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Stability and Hopf bifurcation dynamics of a food chain system: plant–pest–natural enemy with dual gestation delay as a biological control strategy

Vijay Kumar1,2 · Joydip Dhar3 · Harbax Singh Bhatti⁴

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Abstract

In this paper, a food chain system with gestation delay for both pest and the natural enemy is proposed. Here the boundedness of the system is studied. Stability analysis for all possible equilibrium points is carried out. The thresholds for Hopf bifurcation at interior and the natural enemy free equilibrium states are studied and analyzed. It is observed that the natural enemy free steady state is stable if the gestation delay for the pest is sufficiently low otherwise system observed oscillating behavior. Similar observations established for the interior equilibrium. The sensitivity analysis is performed to find the respective sensitive indices of the variables of the proposed system. Further, simulations have been carried out to support our analytic results.

Keywords A food chain model · Gestation delay · Boundedness · Hopf bifurcation · Sensitivity analysis

Mathematics Subject Classification 34C23 · 34D20 · 92B05 · 92D30

Introduction

Since pest species are harmful to plants and their control has become a challenge for us. Pest population is responsible for severe environmental and realistic problems [\[11,](#page-7-0) [26](#page-8-0)]. Also, many authors have discussed the models based on chemical pesticides, which are less harmful to humanity

 \boxtimes Vijay Kumar bhagat21@rediffmail.com Joydip Dhar jdhar.iiitmg@gmail.com

> Harbax Singh Bhatti bhattihs100@yahoo.com

¹ IKG-Punjab Technical University, Kapurthala 144601, India

² Department of Applied Sciences Humanities and Management, Beant College of Engineering and Technology, Gurdaspur, Punjab 143521, India

- ³ Department of Applied Sciences, ABV-Indian Institute of Information Technology and Management, Gwalior, MP 474015, India
- ⁴ Department of Applied Sciences, B.B.S.B. Engineering College, Fatehgarh Sahib, Punjab, India

and environment [\[5](#page-7-1), [9](#page-7-2), [16,](#page-7-3) [17,](#page-7-4) [27\]](#page-8-1). For productive use of biological or natural methods to manage pest populations, without any adverse effects, it is important to understand the biology of beneficial species or natural enemy and pests [\[8](#page-7-5)]. Our most important aim is to control negative impacts of agriculture pests, for both humanity and agriculture, which harms the environment and generating different types of pollution. The irrigation and emissions from the paddy field were the most environmentally burdening stages across all major impact categories [[10\]](#page-7-6). Moreover, they have shown that the manufacture of fertilizer and pesticide also play a significant role in putting environmental load. Researchers must have to produce, the natural systems to control pests by taking into account the communications between solid Allee effect in pests with natural methods: alternative food support for the natural enemy, introduction of infected pests to control healthy pests [[7](#page-7-7), [18](#page-7-8)] The interactions between pests and natural enemies in the same biological environment is an ample exciting area of research as per Lotka and Volterra. Natural enemies are more vulnerable to the infected pest since infectious pest population is weak and less active. Therefore natural enemy efficiently harvests pests. Due to the interaction between infected pests and natural enemy, the natural enemies must be infected. Hence

natural enemy populations may live on other food resources for their growth and survival. Also, the species do not grow instantaneously; some time is taken by the species to give a new generation, called gestation lag period [[20](#page-7-9)]. Functional responses play an important role to develop a predator-prey system in population dynamics. Various factors like hiding technique of pests from the natural enemy, shooting ability of the predator to harvest insect, etc., have a large influence on functional responses. Functional responses are of different types: for example, Holling type I–III, etc. A mathematical model has studied and analyzed to study the effect of toxicant in a three-species food-chain system incorporating delay in toxicant uptake process by prey population [[14](#page-7-10)]. They formulated the model using the system of nonlinear ordinary differential equations. Also, people are more conscious and choose, the modern methods to manage agricultural pests, for example, less harmful chemical pesticides and natural techniques [\[2](#page-7-11), [6](#page-7-12), [21,](#page-7-13) [28,](#page-8-2) [29\]](#page-8-3), whereas biological techniques are simple and safer to control pests than pesticide practices. Also time lag factors are of great significance to produce population models and used by numerous authors [\[1](#page-7-14), [3](#page-7-15), [4](#page-7-16), [12](#page-7-17), [13](#page-7-18), [15](#page-7-19), [19](#page-7-20), [23–](#page-8-4)[25](#page-8-5)]. According to many authors, models with continuous lag factors are practical [[15](#page-7-19)] than instant delays [[13\]](#page-7-18).

Keeping in mind the recent literature, in the present study, the dynamics of a food chain model with the gestation delay for both pest the natural enemy is proposed and analyzed. This paper is organized as follows: Sect. [1,](#page-0-0) consists of an introduction. The proposed modeling approach and the mathematical system is presented, in Sects. [2](#page-1-0) and [3](#page-1-1) respectively. In Sect. [4,](#page-2-0) the boundedness of the system has been given and discussed. Equilibrium points and their stability analysis is carried out for possible steady states, in Sect. [5.](#page-2-1) The sensitivity analysis of the system at interior equilibrium point for system parameters is presented, in Sect. [6.](#page-5-0) In Sect. [7,](#page-6-0) numerical simulations are presented to support our analytic results. Finally, the results have been concluded in the last section.

Modelling approach

In this paper, we propose a food chain dynamics of plant–pest–natural enemies, keeping in view that the natural systems to control pests. The biological dynamics is shown in Fig. [1.](#page-1-2) We will use the compartmental modeling approach considering three compartments of the population, namely, plant, pest, and natural enemy.

Further, in the absence of pest, the particular type of plant grows logistically, and the pest has alternative food for survival, details modeling assumptions are stated in the next section.

The proposed mathematical system

The assumptions of the proposed model are as follows:

- (1) In a particular habitat, there are three types of populations, namely, plant $X(t)$, pest $P(t)$ and natural enemy *N*(*t*).
- (2) Plants grow logistically with α as intrinsic growth rate and *k* being carrying capacity. Thus, per capita growth rate for plants is $\alpha X \left(1 - \frac{X}{k}\right)$) , when system is free from pest species.
- (3) Also, the pest species grow logistically with α_1 as intrinsic growth rate and k_1 being carrying capacity. Thus, per capita growth rate for pests is $\alpha_1 P \left(1 - \frac{P}{k_1}\right)$) .
- (4) Plants are harvested by pests with Holling type-I, response function.
- (5) Pests can hide from the natural enemy, hence the natural enemy harvesting pests with Holling type-II response function.
- (6) Let β be the predation rate of the plant by pest; β_1 is the conversion rate for pest; γ is the harvesting rate of pests by the natural enemy. Let *a* be the half-saturation

Fig. 1 Biological dynamics

constant. Let γ_1 be the conversion rate for the natural enemy; μ be the natural death rate of natural enemy.

(7) Finally, τ_1 and τ_2 are the gestation delays for the pest and the natural enemy.

Keeping in view the assumptions and interactions, the schematic flow of the proposed dynamics shown in Fig. [2](#page-2-2). Hence our proposed dynamics can express as a system of equations of the form:

$$
\frac{dX}{dt} = \alpha X \left(1 - \frac{X}{k} \right) - \beta X P(t - \tau_1),\tag{1}
$$

$$
\frac{dP}{dt} = \alpha_1 P \left(1 - \frac{P}{k_1} \right) + \beta_1 XP - \frac{\gamma PN(t - \tau_2)}{a + P},\tag{2}
$$

$$
\frac{dN}{dt} = \frac{\gamma_1 P N}{a + P} - \mu N,\tag{3}
$$

with initial conditions: $X(0) > 0$, $P(0) > 0$ and $N(0) > 0$.

Boundedness

Here, the boundedness of solution of the system (1) (1) – (3) (3) is discussed below:

Lemma 1 *The solution of proposed model* [\(1\)](#page-2-3)–([3\)](#page-2-4) *is uniformly bounded in* Ω, *where*

$$
\Omega = \left\{ (X, P, N) : 0 \le X(t) + P(t) + N(t) \le W_0 \right\},\
$$

$$
\mu' = \min\{\mu, -\alpha, -\alpha_1\}, \ \gamma_1 << \gamma, \ \beta_1 << \beta, \ W_0 = e^{-\mu' t + c}.
$$

Proof Let $W(t) = X(t) + P(t) + N(t)$. Now, differentiating *W*(*t*) w.r.t. *t*, we have

$$
\frac{dW(t)}{dt} = \alpha X \left(1 - \frac{X}{k}\right) - \beta X P + \alpha_1 P \left(1 - \frac{P}{k_1}\right) + \beta_1 X P - \frac{\gamma P N}{a + P} + \frac{\gamma_1 P N}{a + P} - \mu N.
$$

Since $\mu' = min\{\mu, -\alpha, -\alpha_1\}, \ \beta_1 << \beta, \ \gamma_1 << \gamma$, we have

$$
\frac{dW(t)}{dt} \le -\mu'W - \frac{\alpha X^2}{k} - \frac{\alpha_1 P^2}{k_1}
$$

$$
\frac{dW(t)}{dt} + \mu'W \le 0.
$$

Therefore, $W = W_0 = e^{-\mu' t + c}$ Hence, $W(t)$ is bounded, i.e., the proposed system is bounded. $□$

Theorem 1 *The local behavior of different equilibrium points of the system* ([1\)](#page-2-3)–([3\)](#page-2-4) *is as follows:*

- (i) *The equilibrium point* $E_0(0, 0, 0)$ *is always exist and unstable*.
- (ii) *The equilibrium point* $E_1(k, 0, 0)$ *exists and unstable.*

Proof

(i) The characteristic equation for $E_0(0, 0, 0)$ is

$$
(-\lambda + \alpha)(-\lambda + \alpha_1)(-\lambda - \mu) = 0.
$$
 (5)

Fig. 2 Schematic diagram

Equilibria and their stability analysis

The system of Eqs. (1) (1) – (3) (3) (3) have four feasible equilibrium points:

- (1) The equilibrium point $E_0(0, 0, 0)$ always exists.
- (2) The equilibrium point $E_1(k, 0, 0)$ exists.
- (3) The natural enemy free equilibrium $E_2(X_2, P_2, 0)$ exists only when $(H_1) := \alpha > \beta k_1$ holds, where $X_2 = \frac{k(\alpha - \beta k_1)\alpha_1}{\alpha\alpha_1 + k k_1 \beta \beta_1}$

$$
P_2 = \frac{k_1 \alpha(\alpha_1 + k \beta_1)}{\alpha \alpha_1 + k k_1 \beta \beta_1}.
$$

(4) Interior equilibrium *E*∗(*X*[∗], *P*[∗], *N*∗) exists, when

$$
\gamma_1 > max \bigg\{ \mu, \frac{\mu(\alpha + a\beta)}{\alpha}, \frac{\alpha\alpha_1\mu(a+k) + (\alpha + a\beta)(kk_1\beta_1)}{\alpha k(\alpha_1 + k_1\beta_1)} \bigg\},
$$

where *X*[∗], *P*[∗], *N*[∗] are given by

$$
\begin{cases} X^* = k + \frac{ak\beta\mu}{\alpha(\mu - \gamma_1)}, \\ P^* = \frac{a\mu}{\gamma_1 - \mu}, \\ N^* = -\frac{a\gamma_1(\alpha\alpha_1(a\mu + k_1(\mu - \gamma_1))+kk_1\beta_1((\alpha + a\beta)\mu - a\gamma_1)}{\alpha\gamma k_1(\mu - \gamma_1)^2}.\end{cases}
$$
(4)

Here, the characteristic roots are $\lambda = \alpha$, $\lambda = \alpha_1$, $\lambda = -\mu$. The equilibrium $E_0(0, 0, 0)$ is always unstable, since two of the characteristic roots, i.e., $\lambda = \alpha$ and $\lambda = \alpha_1$ of [\(5](#page-2-5)) are positive.

(ii) The characteristic equation for $E_1(k, 0, 0)$ is

$$
(-\lambda - \alpha)(-\lambda + (k\beta_1 + \alpha_1))(-\lambda - \mu) = 0.
$$
 (6)

The characteristic roots are $\lambda = -\alpha$, $\lambda = k\beta_1 + \alpha_1$, $\lambda = -\mu$. Hence, the equilibrium point $E_1(k, 0, 0)$ is unstable because one of the eigen value, i.e., $\lambda = k\beta_1 + \alpha_1$ of Eq. [\(6](#page-3-0)) is positive.

Now, we state a lemma as similar to [[22](#page-7-21)]:

Lemma 2 *For the polynomial of the form,* $z^3 + pz^2 + qz + r = 0$,

□

- (i) *Ifr <* 0*, then the equation has at least one non negative root;*
- (ii) *If* $r \ge 0$ *and* $\triangle = p^2 3q \le 0$, *the equation has no non-negative value;*
- (iii) *If* $r \ge 0$ *and* $\triangle = p^2 3q > 0$, *the equation has non-negative roots if and only if* $z_1^* = \frac{-p + \sqrt{\Delta}}{3}$ *and* $h(z_1^*) \leq 0$, *where* $h(z) = z^3 + pz^2 + qz + r$.

Theorem 2 *Let* (H_2) *holds. For the system* (1) (1) – (3) (3) ,

- (i) *The natural enemy free equilibrium* $E_2(X_2, P_2, 0)$ *is locally asymptotically stable for all* $\tau_1 \in [0, \tau_{10}^+)$.
- (ii) If $\tau_1 \ge \tau_{10}^+$ then the equilibrium $E_2(X_2, P_2, 0)$ is unsta*ble and undergoes Hopf bifurcation*.

Proof The characteristic equation of the jacobian matrix at $E₂$ can be written as:

$$
(\lambda^3 + A\lambda^2 + B\lambda + C) + (E\lambda + D)e^{-\lambda \tau_1} = 0,\t(7)
$$

where $A = -a_1 - a_4 - a_6$, $B = a_6(a_1 + a_4) + a_1a_4$, $C =$ $-a_1 a_4 a_6$, $E = -a_2 a_3$, $D = a_2 a_3 a_6$ and $a_1 = -\frac{2X_2 a_6}{k} + \alpha - P_2 \beta$, $a_2 = -X_2\beta, a_3 = P_2\beta_1, a_4 = \alpha_1 - \frac{2P_2\alpha_1}{k_1} + X_2\beta_1, a_5 = -\frac{P_2\gamma}{a+P_2},$ $a_6 = -\mu + \frac{P_2 \gamma_1}{a + P_2}.$

In the absence of delay ($\tau_1 = 0$), the transcendental Eq. [\(7](#page-3-1)) reduces to

$$
\lambda^3 + A\lambda^2 + (B + E)\lambda + (C + D) = 0,\tag{8}
$$

where $A = -a_1 - a_4 - a_6$, $B + E = a_6(a_1 + a_4) + a_1a_4 - a_2a_3$, $C + D = -a_1a_4a_6 + a_2a_3a_6$. By Routh–Hurwitz criterion, all the roots of Eq. [\(8\)](#page-3-2) have negative real parts and the equilibrium E_2 is locally asymptotically stable if (H_2) : A , $(B + E)$,

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 $C + D > 0$ and $A(B + E) - (C + D) > 0$ holds. Assume that $\lambda = iw$ is root of [\(7](#page-3-1)), therefore we have

$$
(iw)^3 + A(iw)^2 + B(iw) + C + ((E(iw) + D)e^{-iw\tau_1} = 0.
$$
 (9)

Equating real and imaginary parts from ([9\)](#page-3-3), it can be obtained

$$
Ew\sin w\tau_1 + D\cos w\tau_1 = Aw^2 - C,\tag{10}
$$

$$
Ew\cos w\tau_1 - D\sin w\tau_1 = w^3 - Bw.\tag{11}
$$

Solving (10) (10) and (11) (11) , we get

$$
w^6 + pw^4 + qw^2 + r = 0,
$$
\n(12)

where

$$
p = A2 - 2B,q = B2 - 2(AC) - E2,r = C2 - D2.
$$

By substituting $w^2 = z$ in Eq. ([12](#page-3-6)), we define

$$
F(z) = z^3 + pz^2 + qz + r.
$$

By Lemma [2](#page-3-7), there exists at least one positive root $w = w_0$ of Eq. (12) (12) satisfying (10) (10) and (11) (11) , which implies that Eq. ([7\)](#page-3-1) has a pair of purely imaginary roots $\pm iw_0$. Solving ([10\)](#page-3-4) and [\(11](#page-3-5)) for τ_1 and substituting the value of $w = w_0$, the corresponding $\tau_{1k} > 0$ is given by

$$
\tau_{1k}^+ = \frac{1}{w_0} \left[\cos^{-1} \left(\frac{E w_0^4 + (AD - BE) w_0^2 - CD}{E^2 w_0^2 + D^2 \right)^2} \right) + 2k\pi \right],
$$

where, *k* is a positive integer. Since the existence of Hopf bifurcation at τ_{10}^+ , it is required that the transversality condition $Re\left[\left(\frac{d\lambda}{d\tau_1}\right)\right]$ \setminus ⁻¹] $\tau_1 = \tau_{10}^+$ 10 \neq 0 should hold, therefore taking the

derivative of λ with respect to τ_1 in ([7\)](#page-3-1), we get

$$
\left(\frac{d\lambda}{d\tau_1}\right)^{-1} = \frac{(3\lambda^2 + 2A\lambda + B)e^{\lambda\tau_1} + E}{\lambda(E\lambda + D)} - \frac{\tau_1}{\lambda}.
$$

At $\lambda = iw_0$ and $\tau_1 = \tau_{10}^+$, we have

$$
Re\left[\left(\frac{d\lambda}{d\tau_1}\right)^{-1}\right] = \frac{MQ - NR}{w_0(L^2 + M^2)},
$$

where $K = -3w_0^2 + B$, $L = 2Aw_0$, $M = D$, $N = Ew_0$, $Q = K \sin w_0 \tau_{10} + L \cos w_0 \tau_{10}$ and $R = K \cos w_0 \tau_{10} -$ *L* sin $w_0 \tau_{10} + E$.

Now, we have,

$$
Re\left[\left(\frac{d\lambda}{d\tau_1}\right)^{-1}\right]_{\tau_1=\tau_{10}^+}\neq 0, \text{ if }MQ \neq NR.
$$

Theorem 3 *Let* (H_3) *holds. For the system* (1) (1) – (3) (3) ,

(i) *The interior equilibrium* $E^*(X^*, P^*, N^*)$ *is locally asymptotically stable for all* $\tau_1 \in [0, \tau_{10}^+)$.

□

(ii) If $\tau_1 \ge \tau_{10}^+$, then the equilibrium $E^*(X^*, P^*, N^*)$ is *unstable and undergoes Hopf bifurcation*.

Proof The characteristic equation of the jacobian matrix at *E*∗ can be written as:

$$
(\lambda^3 + A_2 \lambda^2 + A_1 \lambda + A_0) + (B_1 \lambda + B_0)e^{-\lambda \tau_1} + (C_1 \lambda + C_0)e^{-\lambda \tau_2} = 0,
$$
\n(13)

where $A_2 = -(b_1 + b_4 + b_7)$, $A_1 = b_1b_4 + b_1b_7 + b_4b_7$, $A_0 = -b_1b_4b_7$, $B_1 = -b_2b_3$, $B_0 = b_2b_3b_7$, $C_1 = -b_5b_6$, $C_0 = b_1 b_5 b_6$ and

$$
b_1 = -\frac{X^* \alpha}{k} + (1 - \frac{X^*}{k})\alpha - P^* \beta, \quad b_2 = -X^* \beta, \quad b_3 = P^* \beta_1, \n b_4 = \frac{P^* N^* Y}{(a+P^*)^2} - \frac{N^* Y}{a+P^*} + (1 - \frac{P^*}{k_1})\alpha_1 - \frac{P^* \alpha_1}{k_1} + X^* \beta_1, b_5 = -\frac{P^* Y}{a+P^*}, \n b_6 = -\frac{P^* N^* Y_1}{(a+P^*)^2} + \frac{N^* Y_1}{a+P^*}, b_7 = -\mu + \frac{P^* Y_1}{a+P^*}.
$$

In the absence of delay $\tau_1 = 0$ and $\tau_2 = 0$, the transcendental Eq. (13) (13) reduces to

$$
\lambda^3 + A_2 \lambda^2 + (A_1 + B_1 + C_1)\lambda + (A_0 + B_0 + C_0) = 0, \quad (14)
$$

By Routh–Hurwitz criterion, we know that if (H_3) : $A_0 + B_0 + C_0 > 0, A_2(A_1 + B_1 + C_1) > A_0 + B_0 + C_0$ holds, then all the roots of Eq. (14) (14) have negative real parts and the equilibrium *E*[∗] is locally asymptotically stable. Obviously, $iv(\tau_1) v > 0$ is a root of Eq. [\(13\)](#page-4-0) with $\tau_2 = 0$ if and only if $-iv^3 - A_2v^2 + (A_1 + C_1)v^2 + A_0 + C_0 + (iB_1v + B_0)(\cos v\tau_1)$ $-i \sin v\tau_1$ = 0 [\[30](#page-8-6)]. On separating real and imaginary parts from above equation, we have

$$
\begin{cases}\n-v^3 + (A_1 + C_1)v = B_0 \sin v \tau_1 - B_1 v \cos v \tau_1, \\
A_2 v - (A_0 + C_0) = B_0 \cos v \tau_1 + B_1 v \sin v \tau_1,\n\end{cases}
$$
\n(15)

which gives us

$$
v^6 + p_1 v^4 + q_1 v^2 + r_1 = 0,\t\t(16)
$$

Table 1 The sensitive indices $\gamma_{y_u}^{x_v} = \frac{\partial x_v}{\partial y_u} \times \frac{y_u}{x_v}$ of the model [\(1](#page-2-3))–[\(3\)](#page-2-4) to the parameters y_u for the parameter values: $\alpha = 1.1$; $k = 2$; $\beta = 0.05$; $\alpha_1 = 1.6$; $k_1 = 3$; $\beta_1 = 0.01$; $\gamma = 0.5$; $a = 1$; $\gamma_1 = 0.3$; $\mu = 0.2$

Parameter (y_u)	$\gamma^{X^*}_{y_u}$	P^* y_u	γ^{N^*} y_u
α	0.1	0	0.0032967
\boldsymbol{k}	1	0	0.032967
ß	-0.1	0	-0.0032967
α_1	0	0	0.967033
k ₁	0	0	1.93407
β_1	0	0	0.032967
γ	0	0	-1
a	-0.1	1	-0.937363
γ_1	0.3	- 3	3.81209
μ	-0.3	3	-3.81209

where

$$
p_1 = A_2^2 - 2(A_1 + C_1),
$$

\n
$$
q_1 = (A_1 + C_1)^2 - 2A_2(A_0 + C_0) - B_1^2,
$$

\n
$$
r_1 = (A_0 + C_0)^2 - B_0^2.
$$

Let $v^2 = y$, then Eq. ([16\)](#page-4-2) becomes,

$$
F(z) = y^3 + p_1 y^2 + q_1 y + r_1 = 0.
$$

By using Lemma [2](#page-3-7) and proceeding like above Theorem [\(2](#page-3-8)), i.e., to avoid the repetition of mathematical calculations, we get the required existence condition of Hopf bifurcation for equilibrium point *E*^{*} at τ_{10}^+ , we see that if $\tau_1 \ge \tau_{10}^+$, then the equilibrium $E^*(X^*, P^*, N^*)$ is unstable and undergoes Hopf bifurcation. \Box

2.5 X(t) 2 $P(t)$ \cdot N(t) Solution X, P, N Solution X, P, N 1.5 1 0.5 0 10 20 30 40 50 time t

Fig. 3 The natural enemy free equilibrium $E_2(1.24, 2.44, 0)$ is stable for parameter values: $\alpha = 4.5$; $k = 2$; $\beta = 0.7$; $\alpha_1 = 0.6$; $k_1 = 1.2$; $\beta_1 = 0.5; \gamma = 0.05; a = 1; \gamma_1 = 0.044; \mu = 0.05$

Fig. 4 The natural enemy free equilibrium $E_2(2.26, 7.81, 0)$ is stable for the parametric values: $\alpha = 1.8; k = 17;$ $\beta = 0.2; \alpha_1 = 0.3; k_1 = 6;$ $\beta_1 = 0.04; \gamma = 0.6; a = 0.05;$ $\gamma_1 = 0.3; \mu = 0.001;$ $\tau_1 = 8.3 < \tau_{10}^+ = 9.25$

Fig. 5 The natural enemy free equilibrium $E_2(2.26, 7.81, 0)$ is unstable and Hopf bifurcation appears for the parametric values: $\alpha = 1.8$; $k = 17; \beta = 0.2; \alpha_1 = 0.3;$ $k_1 = 6$; $\beta_1 = 0.04$; $\gamma = 0.6$; $a = 0.05; \gamma_1 = 0.3; \mu = 0.001;$ $\tau_1 = 10.25 > \tau_{10}^+ = 9.25$

Sensitivity analysis

In this section, the sensitivity analysis of the system (1) – (3) (3) at the interior equilibrium point is carried out. The respective sensitive parameters of the state variables of the system at interior equilibrium point are given in the Table [1](#page-4-3), using the values of parameters: $\alpha = 1.1$; $k = 2$; $\beta = 0.05$; $\alpha_1 = 1.6$; $k_1 = 3$; $\beta_1 = 0.01$; $\gamma = 0.5$; $a = 1$; $\gamma_1 = 0.3$; $\mu = 0.2$. It is clear that α , k , γ_1 have a positive impact on X^* . Also, the

impact of β , a , μ is negative on X^* , whereas the impact of remaining parameters on X^* is zero. The parameter k is more sensitive to X^* . Also *a*, μ have a positive impact on P^* . The impact of parameter γ_1 on P^* is negative; the remaining parameters have zero impact on *P*[∗]. The more sensitive parameters to P^* are γ_1 and μ . Again, the impact of α , k , α_1 , k_1 , β_1 , γ_1 on N^* is positive. The impact of β , γ , α , μ is negative on N^* . Clearly, γ_1 and μ are more sensitive parameters to N^* .

Fig. 6 The interior equilibrium *E*∗(1.82, 2, 3.31) is stable for parametric values: $\alpha = 1.1$; $k = 2$; $\beta = 0.05$; $\alpha_1 = 1.6$; $k_1 = 3$; $\beta_1 = 0.01$; $\gamma = 0.5; a = 1; \gamma_1 = 0.3; \mu = 0.2$

Numerical simulations

Numerical simulations of the system (1) (1) – (3) are performed to support our analytic findings with the help of MAT-LAB. The natural enemy free equilibrium $E_2(1.24, 2.44, 0)$ is stable for parameter values: $\alpha = 4.5$; $k = 2$; $\beta = 0.7$; $\alpha_1 = 0.6; k_1 = 1.2; \beta_1 = 0.5; \gamma = 0.05; a = 1; \gamma_1 = 0.044;$ $\mu = 0.05$ and result is shown in Fig. [3](#page-4-4). Moreover, the natural enemy free equilibrium $E_2(2.26, 7.81, 0)$ is stable for the parametric values: $\alpha = 1.8$; $k = 17$; $\beta = 0.2$; $\alpha_1 = 0.3$; $k_1 = 6$; $\beta_1 = 0.04$; $\gamma = 0.6$; $a = 0.05$; $\gamma_1 = 0.3$; $\mu = 0.001$; $\tau_1 = 8.3 < \tau_{10}^+ = 9.25$, see Fig. [4](#page-5-1). The natural enemy free equilibrium $E_2(2.26, 7.81, 0)$ is unstable and Hopf

bifurcation appears for the parametric values: $\alpha = 1.8$; $k = 17; \ \beta = 0.2; \ \alpha_1 = 0.3; \ k_1 = 6; \ \beta_1 = 0.04; \ \gamma = 0.6;$ $a = 0.05$; $\gamma_1 = 0.3$; $\mu = 0.001$; $\tau_1 = 10.25 > \tau_{10}^+ = 9.25$ and result is shown in Fig. [5](#page-5-2). The interior equilibrium $E^*(1.82, 2, 3.31)$ is stable for parametric values: $\alpha = 1.1$; $k = 2$; $\beta = 0.05$; $\alpha_1 = 1.6$; $k_1 = 3$; $\beta_1 = 0.01$; $\gamma = 0.5$; $a = 1$; $\gamma_1 = 0.3$; $\mu = 0.2$, see Fig. [6](#page-6-1). It is clear from Fig. [7](#page-6-2) that the interior equilibrium $E[*](1.88, 0.25, 1.23)$ is stable for parametric values: $\alpha = 0.2$; $k = 5$; $\beta = 0.5$; $\alpha_1 = 0.32$; $k_1 = 2$; $\beta_1 = 0.1$; $\gamma = 0.2$; $a = 1$; $\gamma_1 = 0.01$; $\mu = 0.002$; $\tau_1 = 0.6 < \tau_{10}^+ = 1$. It is obvious from Fig. [8](#page-7-22) that the interior equilibrium $E^*(1.88, 0.25, 1.23)$ is unstable and Hopf bifurcation appears for the parametric values: $\alpha = 0.2$; $k = 5; \beta = 0.5; \alpha_1 = 0.32; k_1 = 2; \beta_1 = 0.1; \gamma = 0.2; a = 1;$ $\gamma_1 = 0.01; \mu = 0.002; \tau_1 = 1.5 > \tau_{10}^+ = 1.$

Conclusion

Here, a food chain: a plant–pest–natural enemy system with the gestation delay for pest and the natural enemy is proposed. There are four feasible equilibrium points, and asymptotic stability of the system is studied and analyzed for all equilibria. The steady states $E_0(0, 0, 0)$ and $E_0(K, 0, 0)$ are always unstable. The existence of Hopf bifurcation at the natural enemy free as well as interior equilibrium point is explored and determined the critical limits for gestation delay, τ_1 . It is observed that the natural enemy free steady state is stable if the gestation delay for pest (τ_1) is below a certain threshold otherwise system observed oscillating behavior. A similar oscillating solution exists for the interior steady state. It is studied that the natural enemy free and

Fig. 7 The interior equilibrium *E*[∗](1.88, 0.25, 1.23) is stable for parametric values: $\alpha = 0.2$; $k = 5; \beta = 0.5; \alpha_1 = 0.32;$ $k_1 = 2$; $\beta_1 = 0.1$; $\gamma = 0.2$; $a = 1; \gamma_1 = 0.01; \mu = 0.002;$ $\tau_1 = 0.6 < \tau_{10}^+ = 1$

Fig. 8 The interior equilibrium *E*∗(1.88, 0.25, 1.23) is unstable and Hopf bifurcation appears for the parametric values: $\alpha = 0.2; k = 5; \beta = 0.5;$ $\alpha_1 = 0.32; k_1 = 2; \beta_1 = 0.1;$ $\gamma = 0.2; a = 1; \gamma_1 = 0.01;$ $\mu = 0.002; \tau_1 = 1.5 > \tau_{10}^+ = 1$

interior equilibrium are asymptotically stable under certain conditions. Also, the sensitivity analysis is performed at interior equilibrium point for the system parameters. Numerical simulations of the system are carried out with a particular set of parameter values to verify our analytic results.

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