{dn_2}{dt} = s_1''N_2 - \tau''(N_2)^2 + \gamma''N_1N_2 - \delta''N_2N_3 - q_2E_2N_2 \tag{3.23}
\]
\[
\frac{dn_3}{dt} = \ell''N_3 - \rho''N_3^2 - \psi''N_2N_3 - q_3E_3N_3 \tag{3.24}
\]

Where
\[
r_1'' \in (r_1)^{1-p}(r_1)^p, \quad s_1'' \in (s_1)^{1-p}(s_1)^p, \quad \ell_1'' \in (\ell_1)^{1-p}(\ell_1)^p, \quad \alpha_1'' \in (\alpha_1)^{1-p}(\alpha_1)^p, \quad \gamma_1'' \in (\gamma_1)^{1-p}(\gamma_1)^p, \quad \delta_1'' \in (\delta_1)^{1-p}(\delta_1)^p, \quad \psi_1'' \in (\psi_1)^{1-p}(\psi_1)^p \text{ and } p \in [0, 1].
\]

Therefore the parametric form of the differential equations (3.10) - (3.12) is given by:
\[
\frac{dn_1}{dt} = (r_1)^{1-p}(r_1)^pN_1 - (\alpha_1)^{1-p}(\alpha_1)^pN_1^2 - (\beta_1)^{1-p}(\beta_1)^pN_1N_2 - q_1E_1N_1 \tag{3.25}
\]
\[
\frac{dn_2}{dt} = (s_1)^{1-p}(s_1)^pN_2 - (\tau_1)^{1-p}(\tau_1)^pN_2^2 + (\gamma_1)^{1-p}(\gamma_1)^pN_1N_2 - (\delta_1)^{1-p}(\delta_1)^pN_2N_3 - q_2E_2N_2 \tag{3.26}
\]
\[
\frac{dn_3}{dt} = (\ell_1)^{1-p}(\ell_1)^pN_3 - (\rho_1)^{1-p}(\rho_1)^pN_3^2 - (\psi_1)^{1-p}(\psi_1)^pN_2N_3 - q_3E_3N_3 \tag{3.27}
\]

2.3 Dynamic Behavior of the Harvesting Model

2.3.1 Equilibrium States of Prey-Predator Model with Parametric Interval Coefficient

The system under investigation has eight equilibrium states given by:
\[
\frac{dN_i(t,p)}{dt} = 0, \ i = 1, 2, 3 \tag{3.28}
\]

The possible equilibrium points of the systems are:

I. The extinct state:
\[
N_1 = 0, \ N_2 = 0 \text{ and } N_3 = 0 \tag{3.29}
\]

II. The state in which both the predator and competitor to the predator washed out and prey survive:
That is:
\[
N_1 = N_3 = 0 \text{ and } N_2 = \frac{a-q_1E_1}{b} \tag{3.30}
\]
\[
a = (r_1)^{1-p}(r_1)^p \text{ and } b = ((\alpha_1)^{1-p}((\alpha_1))^p
\]
\[
\frac{a-q_1E_1}{b}, 0, 0 \tag{3.31}
\]

III. The state in which only the predator survives and the prey and competitor to the predator are washed out:
That is:
\[
N_1 = N_3 = 0 \text{ and } N_2 = \frac{d-q_2E_2}{e} \tag{3.32}
\]
\[
d = (s_1)^{1-p}(s_1)^p \text{ and } e = ((\alpha_2)^{1-p}((\alpha_2))^p
\]
\[
(0, \frac{d-q_2E_2}{e}, 0) \tag{3.33}
\]

IV. The state in which both the prey and the predators washed out and competitor to the predator survive:
That is:
\[
N_1 = N_2 = 0 \text{ and } N_3 = \frac{h-q_3E_3}{k} \tag{3.34}
\]
\[
h = (\ell_1)^{1-p}(\ell_1)^p \text{ and } k = ((\rho_3)^{1-p}((\rho_3))^p
\]
\[
(0, 0, \frac{h-q_3E_3}{k}) \tag{3.35}
\]

V. The state in which both the prey and the predators stay alive and competitor to the predator vanishes:
That is:
\[
N_1 = 0 \tag{3.36}
\]
\[
(r_1)^{1-p}(r_1)^pN_1 - ((\alpha_1)^{1-p}((\alpha_1))^pN_1^2 - ((\beta_1)^{1-p}((\beta_1))^pN_1N_2 - q_1E_1N_1 = 0 \tag{3.37}
\]

And
By substituting \((3.33)\) in to equations \((3.1) - (3.3)\) and neglecting second and higher order terms of the Perturbations \(u_1, u_2\) and \(u_3\) we get the equations of the perturbed state:

\[
\frac{du}{dt} = A U
\]

where \(U = (u_1, u_2, u_3)\) and
The equilibrium state is stable, if three roots of the equation (3.36) have negative real parts. Solving for \( a, d \) and \( h \) from the equation (3.1) - (3.3) respectively and substituting in to the equation (3.35) we obtain the variational matrix:

\[
A = \begin{bmatrix}
-a - 2bN1 - cN2 - q1E1 & -cN1 & 0 \\
-2eN2 + fN1 - gN3 - q2E2 & -gN2 & 0 \\
0 & -mN3 & h - 2kN3 - mN2 - q3E3
\end{bmatrix}
\] (3.35)

The characteristic equation for the system is:

\[
d[A - \lambda I] = 0
\] (3.36)

The equilibrium state is stable, if three roots of the equation (3.36) have negative real parts.

### Local Stability Analysis

The local and global stability of the equilibrium states I, II, III and IV are found to be unstable. But the remaining equilibrium states V, VI, VII and VIII.

#### i. Stability of the Equilibrium State (\( \bar{N}_1, \bar{N}_2, 0 \))

The variational matrix at the trivial equilibrium point will become:

\[
A = \begin{bmatrix}
a - 2b\bar{N}1 - c\bar{N}2 - q1E1 & -c\bar{N}1 & 0 \\
f\bar{N}2 & d - 2e\bar{N}2 + f\bar{N}1 - q2E2 & -g\bar{N}2 \\
0 & 0 & h - m\bar{N}2 - q3E3
\end{bmatrix}
\] (3.38)

The Characteristic equation of the above variational matrix is given by:

\[
d[A - \lambda I] = 0
\]

One of the Eigen values of variational matrix \( A \) is \( \lambda = (h - q3E3) - m\bar{N}2 \) and the other two are obtained from the quadratic equation:

\[
\lambda^2 + (b\bar{N}1 + e\bar{N}2)\lambda + (be + cf)\bar{N}1\bar{N}2 = 0
\] (3.39)

In (3.39), the sum of the roots, \((-b\bar{N}1 + e\bar{N}2)\), is negative and the product of the roots, \((be + cf)\bar{N}1\bar{N}2\), is positive. Therefore the roots of (3.39) are real and negative or complex conjugates having negative real parts. Thus the equilibrium point is asymptotically stable when \((h - q3E3) < m\bar{N}2\).

#### ii. Stability of the Equilibrium State at (\( \bar{N}_1, 0, \bar{N}_3 \))

The variational matrix at the trivial equilibrium point (state) is:

\[
A = \begin{bmatrix}
a - 2b\bar{N}1 - c\bar{N}2 - q1E1 & -c\bar{N}1 & 0 \\
0 & d + f\bar{N}1 - g\bar{N}3 - q2E2 & 0 \\
0 & 0 & h - 2k\bar{N}3 - q3E3
\end{bmatrix}
\] (3.40)

The roots of the corresponding characteristic equations are:

\[
\lambda_1 = -(a - q1E1), \lambda_2 = d + \frac{f}{b}(a - q1E1) - \frac{g}{k}(h - q3E3) \text{ and } \lambda_3 = -(h - q3E3)
\]

The equilibrium point is asymptotically stable when \(d + \frac{f}{b}(a - q1E1) < \frac{g}{k}(h - q3E3)\).

#### iii. Stability of the equilibrium state at (0, \( \bar{N}_2, \bar{N}_3 \))

The variational matrix at the trivial equilibrium point (state) is:

\[
A = \begin{bmatrix}
a - c\bar{N}2 - q1E1 & 0 & 0 \\
f\bar{N}2 & d - 2e\bar{N}2 - g\bar{N}3 - q2E2 & -g\bar{N}2 \\
0 & -m\bar{N}3 & h - 2k\bar{N}3 - m\bar{N}2 - q3E3
\end{bmatrix}
\]

One of the Eigen values of variational matrix \( A \) is \( \lambda_1 = (a - q1E1) - c\bar{N}2 \) and the other two are obtained from the quadratic equation:

\[
\lambda^2 + (e\bar{N}2 + k\bar{N}3)\lambda + (ek - gm)\bar{N}2\bar{N}3 = 0
\] (3.41)

In (3.41), the sum of the roots \((-e\bar{N}2 + k\bar{N}3)\) is negative and the product of the roots \((ek - gm)\bar{N}2\bar{N}3\) is positive. Therefore the roots of (3.41) are real and negative or complex conjugates having negative real parts. Thus the state will be asymptotically stable when \((a - q1E1) < c\bar{N}2\).

#### iv. Stability of the positive interior equilibrium point

The variational matrix at the positive interior equilibrium point is:
Now we prove that the function \( J_b \) is negative semi definite. Therefore, \( (\bar{N}_1, \bar{N}_2, \bar{N}_3) \) is globally asymptotically stable.

2.3.2.2 Global Stability

**Theorem 2:** The equilibrium point \((\bar{N}_1, \bar{N}_2, \bar{N}_3)\) is globally asymptotically stable.

**Proof:**

Let consider the function which defined as:

\[
V(\bar{N}_1, \bar{N}_2, \bar{N}_3) = \left\{ \begin{array}{ll}
N_1 - \bar{N}_1 - \bar{N}_1 \ln \left[ \frac{\bar{N}_1}{N_1} \right] + \left\{ N_2 - \bar{N}_2 - \bar{N}_2 \ln \left[ \frac{\bar{N}_2}{N_2} \right] \right\} + \left\{ N_3 - \bar{N}_3 - \bar{N}_3 \ln \left[ \frac{\bar{N}_3}{N_3} \right] \right\} \\
\end{array} \right.
\]

for \( N_1 > \bar{N}_1, N_2 > \bar{N}_2 \) and \( N_3 > \bar{N}_3 \).

Now we prove that the function \( V \) is a Liapunov function. For this we need to show that:

i. \( V \) is continues and positive definite function
ii. \( \frac{dV}{dt} \) is negative semi definite

Now

i. The function \( V \) is positive definite, since \( N_1 > \bar{N}_1, N_2 > \bar{N}_2 \) and \( N_3 > \bar{N}_3 \).

ii. Now, differentiating \( V \) with respect to 't' we obtain:

\[
\frac{dV}{dt} = \left( 1 - \frac{N_1}{\bar{N}_1} \right) \frac{dN_1}{dt} + \left( 1 - \frac{N_2}{\bar{N}_2} \right) \frac{dN_2}{dt} + \left( 1 - \frac{N_3}{\bar{N}_3} \right) \frac{dN_3}{dt}
\]

Substituting (3.1), (3.2) and (3.3) in to (3.43) we obtain:

\[
\frac{dV}{dt} = (N_1 - \bar{N}_1)[a - bN_1 - cN_1 - q_1N_1] + (N_2 - \bar{N}_2)[d - eN_2 + fN_1 - gN_2 - q_2N_2] + (N_3 - \bar{N}_3)[h - kN_3 - mN_2 - q_3N_3]
\]

\[
= -b(N_1 - \bar{N}_1)^2 - \left\{ e + \frac{1}{2}c - f + g + m \right\} (N_2 - \bar{N}_2)^2 - \left\{ k + \frac{1}{2}m + g \right\} (N_3 - \bar{N}_3)^2 < 0
\]

This implies that \( \frac{dV}{dt} \) is negative semi definite. Therefore, \((\bar{N}_1, \bar{N}_2, \bar{N}_3)\) is globally asymptotically stable.

**Theorem 3:** The systems (3.1) – (3.3) cannot have any limit cycle in the interior of the positive quadrant.

**Proof:** Let \( H(N_1, N_2, N_3) = \frac{1}{N_1N_2N_3} \)

\[
h_1(N_1, N_2, N_3) = aN_1 - b(N_1)^2 - cN_1N_2 - q_1E_1N_1
\]

\[
h_2(N_1, N_2, N_3) = dN_2 - e(N_2)^2 + fN_1N_2 - gN_2N_3 - q_2E_2N_2
\]

\[
h_3(N_1, N_2, N_3) = hN_3 - k(N_3)^2 + mN_2N_3 - q_3E_3N_3
\]

Then \( H(N_1, N_2, N_3) > 0 \) in the interior of positive octant of \( \mathbb{X}^3 \)-space. Because \( N_1, N_2, N_3 > 0 \).

Now \( \Delta (N_1, N_2, N_3) = \frac{\partial}{\partial N_1}(Hh_1) + \frac{\partial}{\partial N_2}(Hh_2) + \frac{\partial}{\partial N_3}(Hh_3) \)

\[
= - \frac{b}{N_1N_2N_3} - \frac{e}{N_1N_2N_3} - \frac{k}{N_1N_2N_3}
\]

\[
= - \left\{ \frac{b}{N_1N_2N_3} + \frac{e}{N_1N_2N_3} + \frac{k}{N_1N_2N_3} \right\} < 0
\]

This shows that \( \Delta (N_1, N_2, N_3) \) does not change sign and identically zero in the positive octant of \( \mathbb{X}^3 \)-space. By Bendixon - Dulac criteria, it follows that the system (3.1) – (3.3) has no closed trajectories and hence no periodic solutions in the positive octant of \( \mathbb{X}^3 \)-space. Thus the system (3.1) - (3.3) cannot have any limit cycles in the interior of the positive octant.
2.4 Bionomic equilibrium of the imprecise prey–predator model

The bionomic equilibrium is nothing but the combination of the concepts of biological equilibrium as well as economic equilibrium. The biological equilibrium is given by equation (3.25). It is the study of the dynamics of living resources using economic models. Economic equilibrium is said to be achieved when the total revenue obtained by selling the harvested biomass (TR) equals to the total cost for the effort devoted to the harvesting (TC).

To discuss the bionomic equilibrium of the imprecise prey, predator and competitor to the predator model, we consider the following parameters.

Let \( c_1 \) be the harvesting cost per unit effort for prey species, \( p_1 \) be the price per unit biomass of the prey, \( c_2 \) be the harvesting cost per unit effort for predator species, \( p_2 \) be the price per unit biomass of the predator, \( c_3 \) be the harvesting cost per unit effort for competitor to the predator species and \( p_3 \) be the price per unit biomass of the competitor to the predator species. Then the net economic rent (net revenue) for the prey, predator and competitor to the predator at any time is given by:

\[
R = (p_1 q_1 N_1 - c_1) E_1 + (p_2 q_2 N_2 - c_2) E_2 + (p_3 q_3 N_3 - c_3) E_3 = R_1 + R_2 + R_3
\]

Where \( R_1 = (p_1 q_1 N_1 - c_1) E_1 \), \( R_2 = (p_2 q_2 N_2 - c_2) E_2 \) and \( R_3 = (p_3 q_3 N_3 - c_3) E_3 \). \( R_1, R_2 \) and \( R_3 \) represent the net revenues for the prey, predator and competitor to the predator species respectively.

The bionomic equilibrium of the imprecise prey–predator model is given by the following simultaneous equations.

\[
\begin{align*}
    a N_1 - b (N_1)^2 - c N_1 N_2 - q_1 E_1 N_1 &= 0 \\
    d N_2 - e (N_2)^2 + f N_1 N_2 - g N_2 N_3 - q_2 E_2 N_2 &= 0 \\
    h N_3 - k (N_3)^2 - m N_2 N_3 - q_3 E_3 N_3 &= 0 \\
    (p_1 q_1 N_1 - c_1) E_1 + (p_2 q_2 N_2 - c_2) E_2 + (p_3 q_3 N_3 - c_3) E_3 &= 0 
\end{align*}
\]

(3.44)

In order to determine the bionomic equilibrium we come across the following cases.

**Case (1):** When \( c_2 > p_2 q_2 N_2 \) and \( c_3 > p_3 q_3 N_3 \), the cost is greater than revenue for the predator and competitor to the predator species, then the predator and competitor to predator species are not harvested. The predator and competitor to the predator harvesting is stopped \((E_2 = 0, E_3 = 0)\). Then only prey harvesting remains operational \((c_1 < p_1 q_1 N_1)\). Therefore \( E_2 = E_3 = 0 \) and \( c_1 < p_1 q_1 N_1 \), we have \((N_1)_{\omega} = \frac{c_1}{p_1 q_1} \).

**Case (2):** When \( c_1 > p_1 q_1 N_1 \) and \( c_3 > p_3 q_3 N_3 \) the cost is greater than revenue for the prey and competitor to the predator species, then the prey and competitor predator species are not harvested. The prey and competitor to the predator harvesting is stopped \((E_1 = 0, E_3 = 0)\). Then only predator harvesting remains operational \(c_2 < p_2 q_2 N_2\). Therefore \( E_1 = E_3 = 0 \) and \( c_2 < p_2 q_2 N_2 \), we have \((N_2)_{\omega} = \frac{c_2}{p_2 q_2} \).

**Case (3):** When \( c_1 > p_1 q_1 N_1 \) and \( c_2 > p_2 q_2 N_2 \), the cost is greater than revenue for the prey and predator species, then the prey and predator species are not harvested. The prey and predator harvesting is stopped \((E_1 = E_2 = 0)\). Then only predator harvesting remains operational \(c_3 < p_3 q_3 N_3\). Therefore \( E_1 = E_2 = 0 \) and \( c_3 < p_3 q_3 N_3 \), we have \((N_3)_{\omega} = \frac{c_3}{p_3 q_3} \).

**Case (4):** When \( c_1 > p_1 q_1 N_1 \), \( c_2 > p_2 q_2 N_2 \) and \( c_3 > p_3 q_3 N_3 \) then the cost is greater than revenues for both species and the whole harvesting will be closed.

**Case (5):** When \( c_1 < p_1 q_1 N_1 \), \( c_2 < p_2 q_2 N_2 \) and \( c_3 < p_3 q_3 N_3 \) the cost is less than the revenue for both species, then both species is harvested. This implies that the revenues for both species being positive; so the whole harvesting will be in operation (the system becomes operational). In this case \((N_1)_{\omega} = \frac{c_1}{p_1 q_1}, (N_2)_{\omega} = \frac{c_2}{p_2 q_2}, (N_3)_{\omega} = \frac{c_3}{p_3 q_3}\) and \((E_1)_{\omega} = \frac{c_1}{p_1 q_1}\). By substituting the values of \((N_1)_{\omega}, (N_2)_{\omega}, (N_3)_{\omega}\) in to the 1st, 2nd, and 3rd equation of (3.44) we obtain:

\[
\begin{align*}
    (E_1)_{\omega} &= \frac{1}{q_1} \left[ a p_1 q_1 q_2 x_1 - c p_1 q_1 c_1 \right] \\
    (E_2)_{\omega} &= \frac{1}{q_2} \left[ d p_1 q_2 q_3 q_4 x_2 - c p_2 q_2 q_3 c_1 + f p_1 q_1 q_3 q_4 c_2 \right] \\
    (E_3)_{\omega} &= \frac{1}{q_3} \left[ h p_2 q_3 q_4 x_3 - k p_2 q_2 q_3 c_3 + m p_3 q_3 q_4 c_2 \right] \\
\end{align*}
\]

(3.45) \hspace{1cm} (3.46) \hspace{1cm} (3.47)

But

\[
\begin{align*}
    (E_1)_{\omega} &> 0 \text{ if } a p_1 q_1 q_2 x_1 > b p_2 q_2 q_3 + c p_1 q_1 c_1 \\
    (E_2)_{\omega} &> 0 \text{ if } d p_2 p_3 q_1 q_4 q_3 > f p_1 p_2 q_3 q_5 q_4 c_1 > e p_3 p_4 q_1 q_2 c_2 + g p_2 p_1 q_1 q_4 c_3 \\
    (E_3)_{\omega} &> 0 \text{ if } h p_2 q_3 q_4 x_3 > k p_2 q_2 q_3 c_3 + m p_3 q_3 q_4 c_2 \\
\end{align*}
\]

(3.48) \hspace{1cm} (3.49) \hspace{1cm} (3.50)

Thus the nontrivial bionomic equilibrium point \([(N_1)_{\omega}, (N_2)_{\omega}, (N_3)_{\omega}, (E_1)_{\omega}, (E_2)_{\omega}, (E_3)_{\omega}]\) exist if conditions (3.48) - (3.50) hold.
2.5 Qualitative Analysis Of Optimal Harvesting Policy

In commercial exploitation of renewable resources the fundamental problem from the economic point of view, is to determine the optimal trade-off between present and future harvests. If we look at the problem it is observed that the marine fishery sectors become more important not only for domestic demand but also from the imperatives of exports.

In this section we study optimal harvesting policy of the system of equation (3.1) - (3.3); and also our objective is to maximize, the objective functional form of the harvesting model, with the instantaneous annual rate of discount \( \delta \) is as follows:

\[
J(E_1, E_2, E_3) = \int_0^\infty e^{-\delta t} [(p_1q_1N_1 - c_1)E_1(t) + (p_2q_2N_2 - c_2)E_2(t) + (p_3q_3N_3 - c_3)E_3(t)] \, dt
\]

Subject to the state constraints (3.1) - (3.3) with control constraints (variables):

\[
0 \leq E_i(t) \leq E_i^{max}, i = 1, 2, 3
\]

First we construct the following Hamiltonian function for the problem by:

\[
H = e^{-\delta t}[\lambda_1(\dot{E}_1 - p_1q_1N_1 + c_1) + \lambda_2(\dot{E}_2 - p_2q_2N_2 + c_2) + \lambda_3(\dot{E}_3 - p_3q_3N_3 + c_3)]
\]

where \( \lambda_1, \lambda_2, \lambda_3 \) are additional unknown functions called the adjoint variables.

Now by differentiating \( H \) with respect to \( E_1, E_2 \) and \( E_3 \) respectively, we obtain:

\[
\frac{\partial H}{\partial E_1} = e^{-\delta t}(p_1q_1N_1 - c_1) - \lambda_1q_1N_1 = \varphi_1(t)
\]

\[
\frac{\partial H}{\partial E_2} = e^{-\delta t}(p_2q_2N_2 - c_2) - \lambda_2q_2N_2 = \varphi_2(t)
\]

\[
\frac{\partial H}{\partial E_3} = e^{-\delta t}(p_3q_3N_3 - c_3) - \lambda_3q_3N_3 = \varphi_3(t)
\]

The optimal control \( E_i(t) \) must satisfy the condition:

\[
E_i(t) = \begin{cases} 
E_i^{max} & \text{if } \varphi_i(t) > 0 \\
0 & \text{if } \varphi_i(t) < 0 
\end{cases} \text{ for } i = 1, 2, 3
\]

Since \( \varphi_i(t) \) causes \( E_i(t), \quad (i = 1, 2, 3) \) to switch between 0 and \( E_i^{max} \) so \( \varphi_i(t) \) \( (i = 1, 2, 3) \) are called switching function. Depending on the sign of the switching function \( \varphi_i(t) \), the optimal control \( E_i(t) \) is a bang–bang switching from one extreme point to other one. When \( \varphi_i(t) = 0 \) \( (i = 1, 2, 3) \), the Hamiltonian function \( H \) becomes independent of the control variable \( E_i(t) \), \( (i = 1, 2, 3) \) and the optimal control cannot be determined by the above procedure. It is then called a singular control \( E_i(t), 0 < E_i(t) < E_i^{max} \). Hence the optimal harvesting policy is

\[
E_i(t) = \begin{cases} 
E_i^{max} & \text{if } \varphi_i(t) > 0 \\
0 & \text{if } \varphi_i(t) < 0 
\end{cases} \text{ for } i = 1, 2, 3
\]

The aim is to find an optimal equilibrium \([N_1, N_2, N_3, E_1, E_2, E_3] \) to maximize Hamiltonian \( H \). Since Hamiltonian \( H \) is linear in the control variables \( E_1, E_2 \) and \( E_3 \), the optimal control can be extreme controls or the singular controls.

Thus for singular control \( \varphi_i(t) = 0 \) \( (i = 1, 2, 3) \), from equations (3.53) - (3.55) solving for \( \lambda_i \) \( (i = 1, 2, 3) \) we obtain:

\[
\lambda_1 = e^{-\delta t} \left( p_1 - \frac{c_1}{q_1N_1} \right)
\]

\[
\lambda_2 = e^{-\delta t} \left( p_2 - \frac{c_2}{q_2N_2} \right)
\]

\[
\lambda_3 = e^{-\delta t} \left( p_3 - \frac{c_3}{q_3N_3} \right)
\]

In this case, the optimal control is called the singular control and equations (3.58) - (3.60) are the necessary conditions for the maximization of Hamiltonian \( H \).

By Pontryagin’s Maximum Principle, the adjoint equations are:
Now, by using equation (3.61) we obtain:
\[
\frac{dx_1}{dt} = -\frac{1}{\alpha_1} \frac{d\alpha_1}{dt} \frac{dx_2}{dt} = -\frac{1}{\alpha_2} \frac{d\alpha_2}{dt} \frac{dx_3}{dt} = -\frac{1}{\alpha_3} \frac{d\alpha_3}{dt}
\]  
(3.61)

Substituting \( a = bN_1 + cN_2 + q_1E_1 \) and (3.59) into the equation (3.62) we obtain:
\[
\frac{dx_1}{dt} = -e^{-\delta t} p_1 q_1 E_1 - \lambda_1 (a - 2bN_1 - cN_2 - q_1 E_1) - \lambda_2 f N_2
\]  
(3.62)

To also obtain a singular path:
\[
\frac{dx_1}{dt} = -e^{-\delta t} p_1 q_1 E_1 + bN_1 \lambda_1 - e^{-\delta t} \left( p_2 - \frac{c_2}{q_2 N_2} \right)
\]  
(3.63)

This is of the form:
\[
\frac{dx_1}{dt} + A_1 x_1 = -A_2 e^{-\delta t}
\]

(3.64)

and also:
\[
\frac{dx_2}{dt} = -e^{-\delta t} p_2 q_2 E_2 + cN_1 \lambda_1 - \lambda_2 (d - 2eN_2 + f N_1 - gN_3 - q_2 E_2) + mN_2 \lambda_3
\]  
(3.65)

Substituting \( d = eN_2 - f N_1 + gN_3 + q_2 E_2 \) and equations (3.58) into (3.65) we obtain:
\[
\frac{dx_2}{dt} + B_1 \lambda_2 = B_2 e^{-\delta t}
\]

(3.66)

where \( B_1 = -eN_2 + B_2 = (p_1 - \frac{c_1}{q_1 N_1}) cN_1 + (p_3 - \frac{c_3}{q_3 N_3}) mN_3 - p_2 q_2 E_2 \)

The above equation is linear in \( \lambda_2 \) and its solution is given by:
\[
\lambda_2 = \frac{B_2}{B_1 - \delta} e^{-\delta t}
\]

(3.67)

Similarly,
\[
\frac{dx_3}{dt} = -e^{(\delta - \sigma)} p_3 q_3 E_3 + gN_2 \lambda_2 - \lambda_3 (h - 2kN_3 - mN_2 - q_3 E_3)
\]

(3.68)

Now by substituting \( h = kN_3 + mN_2 + q_3 E_3 \), equation (3.58) and (3.59) into (3.68) we obtain:
\[
\frac{dx_3}{dt} + C_1 \lambda_3 = C_2 e^{-\delta t}
\]

(3.69)

where \( C_1 = -kN_3 \) and \( C_2 = \left( p_2 - \frac{c_2}{q_2 N_2} \right) gN_2 - p_2 q_3 E_3 \)

The above equation is linear in \( \lambda_3 \) and its solution is given by:
\[
\lambda_3 = \frac{C_2}{C_3 - \delta} e^{-\delta t}
\]

(3.70)

It is obviously that \( \lambda_1(t), \lambda_2(t) \) and \( \lambda_3(t) \) are bounded as \( t \to \infty \). From (3.58) and (3.64) we obtain a singular path:
\[
p_1 - \frac{c_1}{q_1 N_1} = -\frac{A_2}{A_1 - \delta}
\]

(3.71)

And also from (3.59) and (3.67), we obtain a singular path:
\[
p_2 - \frac{c_2}{q_2 N_2} = \frac{B_2}{B_1 - \delta}
\]

(3.72)

Similarly, from (3.60) and (3.70) we obtain a singular path:
\[
p_3 - \frac{c_3}{q_3 N_3} = \frac{C_2}{C_3 - \delta}
\]

(3.73)

Let \( F(N_1) = \left( p_1 - \frac{c_1}{q_1 N_1} \right) + \frac{A_2}{A_1 - \delta} \), \( G(N_2) = \left( p_2 - \frac{c_2}{q_2 N_2} \right) - \frac{B_2}{B_1 - \delta} \) and \( M(N_3) = \left( p_3 - \frac{c_3}{q_3 N_3} \right) - \frac{C_2}{C_3 - \delta} \) then there exists a unique positive root \( \tilde{N}_1 = (N_1)_0 \) of \( F(N_1) = 0 \) in the interval \( 0 < \tilde{N}_1 < k_1 \) if the following inequalities hold:
\[
F(0) < 0, F'(k_1) > 0, F'(<N_1) > 0 \text{ for } \tilde{N}_1 > 0
\]

And also there exists a unique positive root \( \tilde{N}_2 = (N_2)_0 \) of \( G(N_2) = 0 \) in the interval \( 0 < \tilde{N}_2 < k_2 \) if the following inequalities hold:
\[
G(0) < 0, G'(k_2) > 0, G'(\tilde{N}_2) > 0 \text{ for } \tilde{N}_2 > 0
\]

Similarly, there exists a unique positive root \( \tilde{N}_3 = (N_3)_0 \) of \( M(\tilde{N}_3) = 0 \) in the interval \( 0 < \tilde{N}_3 < k_3 \) if the following inequalities hold:
\[
M(0) < 0, M'(k_3) > 0, M'(\tilde{N}_3) > 0 \text{ for } \tilde{N}_3 > 0
\]
Now, for $\bar{N}_1 = (N_1)_0$, $\bar{N}_2 = (N_2)_0$ and $\bar{N}_3 = (N_3)_0$ we get:

$$(N_1)_0 = \frac{c_1}{q_1 N_1}, \quad (N_2)_0 = \frac{c_2}{p_2 q_2}, \quad (N_3)_0 = \frac{c_3}{p_3 q_3}$$

$$(E_1)_\delta = \frac{p_1 p_2 (q_1^2 q_2)}{p_1 p_2 p_3 q_3 q_2 (q_2^2 q_3)}$$

$$(E_2)_\delta = \frac{d p_1 p_2 p_3 q_1 q_2 q_3 - e p_1 p_2 q_1 q_3 c_2}{p_1 p_2 p_3 q_3 (q_2^2 q_3)}$$

and

$$(E_3)_\delta = \frac{h p_2 p_3 q_2 q_3 - k p_2 q_2 c_3 - m p_3 q_3 c_2}{p_2 p_3 q_3 (q_3^2)}$$

Hence once the optimal equilibrium $((N_1)_\delta, (N_2)_\delta, (N_3)_\delta)$ is determined, the optimal harvesting effort $((E_1)_\delta, (E_2)_\delta, (E_3)_\delta)$ can be determined. From (3.64), (3.67) and (3.70) we found that $\lambda_i(t)$ where $i = 1, 2, 3$ do not vary with time in optimal equilibrium. Hence they remains bounded as $t \to \infty$.

From (3.71), (3.72) and (3.73) we note that:

$$p_1 - \frac{c_1}{q_1 N_1} = -\frac{A_2}{A_1 - \delta} \to 0 \text{ as } \delta \to 0$$

$$p_2 - \frac{c_2}{q_2 N_2} = \frac{B_2}{B_1 - \delta} \to 0 \text{ as } \delta \to 0$$

and

$$p_3 - \frac{c_3}{q_3 N_3} = \frac{C_2}{C_1 - \delta} \to 0 \text{ as } \delta \to 0$$

Thus, the net economic revenue for the Prey population $R_1 = 0$, the net economic revenue for the predator population $R_2 = 0$ and the net economic revenue for the competitor to the Predator population $R_3 = 0$. From this we conclude that, if the discount rate increases, then the net economic revenue decrease and even may tend to zero if the discount rate tends to infinity. Hence finally we remarked that high interest rate will cause high inflation rate.

3. Numerical simulation

In this section, we substantiate as well as augment our analytical findings through numerical simulations considering the interval parameters. For this, numerical examples are obtained to illustrate the proposed methodology presented in this paper.

**Example:** Let us consider a set of artificial values of parameters as follows in appropriate units:

$r = [2.5, 3]$, $\alpha_1 = [0.2, 0.6]$, $\alpha_2 = [0.7, 0.8]$, $\alpha_3 = [0.5, 0.9]$, $s = [4.5, 5]$, $\beta_1 = [0.8, 0.9]$, $\beta_2 = [0.2, 0.4]$, $\beta_3 = [0.5, 0.6]$, $\ell = [5.5, 6]$, $\delta_1 = [0.2, 0.5]$, $q_1 = 0.2$, $q_2 = 0.5$, $q_3 = 0.8$, $E_1 = 10$, $E_2 = 8$, $E_3 = 6$ and $p \in [0, 1]$.

The trivial equilibrium point $(0,0,0)$ always exists for all values of $P \in [0,1]$. The non trivial equilibrium points and the Eigen values of variational matrices at the corresponding points of co-existence equilibriums are given in table1 below for different values of $P$ in $[0,1]$.  

---

**Table 1:**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Value</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>$r$</td>
<td>$[2.5, 3]$</td>
<td>Rate of change</td>
</tr>
<tr>
<td>$\alpha_1$</td>
<td>$[0.2, 0.6]$</td>
<td>Birth rate of Prey</td>
</tr>
<tr>
<td>$\alpha_2$</td>
<td>$[0.7, 0.8]$</td>
<td>Birth rate of Predator</td>
</tr>
<tr>
<td>$\alpha_3$</td>
<td>$[0.5, 0.9]$</td>
<td>Birth rate of Competitor</td>
</tr>
<tr>
<td>$s$</td>
<td>$[4.5, 5]$</td>
<td>Survival rate of Prey</td>
</tr>
<tr>
<td>$\beta_1$</td>
<td>$[0.8, 0.9]$</td>
<td>Death rate of Predator</td>
</tr>
<tr>
<td>$\beta_2$</td>
<td>$[0.2, 0.4]$</td>
<td>Death rate of Prey</td>
</tr>
<tr>
<td>$\beta_3$</td>
<td>$[0.5, 0.6]$</td>
<td>Death rate of Competitor</td>
</tr>
<tr>
<td>$\ell$</td>
<td>$[5.5, 6]$</td>
<td>Carrying capacity</td>
</tr>
<tr>
<td>$\delta_1$</td>
<td>$[0.2, 0.5]$</td>
<td>Infection rate</td>
</tr>
<tr>
<td>$q_1$</td>
<td>$0.2$</td>
<td>Immersion rate of Prey</td>
</tr>
<tr>
<td>$q_2$</td>
<td>$0.5$</td>
<td>Immersion rate of Predator</td>
</tr>
<tr>
<td>$q_3$</td>
<td>$0.8$</td>
<td>Immersion rate of Competitor</td>
</tr>
<tr>
<td>$E_1$</td>
<td>$10$</td>
<td>Environmental impact on Prey</td>
</tr>
<tr>
<td>$E_2$</td>
<td>$8$</td>
<td>Environmental impact on Predator</td>
</tr>
<tr>
<td>$E_3$</td>
<td>$6$</td>
<td>Environmental impact on Competitor</td>
</tr>
<tr>
<td>$p$</td>
<td>$[0, 1]$</td>
<td>Discount rate</td>
</tr>
</tbody>
</table>

---

21
The fluctuation of prey, predator and competitor to the predator population with respect to time beginning with that the eigen values corresponding to different equilibrium points are complex conjugate with negative real part.

From the above table we observe that there are different equilibrium points for different values of p. We also see that the eigen values corresponding to different equilibrium points are complex conjugate with negative real part. Therefore the equilibrium points are stable.

The fluctuation of prey, predator and competitor to the predator population with respect to time beginning with \( N_1 = 1, N_2 = 1.09 \) and \( N_3 = 1.1 \) for \( p = 0.7, p = 0.8, p = 0.9 \) and \( p = 1 \) are depicted in following Figures respectively.

### Table 3.2 Equilibrium states, eigen values, nature of the equilibrium points.

<table>
<thead>
<tr>
<th>Values of p</th>
<th>Equilibrium states</th>
<th>Eigen values</th>
<th>Nature of the equilibrium states</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>0.29, 0.362, 0.536</td>
<td>(-0.636, -0.156 - 0.111i, -0.156 + 0.111i)</td>
<td>Stable</td>
</tr>
<tr>
<td>0.1</td>
<td>0.294, 0.435, 0.578</td>
<td>(-0.674, -0.144 - 0.117i, -0.144 + 0.117i)</td>
<td>Stable</td>
</tr>
<tr>
<td>0.2</td>
<td>0.292, 0.515, 0.623</td>
<td>(-0.665, -0.18 - 0.126i, -0.18 + 0.126i)</td>
<td>Stable</td>
</tr>
<tr>
<td>0.3</td>
<td>0.275, 0.601, 0.661</td>
<td>(-0.793, -0.164 - 0.125i, -0.164 + 0.125i)</td>
<td>Stable</td>
</tr>
<tr>
<td>0.4</td>
<td>0.281, 0.674, 0.729</td>
<td>(-0.783, -0.188 - 0.156i, -0.188 + 0.156i)</td>
<td>Stable</td>
</tr>
<tr>
<td>0.5</td>
<td>0.275, 0.755, 0.791</td>
<td>(-0.812, -0.179 - 0.155i, -0.179 + 0.155i)</td>
<td>Stable</td>
</tr>
<tr>
<td>0.6</td>
<td>0.261, 0.842, 0.856</td>
<td>(-0.846, -0.168 - 0.148i, -0.168 + 0.148i)</td>
<td>Stable</td>
</tr>
<tr>
<td>0.7</td>
<td>0.322, 0.929, 0.967</td>
<td>(-0.943, -0.193 - 0.178i, -0.193 + 0.178i)</td>
<td>Stable</td>
</tr>
<tr>
<td>0.8</td>
<td>0.149, 1.079, 0.928</td>
<td>(-0.962, -0.254, -0.128)</td>
<td>Stable</td>
</tr>
<tr>
<td>0.9</td>
<td>0.195, 1.134, 1.162</td>
<td>(-1.017, -0.197 - 0.124i, -0.197 + 0.124i)</td>
<td>Stable</td>
</tr>
<tr>
<td>1</td>
<td>0.2, 1.2, 1.2</td>
<td>(-1.072, -0.204 - 0.122i, -0.204 + 0.122i)</td>
<td>Stable</td>
</tr>
</tbody>
</table>

Stable, Stable, Stable, Stable, Stable, Stable, Stable, Stable, Stable, Stable

![Graph](image_url)
Population sizes at $p=0.8$ and $p=0.9$. The figure shows the predation model with three populations: prey, predator, and competitor.
4. CONCLUSIONS AND RECOMMENDATIONS

4.1 Conclusions

Prey–predator (competitor) harvesting model has undergone different development in theoretical and practical applications in the field of biomathematics. Most of the researchers have developed the prey, predator and competitor to the predator harvesting model based on the assumption that the biological parameters are precisely known but the scenario is different in real life situation. In this paper, we developed a method to find the biological equilibrium points, bio-economic equilibrium points and optimal harvesting policy when some biological parameters are imprecise in nature. Here we develop the concepts imprecise parameters to the prey, predator and competitor to the predator harvesting model by considering the prey population growth rate, predator population growth rate, competitor to the predator growth rate and predation coefficients are imprecise in nature for the lack of precise numerical information. The ability of calculating the biological equilibrium points, bio-economic equilibrium points and optimal harvesting policy developed in this paper might help to develop more realistic mathematical models in the area of mathematical biology. Before ending this article we would like to mention that one may consider Lotka–Volterra model with logistic growth under imprecise biological parameters. Impreciseness of the harvesting cost and price of the biomass of the species of the harvesting model are also important characteristic to be considered.

4.2 Recommendation

Basing on the results of qualitative analysis and numerical simulation of the model, we recommend that;

i. Prey-predator (competitor) should not be harvested at a rate higher than their growth rate. However optimal harvesting of the prey-predator (competitor) at a rate much lower than their growth rate is permissible, since this would not lead to collapse of the system in the long term.

ii. The population density of the predator can be increased drastically by increasing the growth rate of the prey species e.g. regular recruiting more prey into the area. Since regular recruiting of prey may not be realistic, the best alternative is to minimize or stop poaching of the preys so as to greatly increase the number of their population in that area, which will in turn result in an increase in the population of the predator. But the number of population of competitor to the predator does not dependent on the number of prey population; it depends on the number of predator population that competes’ with them for common resource. This common resource may be additional resources for predator population.

iii. The population density of the predator depends mainly on the biomass of the prey than that of
competitor, hence any attempt to control the population density of the predator should be based on controlling the population density of the prey.

Fig.2 prey-predator examples

References

- B. Dubey, P. Chandra, P. Sinha, A model for fishery resource with reserve area, Nonlinear Anal. Real


