Modeling and Stability Analysis of Host-parasite Population Dynamics

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To cite this article:
Geremew Kenassa Edessa, Purnachandra Rao Koya. Modeling and Stability Analysis of Host-parasite Population Dynamics. Mathematical Modelling and Applications. Vol. 5, No. 2, 2020, pp. 118-128. doi: 10.11648/j.mma.20200502.17

Received: January 1, 2020; Accepted: May 5, 2020; Published: May 28, 2020

Abstract: In this study, a mathematical model is developed to explore the population dynamics of two host species. Both the hosts depend on the same resources and the availability of such resources is limited in nature. If the host populations increase abnormally the limited natural resources will be used up. Hence, the concept of parasite is brought in to the picture to regulate the host populations. The parasite is a mechanism that reduces the host populations. However, on one hand if the parasite attacks more the hosts may extinct and on the other hand if the parasite do not attack then the host populations may increase and resource may be used up. Hence, the parasite is expected to maintain a balance so that neither the host populations nor the resources extinct. Here, both the hosts are classified in to susceptible and infected and hence the model comprises of four populations: Susceptible Host–1, Infected Host–1, Susceptible Host–2 and Infected Host–2. Thus, the mathematical model comprises of a system of four first order non-linear ordinary differential equations. Mathematical analysis of the model is conducted. Positivity and boundedness of the solution have been verified and thus shown that the model is physically meaningful and biologically acceptable. Equilibrium points of the model are identified and stability analysis is conducted. Simulation study is conducted in order to support the mathematical analysis using software packages Mat lab and DeDiscover.

Keywords: Modeling, Hosts, Parasite-mediated Interactions, Stability, Numerical Simulation

1. Introduction
Parasites can theoretically encourage indirect interactions among ecological communities those live in the same trophic level. These interactions lead to some kind of effects on the competing species. Such interactions are also called as parasite–mediated competition [1-2, 4, 13-15, 19]. However, parasites induce competition–like indirect interactions among species that would not otherwise happen at all [5, 8]. These ideas reflect indirect effects of parasites on community composition leading to coexistence or promoting exclusion.

1.1. Parasite–mediated Competition
Parasites may affect competitive dynamics between native and introduced animals and plants through density and trait effects [3, 6, 13-15]. Models of parasite-mediated competition have provided insight into the role of shared parasites in shaping communities of competing species. Parasites can regulate the population density of their hosts and, as a result, affect the dynamics of the community, and can cause the extinction of local populations and of whole species [2, 3, 13-15].

In this present study, parasites are predicted to enhance the range of conditions leading to coexistence if, for example, one of the host are more heavily impacted by the parasite. However, some arguments argued that the predominant pressure exerted by parasites on communities might not be the result of catastrophic outbreaks, but rather of less virulent and persistent where the parasites moderate or enhance the competitive capabilities of their hosts [3, 13].

When viewed as part of an ecosystem, the parasitized host is subject to numerous other interactions that convey benefits, no apparent benefits, or harm or not benefits to it [11, 12, 5, 20]. Argued that the direct outlay to the host that reduce its fitness can be either aggravated by indirect consequences (i.e. where parasitism renders predation more likely), or improved by indirect consequences.
1.2. Apparent Competition

In the context of parasitism, apparent competition is predicted to occur when two species that do not otherwise interact both host with the same parasite species [16-18, 22]. As both host species are a resource for the parasite, population density increases in either host lead to reductions in the other, via the, density mediated, negative effects of the parasite. Apparent competition can theoretically lead to the elimination of one host, indirectly coupling the dynamics of host species with different habitat or resource requirements.

In view of these, this study targeted to present these synthesized empirical and theoretical work using mathematical modeling and analysis qualitatively how parasites influences competitive and predatory interactions between species [1, 7, 10, 13, 21, 23]. We highlight the sensitive parameters to assess the impact of parasites on communities by incorporating the theoretical and empirical studies so as to examine how the effects of parasitism. Moreover, parasitism scales up the community-level processes.

In this paper, comparison of the model system would be carried out through the investigation of the interaction made between the populations directly and indirectly to each other and with parasite.

2. Assumption and Model Formulation

Accordingly, Mathematical epidemiology to study the dynamics of diseases spread has become an interesting topic of research study and received much attention from scientists after the pioneering work of [2, 3, 13-15, 20].

In this recent study, by applying a deterministic framework of an ecological model of the infection of parasite, the analysis of the population dynamics of eco-epidemiology would be investigated.

To start with, the mathematical model of a single type of host and its parasite would be defined by considering the following descriptions. These are the changes in the numbers of free-living parasite. These changes are due to three minimum processes: host birth, mortality due to parasite infection and natural death of a host. Hence, the classical model of the combination of host and parasite population growth with carrying capacity of host which follows the general deterministic trajectories defined as

\[
\frac{dH}{dt} = rH \left(1 - \frac{H}{k}\right) - \delta HP 
\]

(1)

\[
\frac{dP}{dt} = \delta mHP - dP;
\]

(2)

This model is shown with simple diagram in figure 1. This system is similar to a predator-prey system but differing by the quantity \(m\) and called in host-parasite system burst size where as in predator-prey system ecological efficiency. Regardless of the potentially widespread effects of parasites on the community, most studies of host-parasite dynamics focus on the effects of a single parasite on a single host, neglecting the host’s interactions with each other and with other species.

The present model considers interaction of two hosts. These hosts do not directly interact with each other but they are mediated by a common enemy or parasite.

In this paper, the recovery of the hosts would not be considered. Moreover, in this study the interactions involving parasites that feed on a host individual usually living on or in it and often causing harm but not immediate death and the carrying capacity is only allowed for the uninfected hosts. i.e., for infected hosts limited resources are available. Further, responses of the hosts are linear rather than saturated. In this assumption, the model for free living parasite is not explicitly modeled.

The interaction between the states of the present model is illustrated in Figure 2 below;

The present mathematical model is formulated based on the above schematic model, and is given by the following system of differential equation:

\[
\frac{dx_1}{dt} = ax_1 \left(1 - \frac{x_1 + x_2}{k}\right) - \mu x_1 x_2 - ax_1 
\]

(3)

\[
\frac{dx_2}{dt} = \mu x_1 x_2 - Ax_2; A = \alpha + \delta
\]

(4)

\[
\frac{dy_1}{dt} = by_1 \left(1 - \frac{y_1 + y_2}{k'}\right) - ry_1 y_2 - \theta y_1
\]

(5)

\[
\frac{dy_2}{dt} = ry_1 y_2 - By_2; B = \beta + \theta
\]

(6)

Here, \(X(t) = x_1(t) + x_2(t)\) and \(Y(t) = y_1(t) + y_2(t)\) denote total population sizes of the first and second hosts respectively. However, the total size of both the hosts in the system is given by \(W(t) = X(t) + Y(t)\).

Descriptions of notations, parametric values of the system
are given in the following tables.

### Table 1. Notations and descriptions of the model variables.

| Variables | Descriptions |
|-----------|--------------|
| $x_1(t)$ | Density of first susceptible host population |
| $x_2(t)$ | Density of first infectious host population |
| $y_1(t)$ | Density of second susceptible host population |
| $y_2(t)$ | Density of second infectious host population |

### Table 2. Notations and descriptions of the parameters used in the model.

| Parameters | Descriptions |
|-----------|--------------|
| $a$ | Growth rate of host–1 |
| $b$ | Growth rate of host–2 |
| $μ$ | Infection transmission rate of host–1 |
| $r$ | Infection transmission rate of host–2 |
| $α$ | Total death rate of host–1 |
| $β$ | Total death rate of host–2 |
| $k_1$ | Environmental carrying capacity for host–1 |
| $k_2$ | Environmental carrying capacity for host–2 |

### 3. Qualitative Analysis of the Model Systems

#### 3.1. Positivity and Boundedness of the System

For the model to be meaningful and well posed, it is necessary that the solutions of (3)–(6) with positive initial data are positive and bounded for all $t > 0$. This fact has been stated and proved in the following:

Proposition 1: All solutions of the model equations (3)–(6) are non-negative and bounded.

**Proof:**

**Positivity of the populations:** consider the equation of system (3)–(6).

Up on integrating the model equations (3)–(6) with respect to $t$ the analytical solutions of the model variables are obtained as

$$x_1(t) = x_{10} \exp \left\{ \int_0^t \left[ a \left( 1 - \frac{x_1(s) + x_2(s)}{k} \right) - \mu x_2(s) - \alpha \right] ds \right\}$$

$$x_2(t) = x_{20} \exp \left\{ \int_0^t (\mu x_4(q) - \alpha) dq \right\}$$

$$y_1(t) = y_{10} \exp \left\{ \int_0^t \left[ b \left( 1 - \frac{y_1(u) + y_2(u)}{k} \right) - r y_2(u) - \beta \right] du \right\}$$

$$y_2(t) = y_{20} \exp \left\{ \int_0^t (r y_4(v) - \beta) dv \right\}$$

Here, it can be observed that each solution is a product of an initial condition and an exponential function. However, by definition every initial condition and exponential function are non-negative quantities, so are their products.

Therefore, all solutions of the system of equations (3)–(6) are non-negatives for all $t \geq 0$.

**Boundedness of the system**

In theoretical eco-epidemiology, a system is bounded implies that the system is biologically valid and well behaved. Here in the present study, the biological validity of the model is achieved by showing that the solutions of (3)–(6) are bounded.

In order to show the system is bounded, let the expression for the total population of the system as $W(t) = x_1(t) + x_2(t) + y_1(t) + y_2(t)$ be considered. Also, let $mW = m(x_1 + x_2 + y_1 + y_2)$ and $(dW/dt) = (dx_1/dt) + (dx_2/dt) + (dy_1/dt) + (dy_2/dt)$ where $m$ is any positive constant. Now, using the fore going expressions in the addition of (3)–(6) reduces the result as:

$$\frac{dW}{dt} + mW = ax_1 \left( 1 - \frac{x_1 + x_2}{k} \right) + by_1 \left( 1 - \frac{y_1 + y_2}{k} \right) + by_1 \left( 1 - \frac{y_1 + y_2}{k} \right) + m(x_1 + y_1) + my_1,$$

For $A$ and $B > m$

$$\frac{dW}{dt} + mW \leq ax_1 \left( 1 - \frac{x_1 + x_2}{k} \right) + by_1 \left( 1 - \frac{y_1 + y_2}{k} \right) + m(x_1 + y_1),$$

$$\frac{dW}{dt} + mW \leq (a + m)x_1 - a \frac{x_1}{k} - a \frac{x_2}{k} + (b + m)y_1 - b \frac{y_1}{k} - b \frac{y_2}{k}$$

$$\leq (a + m)x_1 - a \frac{x_1}{k} + (b + m)y_1 - b \frac{y_2}{k} - b \frac{y_2}{k}$$

$$\leq \frac{k}{4a} (a + m)^2 - \frac{a}{k} \left( x_1 - \frac{k}{2a} (a + m) \right)^2 + \frac{k'}{4b} (b + m)^2$$

$$\leq (k/4a)(a + m)^2 + (k'/4b)(b + m)^2$$

Now, introducing notation $\omega = (k/4a)(a + m)^2 + (k'/4b)(b + m)^2$ the differential inequality reduces to the form $(dW/dt) + mW \leq \omega$. It is a first order ordinary differential equation with constant coefficients and its analytical solution is given by $W(t) \leq (\omega/m)(1 - e^{-mt}) + W_0 e^{-mt}$. However, in the limit as $t \to \infty$, the solution takes the form as $W(t) \leq (\omega/m)$showing that the total population size is bounded. Hence, each population size is also bounded. Thus, the statement holds true.

#### 3.2. Existence of Steady States of the Model

The model equations (3)–(6) possess equilibrium points which are biologically feasible. These are obtained by setting $(dx_1/dt) = (dx_2/dt) = (dy_1/dt) = (dy_2/dt) = 0$ and solving the resultant expressions. Thus, the nine equilibrium points of the model and the corresponding existence conditions are listed as follows:

1. Trivial equilibrium $E_0 = (0, 0, 0, 0, 0)$ always exists.
   At this all the populations are absent.
2. Axial equilibrium $E_1 = (x_1^*, 0, 0, 0, 0)$ exists if $a > \alpha$. Here, only susceptible host–1 is present.
3. Axial equilibrium $E_2 = (0, 0, y_1^*, 0, 0)$ exists...
if $b > 0$. Here, only susceptible host–2 is present.

4. Disease free equilibrium $E_3 = (x_1^*, 0, y_1^*, 0)$ exists.

At this equilibrium both susceptible host–1 and host–2 are present.

5. Equilibrium $E_4 = (x_1^{**}, x_2^{**}, 0, 0)$ exists if $a[1 - (A/\mu k)] > \alpha$ and $\mu k > A$. Here both susceptible and infected populations of host–1 are present.

6. Equilibrium $E_5 = (0, 0, y_1^{**}, y_2^{**})$ exists if $b[1 - (B/rk')] > \theta$ and $rk' > B$. Here both susceptible and infected populations of host–2 only exist.

7. Equilibrium $E_6 = (x_1^*, 0, y_1^*, y_2^{**})$ exists. This is a disease–free equilibrium of host–1.

8. Equilibrium $E_7 = (x_1^{**}, x_2^*, y_1^*, 0)$ exists. This is a disease–free equilibrium of host–2.

9. Equilibrium $E_8 = (x_1^{**}, x_2^{**}, y_1^{**}, y_2^{**})$ exists. This is a co–existence equilibrium i.e., all types of hosts present here.

Here in the equilibrium points some notations are used in the coordinates that represent the following expressions:

$$x_1^* = [k(a - \alpha)/a], x_1^{**} = (A/\mu)x_2^{**} = [k/(\mu k + a)][a - \alpha - (aA/\mu k)],$$

$$y_1^* = [k'(b - \theta)/b], y_1^{**} = B/r, y_2^{**} = [k'/(b + rk')][b - \theta - (bB/rk')].$$

### 3.3. Variational Matrix of the Model Equations

Set the model equations (3)–(6) as

$$dx_1/dt = f(x_1, x_2, y_1, y_2),$$

where the variational matrix $M$ is defined by

$$M = \begin{pmatrix}
\frac{\partial f}{\partial x_1} & \frac{\partial f}{\partial x_2} & \frac{\partial f}{\partial y_1} & \frac{\partial f}{\partial y_2} \\
\frac{\partial g}{\partial x_1} & \frac{\partial g}{\partial x_2} & \frac{\partial g}{\partial y_1} & \frac{\partial g}{\partial y_2} \\
\frac{\partial h}{\partial x_1} & \frac{\partial h}{\partial x_2} & \frac{\partial h}{\partial y_1} & \frac{\partial h}{\partial y_2} \\
\frac{\partial k}{\partial x_1} & \frac{\partial k}{\partial x_2} & \frac{\partial k}{\partial y_1} & \frac{\partial k}{\partial y_2}
\end{pmatrix}.$$
\[ M_1 = \begin{bmatrix} a - a & 0 & 0 \\ \mu k(a - a) & -A & 0 \\ 0 & 0 & \theta - b - B \end{bmatrix} \]

Its characteristic equation is given by \( \det(M_1 - \lambda I) = 0 \). On solving which the eigenvalues of \( M_1 \) are obtained as \( \lambda_1 = \alpha - a, \lambda_2 = (\mu k(a - a)/a - A), \lambda_3 = (\theta - b), \lambda_4 = [rk'(b - \theta)/b] - B \). Clearly, \( \lambda_1 \) and \( \lambda_2 \) are negative quantities. Also, the values \( \lambda_3 \) and \( \lambda_4 \) are also negative if both the conditions \( \mu k(a - a)/a > 1 \) and \( (\theta - b)/b > 1 \) are satisfied. Hence, the equilibrium point \( E_1 \) is stable under the considered assumption.

Theorem 3: The equilibrium point \( E_2 = (0, 0, y_1^*, y_2^*) \) is stable if \( Q^2 - 4K > 0 \).

Proof: The variational matrix \( M_2 \) at equilibrium point \( E_2 \) is denoted by \( M_1 \) and given by

\[ M_2 = \begin{bmatrix} a - a & 0 & 0 \\ \mu k(a - a) & -A & 0 \\ 0 & 0 & (r k'/b + r k' - b K/rk') \end{bmatrix} \]

Its characteristic equation is given by \( \det(M_2 - \lambda I) = 0 \). By solving the eigenvalues of \( M_2 \) becomes \( \lambda_1 = \alpha - a, \lambda_2 = -(\mu k(a - a)/a - A), \lambda_3 = (\theta - b)/b \). Clearly, \( \lambda_1 \) and \( \lambda_2 \) are negative quantities. Also, the values \( \lambda_3 \) and \( \lambda_4 \) are also negative if both the conditions \( Q^2 - 4K > 0 \) and \( K > 0 \) hold true simultaneously.

Thus, equilibrium point \( E_2 \) is stable under the considered condition.

Theorem 4: The equilibrium point \( E_3 = (x_1^*, x_2^*, 0, 0) \) is stable if \( b < \theta \) and \( H > 1 \) otherwise unstable.

Proof: Consider
\[ M_3 = \begin{bmatrix} -a/A \mu k & -a(a + k\mu)/\mu \\ (\mu k(a - a)/a - A) & 0 \end{bmatrix} \]

But applying the elementary linear algebra on interchanging column operation this \( M_3 \) is equivalent to;
\[ M_4 = \begin{bmatrix} -A(a + k\mu)/\mu & -a(a + k\mu)/\mu \\ (\mu k(a - a)/a - A) & 0 \end{bmatrix} \]

Now, the variational matrix \( M_3 \) is equivalent denoted by \( M_4 \) and its characteristic equation is given by \( \det(M_3 - \lambda I) = 0 \). This gives the eigenvalues \( \lambda_1 = -A(a + k\mu)/\mu, \lambda_2 = H, \lambda_3 = b - \theta, \lambda_4 = -B \).

Where, \( H = (\mu k(a - a)/a - A) \). Clearly, \( \lambda_1 \) is negative if \( b < \theta \) and \( \mu k(a - a)/a > 1 \). Hence, the equilibrium \( E_3 \) is stable under the stated conditions.

Theorem 5: The equilibrium point \( (0, 0, y_1^*, 0) \) is stable if \( rk'(b - \theta)/b < B \).

Proof: The variational matrix \( M_5 \) at this equilibrium is denoted by \( M_6 \) and given by

\[ M_5 = \begin{bmatrix} a - a & 0 & 0 \\ \mu k(a - a) & -A & 0 \\ 0 & 0 & (r k'/b + r k' - b K/rk') \end{bmatrix} \]

The characteristic equation \( \det(M_5 - \lambda I) = 0 \) of the variation matrix \( M_5 \) at the equilibrium point \( E_6 \) takes the form as \( (a - a - \lambda)(-A - \lambda)(b - \theta - \lambda)(B - \lambda) = 0 \) giving the eigenvalues \( \lambda_1 = a - a, \lambda_2 = -A, \lambda_3 = \theta - b, \lambda_4 = r k'(b - \theta)/b - B \). Here, \( \lambda_2 < 0 \) unconditionally. However, the remaining eigenvalues are conditionally negatives with the conditions

(i) \( \lambda_1 < 0 \) if \( a < a \)
(ii) \( \lambda_3 < 0 \) if \( \theta < b \)
(iii) \( \lambda_4 < 0 \) if \( rk'(b - \theta)/b < B \).

Here, under the considered condition the equilibrium point \( E_6 \) is stable.

Theorem 6: The equilibrium point \( E_7 = (x_1^*, 0, 0, 0) \) is stable if \( [\mu k(a - a)/a] < A \).

Proof: The characteristic equation \( \det(M_7 - \lambda I) = 0 \) of the variation matrix \( M_7 \) at the equilibrium point \( E_8 \) takes the form as \( (a - a - \lambda)(-A - \lambda)(b - \theta - \lambda)(-B - \lambda) = 0 \) giving the eigenvalues \( \lambda_1 = a - a, \lambda_2 = -A, \lambda_3 = b - \theta, \lambda_4 = -B \). Here, it can be observed that \( \lambda_4 \) is unconditionally negative while the other eigenvalues are negatives conditionally with the following conditions:

(i) \( \lambda_1 < 0 \) if \( a < a \)
(ii) \( \lambda_2 < 0 \) if \( \mu k(a - a)/a < A \)
(iii) \( \lambda_3 < 0 \) if \( b < \theta \).

3.4.2. Global Stability of the System

To determine the global stability of the system Lyapunov function method is followed in this study.

Theorem 7: The interior equilibrium point
\((x_1^*, x_2^*, y_1^*, y_2^*)\) is globally stable.

**Proof:**

Construct the Lyapunov function as

\[
V(x_1, x_2, y_1, y_2) = x_1 - x_1^* - x_1^* \ln(x_1/x_1^*) + l[x_2 - x_2^* - x_2^* \ln(x_2/x_2^*)] + m[y_1 - y_1^* - y_1^* \ln(y_1/y_1^*)] + s[y_2 - y_2^* - y_2^* \ln(y_2/y_2^*)].
\]

Where \(m, l, s\) are chosen constants.

Now, the differentiation of \(V\) with respect to \(t\) and after some algebraic manipulation reduces to the form as follow;

\[
\frac{dV}{dt} = \left(\frac{x_1 - x_1^*}{x_1}\right) \left(\frac{dx_1}{dt}\right) + \left(\frac{x_2^* - x_2}{x_2}\right) \left(\frac{dx_2}{dt}\right) + m \left(\frac{y_1 - y_1^*}{y_1}\right) \left(\frac{dy_1}{dt}\right) + s \left(\frac{y_2 - y_2^*}{y_2}\right) \left(\frac{dy_2}{dt}\right) + \frac{x_1 - x_1^*}{x_1} a x_1 \left(1 - \frac{x_1 + x_2}{k}\right) - \mu(x_1 x_2) - \alpha x_1 + l \left(\frac{x_2 - x_2^*}{x_2}\right) \mu(x_1 x_2) - A x_2 + m \left(\frac{y_1 - y_1^*}{y_1}\right) b y_1 \left(1 - \frac{y_1 + y_2}{k'}\right) - r(y_1 y_2) - \theta y_1
\]

For simplicity, choosing \(l = 1\) and \(m = s\) the foregoing time derivative reduces to \(dV/dt = -(a/k)(x_1 - x_1^*)^2 - (bm/k')(y_1 - y_1^*)^2\). Now, it can be observed that \(dV/dt < 0\).

Therefore, the interior equilibrium point is globally stable.

### 3.5. Numerical Simulations

In this section, numerical simulations of the dynamics of the model equation around some of the steady state for certain range of parametric values are used to support the analytical results.

The parameters and its values used in this study are mentioned in the following table as follow;

| Figures | Parameters and their values |
|---------|-----------------------------|
| \(a\) | \(b\) | \(\mu\) | \(r\) | \(\alpha\) | \(\theta\) | \(\delta\) | \(\beta\) | \(k\) | \(k'\) |
| 3 | 1.1940 | 0.2550 | 0.3900 | 3.3920 | 2.4150 | 0.6000 | 2.1600 | 1.0900 | 0.6250 | 2.3920 |
| 4 | 3.1260 | 3.9550 | 0.1600 | 6.2000 | 3.1050 | 0.4800 | 15.2200 | 1.6470 | 0.5450 | 6.9200 |
| 5 | 3.0660 | 2.6200 | 6.0300 | 4.1440 | 8.0700 | 0.4800 | 11.9400 | 2.1960 | 2.6350 | 7.6400 |
| 6 | 5.5380 | 0.8700 | 8.4600 | 1.1040 | 1.7250 | 5.3820 | 1.0400 | 6.0210 | 3.5100 | 1.6240 |
| 7 | 5.5380 | 1.1250 | 8.4600 | 5.9060 | 1.7250 | 5.2680 | 4.4000 | 7.9020 | 3.5100 | 6.0160 |
| 8 | 5.4960 | 1.4950 | 0.2300 | 1.8240 | 0.5850 | 0.2520 | 2.4200 | 3.9060 | 1.2500 | 2.8320 |
| 9 | 5.4960 | 2.5400 | 0.2300 | 3.1600 | 0.5850 | 0.2700 | 2.4200 | 8.0730 | 1.2500 | 5.0640 |

**Figure 3.** All populations vanish together over time.
This figure shows that when growth rate of both populations is less than the natural death rate and attacked by disease all the populations become completely extinct.

Figure 4. The populations of the second host exist with disease.

This figure shows that the growth rate of host-2 is greater than its natural death rate and the first host more affected and hence declines more.

Figure 5. The population of the second host dominant over the other.

This figure shows that the infected host declines more due to the infection rate and natural death rate of host-1 is increased keeping constant death rate of host-2.
One can observe from this figure that host-2 is more affected by disease and die due to natural death. But once the populations of the infected starts to decline, the first host become slightly increasing and regulate each other by forcing the second host to extinct without recovery.

This figure shows that host-2 is more infected and hence the mortality rate due to disease increases while the first host keeps increasing up to certain time.
This figure shows that both host populations would live together without harming each other when the infected hosts are decline and extinct without recovery.

This figure shows that when infection transmission rate of host-1 and mortality rate due to disease of host-2 increased keeping the other parameters constant the populations become stabilized.

4. Conclusion

In the present study the population dynamics of four dimensional systems has been taken up for investigation. A
The mathematical model for this four dimensional system is developed by studying carefully and incorporating the futures of each system together. The positivity and boundedness of the model variables are verified and hence shown that the developed four-dimensional system is biologically well behaved.

In this paper, species have strong indirect effects on others, and predicting these effects is a central challenge in ecology. Host species sharing an enemy can be linked by apparent competition, that is interaction between two hosts and parasite is carried out for investigation. In the case where infection was permitted, parasite coexists in the population, but only under particular conditions, namely when rates of infection were varies both hosts.

The results from the model system demonstrate that the important role that parasite may play in the establishment of community structures within host populations’ dynamics. This was indicated from the stability of interior equilibrium point in which the populations are coexisting.

We need to point out here that, although the model considered in this study is two host populations via shared enemy model. Even though there is no direct interactions specifically associated with the hosts populations, the effects of each host was investigated by altering parameters involved in the system. It has been shown that, this process is strong enough to be a community-wide structuring mechanism that could be used to predict future states of diversification. To show this, numerical study of the model was carried out to support the finding. Moreover, the model system would be fully screened when there is a competition between the hosts that would be mediated by parasite interaction providing shared resources. This would be our next further investigation.

Acknowledgements

The authors would like to thank the editors and anonymous referee for their valuable comments and suggestions which led to a significant improvement of the work.

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