

## Research Article

# Stability and Sensitive Analysis of a Model with Delay Quorum Sensing

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This paper formulates a delay model characterizing the competition between bacteria and immune system. The center manifold reduction method and the normal form theory due to Faria and Magalhaes are used to compute the normal form of the model, and the stability of two nonhyperbolic equilibria is discussed. Sensitivity analysis suggests that the growth rate of bacteria is the most sensitive parameter of the threshold parameter  $R_0$  and should be targeted in the controlling strategies.

## 1. Introduction

Quorum sensing is a process that enables bacteria to communicate using secreted signaling molecules called *autoinducers* [1]. It makes bacteria regulate their gene expression collectively and control their behaviors on community scale. Quorum sensing was initially observed in the marine bacterium *Vibrio fischeri* about 30 years ago [2, 3]. Now, many other species are observed to exhibit quorum sensing behavior, including major human pathogens such as *Staphylococcus aureus* and *Pseudomonas aeruginosa*. Quorum sensing has received more and more attention (see [4–15] and the references therein) and some models are formulated to investigate its effect on the transmission of disease. Braselton and Waltman [4] formulated the dynamically allocated inhibitor production. Dockery and Keener [5] were devoted to developing and studying an ODE and a PDE mathematical models for quorum sensing in *Pseudomonas aeruginosa* and found that quorum sensing works because of a biochemical switch between two stable steady solutions, one with low levels of autoinducer and one with high levels of autoinducer. Koerber et al. [6] presented a mathematical model for the early stages of the infection process by *Pseudomonas aeruginosa* in burn wounds which accounts for the quorum sensing and the diffusion of signalling molecules in the burn-wound environment, and the effects of important parameters on

the dynamic properties of the model are discussed in detail. They gave some sufficient conditions for the global asymptotic stability of two boundary equilibria which, respectively, correspond to the survival of the *allelochemical* producer species or the susceptible one. Fergola et al. [7] formulated an *allelopathic* competition model in which a distributed delay term simulates quorum sensing which regulates the delay production process of *allelochemicals*, and they proved the unique existence of the positive solution and the stability of biologically meaningful steady-state solutions. Anguige et al. [9] constructed a multiphase mathematical model of quorum sensing in a maturing *Pseudomonas aeruginosa* biofilm to investigate the effect of antiquorum sensing and antibiotic treatments on the *exopolysaccharide* concentration, signal level, bacterial numbers, and biofilm growth rate. The above articles leave out the immune response to the bacterial invasion. However, the immune status of the hosts has a significant impact on the transmission of an infection in a population. Literatures [11–15] employed a quadratic function to describe the quorum sensing of bacteria and formulate some models to characterize the competition between bacteria and the immune system, in which the existence of periodical solution, chaotic motion, and subharmonic bifurcation, the properties of Hopf bifurcation, and the stability of equilibrium et al. were investigated. As a novelty of this paper, a cubic function is used to express quorum sensing. This makes

the model own more general nonlinearity, which results in much wider set of outcomes including the coexistence of multiple positive equilibria and the existence of critical equilibrium with simple zero singularity.

### 2. Model Formulation

We denote by  $X_U(t)$  the concentration of the uninfected target cells,  $X_I(t)$  the concentration of the infected target cells,  $B(t)$  the concentration of the bacteria,  $I_R(t)$  the concentration of the innate cells, and  $I_A(t)$  the concentration of the adaptive cells. The dynamic relations among them are as follows: the uninfected target cells have a natural turnover  $S_U$  and a half-life  $\mu_{X_U}$  and they are infected by bacteria with mass-action term  $\alpha_1 X_U B$ ; the infected target cells are cleared by half-life  $\mu_{X_I}$  or adaptive immune cells with mass action term  $\alpha_2 X_I I_A$ ; both the innate and the adaptive immune cells have a source term and a half-life time; for the innate immunity, the source term  $S_{I_R}$  includes a wide range of cells involved in the first wave of defense of the host such as natural killer cells, *polymorphonuclear cells*, *macrophages*, and *dendritic cells*, and for the adaptive immunity, the source term  $S_{I_A}$  represents the memory cells, derived from a previous infection or vaccination, a zero source means the first infection with this pathogen and there are no memory cells; both of the two kinds of cells are increased by the signals captured by the bacteria load; the bacteria population has a net growth term represented by a logistic function  $\alpha_{20} B(1 - B/\sigma)$  and it is cleared by the innate immunity with mass action term  $\alpha_3 B I_R$ . Here, we use a function

$$\frac{\alpha_{20}}{B_0} B(t)^2 B(t - \tau) \tag{1}$$

to formulize the bacteria that compete with the immune cells at time  $t$ , which receive signal molecules  $\tau$  time units ago.  $B_0$  is a positive constant,  $\alpha_{20}$  is the growth rate of bacteria, and  $\sigma$  is the effective carrying capacity of the environment. Consequently, the vital dynamics are governed by

$$\begin{aligned} \frac{dB(t)}{dt} &= \alpha_{20} B(t) \left( 1 + \frac{B^2(t - \tau)}{B_0} - \frac{B(t)}{\sigma} \right) \\ &\quad - \alpha_3 B(t) I_R(t) - \alpha_4 B(t) I_A(t), \\ \frac{dX_U(t)}{dt} &= S_U - \alpha_1 X_U(t) B(t) - \mu_{X_U} X_U(t), \\ \frac{dX_I(t)}{dt} &= \alpha_1 X_U(t) B(t) - \alpha_2 I_A(t) X_I(t) - \mu_{X_I} X_I(t), \\ \frac{dI_R(t)}{dt} &= S_{I_R} + \beta_1 B(t) - \mu_{I_R} I_R(t), \\ \frac{dI_A(t)}{dt} &= S_{I_A} + \beta_2 B(t) - \mu_{I_A} I_A(t). \end{aligned} \tag{2}$$

*Remark 1.* The first equation of system (2) suggests that bacteria are controlled and increased by quorum sensing except for their net growth and they are cleared by the innate immune cells. The second equation of system (2) characterizes the dynamics of the uninfected target cells, and the third one reflects the dynamics of the infected target cells. The uninfected target cells are infected by bacteria in mass action law and they have their own constant input flow, and the infected target cells are killed by the adaptive immune cells. The last two equations of system (2) show that each kind of the immune cells has a special source term and their responses are enhanced by the bacteria load. The target cells and the immune cells have their own half-life terms.

### 3. The Existence and Stability of Equilibria

We introduce

$$\begin{aligned} R_0 &= \frac{\alpha_3 S_{I_R}}{\alpha_{20} \mu_{I_R}} + \frac{\alpha_4 S_{I_A}}{\alpha_{20} \mu_{I_A}}, \\ R_1 &= \frac{4\alpha_{20}^2 (1 - R_0)}{B_0 (\alpha_{20}/\sigma + \alpha_3 \beta_1 / \mu_{I_R} + \alpha_4 \beta_2 / \mu_{I_A})^2}. \end{aligned} \tag{3}$$

**Theorem 2.** *System (2) always admits a bacteria-free equilibrium  $E_0 = (0, S_U/\mu_{X_U}, 0, S_{I_R}/\mu_{I_R}, S_{I_A}/\mu_{I_A})$ . If  $R_0 > 1$ , then system (2) admits a unique positive equilibrium  $E_1$ . If  $R_0 = 1$ , then system (2) admits a unique positive equilibrium  $E_2$ . If  $R_0 < 1$  and  $R_1 > 1$ , then system (2) admits no positive equilibrium. If  $R_0 < 1$  and  $R_1 = 1$ , then system (2) admits a unique positive equilibrium  $E_4$ . If  $R_0 < 1$  and  $R_1 < 1$ , then system (2) has two positive equilibria  $E_1$  and  $E_3$ .*

Specifically,

$$\begin{aligned} E_j &= \left( B_j^*, \frac{S_U}{\alpha_1 B_j^* + \mu_{X_U}}, \right. \\ &\quad \left. \frac{\alpha_1 S_U B_j^* \mu_{I_A}}{(\alpha_2 (S_{I_A} + \beta_2 B_4^*) + \mu_{X_I} \mu_{I_A}) (\alpha_1 B_4^* + \mu_{X_U})}, \right. \\ &\quad \left. \frac{S_{I_R} + \beta_1 B_j^*}{\mu_{I_R}}, \frac{S_{I_A} + \beta_2 B_j^*}{\mu_{I_A}} \right), \\ B_{1,3}^* &= \frac{B_0}{2\alpha_{20}} \left( \frac{\alpha_{20}}{\sigma} + \frac{\alpha_3 \beta_1}{\mu_{I_R}} + \frac{\alpha_4 \beta_2}{\mu_{I_A}} \right) \left( 1 \pm \sqrt{1 - R_1} \right), \\ B_2^* &= \frac{B_0}{\alpha_{20}} \left( \frac{\alpha_{20}}{\sigma} + \frac{\alpha_3 \beta_1}{\mu_{I_R}} + \frac{\alpha_4 \beta_2}{\mu_{I_A}} \right), \\ B_4^* &= \frac{B_0}{2\alpha_{20}} \left( \frac{\alpha_{20}}{\sigma} + \frac{\alpha_3 \beta_1}{\mu_{I_R}} + \frac{\alpha_4 \beta_2}{\mu_{I_A}} \right), \end{aligned} \tag{4}$$

$j = 1, 2, 3, 4.$

**Theorem 3.** *If  $R_0 > 1$ , the bacteria-free equilibrium  $E_0$  is asymptotically stable, while if  $R_0 < 1$ ,  $E_0$  is unstable.*

The proofs for Theorems 2 and 3 are trivial, so omit them.

In the sequel, we study the stability of the positive equilibrium  $E_j$ . First of all, we transfer it to the origin and get

$$\frac{dx(t)}{dt} = Mx(t) + Nx(t - \tau) + f(x(t)), \tag{5}$$

where

$$f(x) = \begin{pmatrix} f_1(x) \\ -\alpha_1 x_1 x_2 \\ \alpha_1 x_1 x_2 - \alpha_2 x_3 x_5 \\ 0 \\ 0 \end{pmatrix}, \quad N = \begin{pmatrix} \frac{2\alpha_{20}(B_j^*)^2}{B_0} & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 \end{pmatrix},$$

$$M = \begin{pmatrix} -\frac{\alpha_{20}B_j^*}{\sigma} & 0 & 0 & -\alpha_3 B_j^* & -\alpha_4 B_j^* \\ \frac{-\alpha_1 S_U}{\alpha_1 B_j^* + \mu_{X_U}} & -\mu_{X_U} - \alpha_1 B_j^* & 0 & 0 & 0 \\ \frac{\alpha_1 S_U}{\alpha_1 B_j^* + \mu_{X_U}} & \alpha_1 B_j^* & -\mu_{X_I} - \frac{\alpha_2(S_{I_A} + \beta_2 B_j^*)}{\mu_{I_A}} & 0 & \Theta \\ \beta_1 & 0 & 0 & -\mu_{I_R} & 0 \\ \beta_2 & 0 & 0 & 0 & -\mu_{I_A} \end{pmatrix}, \tag{6}$$

$$f_1(x) = \frac{\alpha_{20}B_j^*}{B_0} (2x_1 + x_1(t - \tau)) x_1(t - \tau) - x_1(\alpha_3 x_4 + \alpha_4 x_5) - \alpha_{20} x_1 \left( \frac{x_1}{\sigma} - \frac{x_1^2(t - \tau)}{B_0} \right),$$

$$\Theta = -\frac{\alpha_1 \alpha_2 S_U \mu_{I_A} B_j^*}{(\mu_{X_U} + \alpha_1 B_j^*)(\mu_{X_I} \mu_{I_A} + \alpha_2(S_{I_A} + \beta_2 B_j^*))}.$$

The characteristic equation of (5) at the origin is

$$\left( \lambda + \mu_{X_I} + \frac{\alpha_2(S_{I_A} + \beta_2 B_j^*)}{\mu_{I_A}} \right) (\lambda + \mu_{X_U} + \alpha_1 B_j^*) \cdot (\lambda^3 + p_1 \lambda^2 + p_2 \lambda + p_3 + (q_1 \lambda^2 + q_2 \lambda + q_3) e^{-\lambda \tau}) = 0, \tag{7}$$

where

$$p_1 = \mu_{I_R} + \mu_{I_A} + \frac{B_j^* \alpha_{20}}{\sigma},$$

$$p_2 = \mu_{I_R} \mu_{I_A} + \frac{(\mu_{I_R} + \mu_{I_A}) B_j^* \alpha_{20}}{\sigma} + \beta_1 \alpha_3 B_j^* + \beta_2 \alpha_4 B_j^*,$$

$$p_3 = \frac{\mu_{I_A} \mu_{I_R} B_j^* \alpha_{20}}{\sigma} + \beta_1 \alpha_3 B_j^* \mu_{I_A} + \beta_2 \alpha_4 \mu_{I_R} B_j^*,$$

$$q_1 = -\frac{2\alpha_{20}(B_j^*)^2}{B_0}, \quad q_2 = -\frac{2\alpha_{20}(B_j^*)^2(\mu_{I_A} + \mu_{I_R})}{B_0},$$

$$q_3 = -\frac{2\alpha_{20}(B_j^*)^2 \mu_{I_A} \mu_{I_R}}{B_0}. \tag{8}$$

Equation (7) has two negative roots  $-\mu_{X_I} - \alpha_2(S_{I_A} + \beta_2 B_j^*)/\mu_{I_A}$ ,  $-\mu_{X_U} - \alpha_1 B_j^*$ , and the other roots can be obtained by solving the following equation:

$$\lambda^3 + p_1 \lambda^2 + p_2 \lambda + p_3 + (q_1 \lambda^2 + q_2 \lambda + q_3) e^{-\lambda \tau} = 0. \tag{9}$$

It can be seen that

$$p_3 + q_3 = \frac{B_j^*}{B_0 \sigma} (B_0 (\alpha_{20} \mu_{I_R} \mu_{I_A} + \beta_1 \alpha_3 \sigma \mu_{I_A} + \beta_2 \alpha_4 \sigma \mu_{I_R}) - 2\alpha_{20} B_j^* \mu_{I_R} \mu_{I_A} \sigma),$$

$$\begin{aligned}
 p_3 + q_3 &< 0, & j = 1, 2; \\
 p_3 + q_3 &> 0, & j = 3; \\
 p_3 + q_3 &= 0, & j = 4.
 \end{aligned}
 \tag{10}$$

**Theorem 4.** For  $\tau = 0$ , both  $E_1$  and  $E_2$  are unstable when they exist.

Obviously,  $E_3$  does not own zero eigenvalue singularity; namely, zero is not a eigenvalue of the Jacobi matrix at  $E_3$ . By using the Routh-Hurwitz stability criterion,  $E_3$  is locally asymptotically stable if  $p_1 + q_1 > 0$ ,  $p_2 + q_2 > 0$ , and  $(p_1 + q_1)(p_2 + q_2) > (p_3 + q_3)$  are satisfied together.

**Theorem 5.** For  $\tau = 0$ ,  $E_3$  is asymptotically stable if  $R_0 < 1$ ,  $R_1 < 1$ , and  $1 - R_0$  are small sufficiently.

*Proof.* Let  $k = 1 - \sqrt{1 - R_1}$  and  $\varepsilon = 1 - R_0$ . From Theorem 2,  $E_3$  exists if and only if  $R_0 < 1$  and  $R_1 < 1$ , which leads to  $0 < 1 - R_0 < 1$  and  $0 < 1 - R_1 < 1$ ; that is,  $0 < \varepsilon < 1$  and  $0 < k < 1$ . By using  $R_1 = 4\alpha_{20}^2\varepsilon/B_0(\alpha_{20}/\sigma + \alpha_3\beta_1/\mu_{I_R} + \alpha_4\beta_2/\mu_{I_A})^2$ , it follows that  $k$  approaches 0 if  $\varepsilon$  does so. Because

$$\begin{aligned}
 \frac{(B_3^*)^2}{B_0} &= \frac{B_3^*}{\sigma} + \frac{\alpha_3}{\alpha_{20}} I_R^* + \frac{\alpha_4}{\alpha_{20}} I_A^* - 1, \\
 I_R^* &= \frac{S_{I_R} + \beta_1 B_3^*}{\mu_{I_R}}, & I_A^* &= \frac{S_{I_A} + \beta_2 B_3^*}{\mu_{I_A}},
 \end{aligned}
 \tag{11}$$

we have

$$\begin{aligned}
 p_1 + q_1 &= 2\alpha_{20}\varepsilon - \left( \frac{\alpha_{20}}{\sigma} + \frac{2\beta_1\alpha_3}{\mu_{I_R}} + \frac{2\beta_2\alpha_4}{\mu_{I_A}} \right) B_3^* + \mu_{I_R} + \mu_{I_A}, \\
 p_2 + q_2 &= 2\alpha_{20}(\mu_{I_R} + \mu_{I_A})\varepsilon + \mu_{I_R}\mu_{I_A} \\
 &\quad + \left( \frac{\alpha_{20}}{\sigma}(\mu_{I_R} + \mu_{I_A}) + \beta_1\alpha_3 + \beta_2\alpha_4 \right. \\
 &\quad \left. + \frac{2\beta_1\alpha_3\mu_{I_A}}{\mu_{I_R}} + \frac{2\beta_2\alpha_4\mu_{I_R}}{\mu_{I_A}} \right) B_3^*, \\
 p_3 + q_3 &= 2\alpha_{20}\mu_{I_R}\mu_{I_A}\varepsilon \\
 &\quad - \left( \frac{\alpha_{20}}{\sigma}\mu_{I_R}\mu_{I_A} + \beta_1\alpha_3\mu_{I_A} + \beta_2\alpha_4\mu_{I_R} \right) B_3^*.
 \end{aligned}
 \tag{12}$$

Note that  $B_3^* = (B_0/2\alpha_{20})(\alpha_{20}/\sigma + \alpha_3\beta_1/\mu_{I_R} + \alpha_4\beta_2/\mu_{I_A})k$ . We have

$$\begin{aligned}
 &(p_1 + q_1)(p_2 + q_2) - (p_3 + q_3) \\
 &= \frac{-B_0}{2\alpha_{20}} \left[ \left( \frac{2\alpha_{20}}{\sigma}(\mu_{I_R} + \mu_{I_A}) + \frac{\beta_1\alpha_3}{\mu_{I_R}}(3\mu_{I_R} + 4\mu_{I_A}) \right. \right. \\
 &\quad \left. \left. + \frac{\beta_2\alpha_4}{\mu_{I_A}} \cdot (3\mu_{I_A} + 4\mu_{I_R}) \right) \varepsilon + \frac{\alpha_{20}B_0}{\sigma^2} \right.
 \end{aligned}$$

$$\begin{aligned}
 &\cdot \left( \frac{\alpha_{20}}{\sigma}(\mu_{I_R} + \mu_{I_A}) + \frac{\beta_1\alpha_3}{\mu_{I_R}}(4\mu_{I_R} + 5\mu_{I_A}) \right. \\
 &\quad \left. + \frac{\beta_2\alpha_4}{\mu_{I_A}}(4\mu_{I_A} + 5\mu_{I_R}) \right) \\
 &+ \frac{B_0}{\sigma} \left( \frac{\beta_1^2\alpha_3^2}{\mu_{I_R}^2}(5\mu_{I_R} + 8\mu_{I_A}) + \frac{13\beta_1\beta_2\alpha_3\alpha_4}{\mu_{I_R}\mu_{I_A}} \right. \\
 &\quad \left. \cdot (\mu_{I_R} + \mu_{I_A}) + \frac{\beta_2^2\alpha_4^2}{\mu_{I_A}^2}(5\mu_{I_A} + 8\mu_{I_R}) \right) \Big] \\
 &\cdot \left( \frac{\alpha_{20}}{\sigma} + \frac{\alpha_3\beta_1}{\mu_{I_R}} + \frac{\alpha_4\beta_2}{\mu_{I_A}} \right) k + (\mu_{I_R} + \mu_{I_A})\varepsilon^2 \\
 &+ \left( \frac{\alpha_{20}B_0}{2\sigma^2}(\mu_{I_R} + \mu_{I_A}) + \frac{\beta_1\alpha_3B_0}{2\sigma\mu_{I_R}}(3\mu_{I_R} + 4\mu_{I_A}) \right. \\
 &\quad \left. + \frac{\beta_2\alpha_4B_0}{2\sigma\mu_{I_A}}(3\mu_{I_A} + 4\mu_{I_R}) + (\mu_{I_R} + \mu_{I_A})^2 \right. \\
 &\quad \left. + \frac{\beta_1^2\alpha_3^2B_0}{\alpha_{20}\mu_{I_R}^2}(\mu_{I_R} + 2\mu_{I_A}) + \frac{\beta_2^2\alpha_4^2B_0}{\alpha_{20}\mu_{I_A}^2}(\mu_{I_A} + 2\mu_{I_R}) \right. \\
 &\quad \left. + \frac{3\beta_1\beta_2\alpha_3\alpha_4B_0}{\alpha_{20}\mu_{I_R}\mu_{I_A}}(\mu_{I_R} + \mu_{I_A}) \right) \varepsilon + (\mu_{I_R} + \mu_{I_A})\mu_{I_R}\mu_{I_A}.
 \end{aligned}
 \tag{13}$$

Seen from above formulae,  $p_1 + q_1$ ,  $p_2 + q_2$ ,  $p_3 + q_3$ , and  $(p_1 + q_1)(p_2 + q_2) - (p_3 + q_3)$  are positive when  $\varepsilon$  is small, which results in the locally asymptotical stability of  $E_3$ .  $\square$

Clearly, the left side of (9) is continuous in  $\tau$  and has roots with positive real parts if and only if it has purely imaginary roots. We will determine whether (9) has purely imaginary roots or not, from which we then will be able to get conditions for all eigenvalues to have negative real parts.

Denote the eigenvalue of the characteristic equation (5) by  $\lambda = \rho(\tau) + i\omega(\tau)$ , where  $\rho(\tau)$ ,  $\omega(\tau)$  continually depend on the delay  $\tau$ . Under the same conditions as Theorem 5, we have  $\rho(0) < 0$ . Since  $\lambda$  is continuous in  $\tau$ , one still has  $\rho(\tau) < 0$  and  $E_3$  remains stable if  $\tau$  is sufficiently small. If there exists a positive value  $\tau_0$  satisfying  $\rho(\tau_0) = 0$ , that is,  $\lambda = i\omega(\tau_0)$  is a purely imaginary root of (9), then  $E_3$  loses its stability and eventually becomes unstable when  $\rho(\tau)$  becomes positive. On the other hand, if such a  $\rho(\tau_0)$  does not exist  $E_3$  is always stable.

Obviously, (9) has a purely imaginary root  $i\omega$ ,  $\omega > 0$ , if and only if

$$-i\omega^3 - p_1\omega^2 + ip_2\omega + p_3 + (-q_1\omega^2 + iq_2\omega + q_3)e^{-i\omega\tau} = 0.
 \tag{14}$$

Separating the real and imaginary parts of (14) and adding up the squares of them lead to

$$z^3 + A_1z^2 + A_2z + A_3 = 0,
 \tag{15}$$

where  $z = \omega^2$ ,  $A_1 = p_1^2 - 2p_2 - q_1^2$ ,  $A_2 = p_2^2 - 2p_1p_3 + 2q_1q_3 - q_2^2$ , and  $A_3 = p_3^2 - q_3^2$ .

It can be verified that  $A_3 < 0$  and (15) has positive roots. Without loss of generality, one assumes that (15) has three positive roots defined by  $z_1, z_2,$  and  $z_3,$  respectively. Then, (14) has three positive roots  $\omega_j = \sqrt{z_j}, j = 1, 2, 3.$  And then, we have

$$\cos(\omega_k \tau) = \frac{(p_1 \omega_k^2 - p_3)(q_3 - q_1 \omega_k^2) + q_2 \omega_k^2 (\omega_k^2 - p_2)}{q_2^2 \omega_k^2 + (q_3 - q_1 \omega_k^2)^2}. \tag{16}$$

Thus, if we denote

$$\tau_k^j = \frac{1}{\omega_k} \arccos \left\{ \frac{(p_1 \omega_k^2 - p_3)(q_3 - q_1 \omega_k^2) + q_2 \omega_k^2 (\omega_k^2 - p_2)}{q_2^2 \omega_k^2 + (q_3 - q_1 \omega_k^2)^2} + \frac{2j\pi}{\omega_k} \right\}, \tag{17}$$

where  $k = 1, 2, 3, j = 0, 1, 2, \dots,$  then  $\pm i\omega_k$  is a pair of purely imaginary roots of (9). Define  $\tau_0 = \min\{\tau_k^0, k = 1, 2, 3\}.$  We have the following.

**Theorem 6.** *Under the same conditions as Theorem 5,  $E_3$  is asymptotically stable for  $\tau \in [0, \tau_0).$*

**Theorem 7.** *If  $3\omega_k^4 + 2A_1\omega_k^2 + A_2 \neq 0,$  system (2) undergoes Hopf bifurcation at the positive equilibrium  $E_3$  when  $\tau = \tau_k^j.$*

### 4. Normal Forms on the Center Manifold

From the discussions in the above section, it can be seen that the Jacobi matrix at  $E_4$  has a uniquely simple zero eigenvalue if  $R_1 = 1$  and  $\tau \neq (p_2 + q_2)/q_3.$  To determine the dynamic properties of  $E_4,$  we have to compute the normal forms on the center manifold. The method used is based on the center manifold reduction and normal form theory due to Faria and Magalhaes; see [16, 17].

By means of  $R_1 = 1,$  it obtains

$$a_2 \alpha_3^2 + a_1 \alpha_3 + a_0 = 0, \tag{18}$$

where

$$\begin{aligned} a_2 &= \alpha_{20} \mu_{I_R} \mu_{I_A}^2 B_0 \beta_1^2 \sigma^2, \\ a_1 &= 2\alpha_{20} \mu_{I_R}^2 \mu_{I_A}^2 \sigma (2\mu_{I_A} \sigma \alpha_{20} S_{I_R} + B_0 \alpha_{20} \beta_1 \mu_{I_A} + \sigma B_0 \beta_1 \beta_2 \alpha_4), \\ a_0 &= \alpha_{20} \mu_{I_R}^3 \mu_{I_A} (B_0 \alpha_{20}^2 \mu_{I_A}^2 + B_0 \alpha_4^2 \beta_2^2 \sigma^2 + 2B_0 \alpha_{20} \mu_{I_A} \alpha_4 \beta_2 \sigma \\ &\quad + 4\alpha_{20} \sigma^2 \mu_{I_A} \alpha_4 S_{I_A} - 4\alpha_{20}^2 \sigma^2 \mu_{I_A}^2). \end{aligned} \tag{19}$$

It is seen from (19) that  $a_0 < 0$  if  $S_{I_A}$  and  $B_0$  are small enough, which means there exists a unique positive  $\alpha_3$  solving (18). Denote it by  $d$  and define  $\alpha_3$  as

$$\alpha_3 = d + \mu, \tag{20}$$

where  $\mu$  is a small parameter. Obviously,  $R_1 = 1$  if  $\mu = 0.$  Next, we transfer  $E_4$  to the origin by

$$\begin{aligned} x_1(t) &= B(t) - B_4^*, & x_2(t) &= X_U(t) - \frac{S_U}{\alpha_1 B_4^* + \mu_{X_U}}, \\ x_3(t) &= X_I(t) - \frac{\alpha_1 S_U B_4^* \mu_{I_A}}{(\alpha_2 (S_{I_A} + \beta_2 B_4^*) + \mu_{X_I} \mu_{I_A})(\alpha_1 B_4^* + \mu_{X_U})}, \\ x_4(t) &= I_R(t) - \frac{S_{I_R} + \beta_1 B_4^*}{\mu_{I_R}}, \\ x_5(t) &= I_A(t) - \frac{S_{I_A} + \beta_2 B_4^*}{\mu_{I_A}}. \end{aligned} \tag{21}$$

Normalizing the delay by  $t \rightarrow t/\tau,$  denoting  $x_i(\tau t)$  by  $u_i(t),$  and neglecting the higher order terms  $O(\mu^2),$  we have

$$\frac{du(t)}{dt} = A(\mu)u(t) + B(\mu)u(t-1) + f(u(t), \mu), \tag{22}$$

where

$$\begin{aligned} f(u, \mu) &= \tau \begin{pmatrix} f_1(u, \mu) \\ -\alpha_1 u_1 u_2 \\ \alpha_1 u_1 u_2 - \alpha_2 u_3 u_5 \\ 0 \\ 0 \end{pmatrix}, & B(\mu) &= \tau \begin{pmatrix} \frac{2\alpha_{20} b_0}{B_0} (b(\mu) + b_1 \mu) & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 \end{pmatrix}, \\ A(\mu) &= \tau \begin{pmatrix} \frac{\alpha_{20}}{\sigma} b(\mu) & 0 & 0 & -(d + \mu)b(\mu) & -\alpha_4 b(\mu) \\ -\alpha_1 d(\mu) & -\mu_{X_U} - \alpha_1 b(\mu) & 0 & 0 & 0 \\ \alpha_1 d(\mu) & \alpha_1 b(\mu) & -\mu_{X_I} - \alpha_2 c(\mu) & 0 & \alpha_2 k(\mu) \\ \beta_1 & 0 & 0 & -\mu_{I_R} & 0 \\ \beta_2 & 0 & 0 & 0 & -\mu_{I_A} \end{pmatrix}, \end{aligned}$$

$$\begin{aligned}
 f_1(u, \mu) &= u_1 \left( \frac{2\alpha_{20}b(\mu)}{B_0} u_1(t-1) - (d + \mu) u_4 \right. \\
 &\quad \left. - \alpha_4 u_5 + \frac{\alpha_{20}}{B_0} u_1^2(t-1) - \frac{\alpha_{20}}{\sigma} u_1 \right) + \frac{\alpha_{20}b(\mu)}{B_0} u_1^2(t-1), \\
 b_0 &= \frac{B_0}{2\alpha_{20}} \left( \frac{\alpha_{20}}{\sigma} + \frac{d\beta_1}{\mu_{I_R}} + \frac{\alpha_4\beta_2}{\mu_{I_A}} \right), & b_1 &= \frac{B_0\beta_1}{2\alpha_{20}\mu_{I_R}}, & b(\mu) &= b_0 + \mu b_1, \\
 c_0 &= \frac{S_{I_A} + \beta_2 b_0}{\mu_{I_A}}, & c_1 &= \frac{\beta_2 b_1}{\mu_{I_A}}, & c(\mu) &= c_0 + \mu c_1, & d_0 &= \frac{S_U}{\mu_{X_U} + \alpha_1 b_0}, \\
 d_1 &= -\frac{S_U \alpha_1 b_1}{(\mu_{X_U} + \alpha_1 b_0)^2}, & d(\mu) &= d_0 + \mu d_1, & k(\mu) &= k_0 + \mu k_1, \\
 k_0 &= -\frac{\alpha_1 d_0 b_0}{\mu_{X_I} + \alpha_2 c_0}, & k_1 &= \frac{d_1^2 (\alpha_2 c_1 b_0 (\mu_{X_U} + b_0 \alpha_1) - \mu_{X_U} (\mu_{X_I} + \alpha_2 c_0))}{b_1 (\mu_{X_I} + \alpha_2 c_0)^2}.
 \end{aligned}
 \tag{23}$$

Let  $\mathcal{C} = \mathcal{C}([-1, 0], \mathbb{R}^5)$  be the Banach space of continuous functions from  $[-1, 0]$  into  $\mathbb{R}^5$  with supremum norm. Define  $z_t \in \mathcal{C}$  as  $z_t(\theta) = z(t + \theta)$ ,  $\theta \in [-1, 0]$ . Equation (22) can be written as the functional differential equation

$$\dot{z}(t) = L(\mu)(z_t) + F(z_t, \mu), \tag{24}$$

where  $V$  is a neighborhood of zero in space of real numbers,  $L : \mathcal{C} \times V \rightarrow \mathbb{R}^5$  is a parameterized family of bounded linear operators, and  $F : \mathcal{C} \times V \rightarrow \mathbb{R}^5$  is a function with

$F(0, \mu) = 0$ ,  $\partial F(0, \mu)/\partial z = 0$  for all  $\mu \in \mathbb{R}$ . They have the following respective forms:

$$\begin{aligned}
 L(\mu)(\varphi) &= L(0)\varphi + L_1(\mu)\varphi, \\
 F(\varphi, \mu) &= F_2(\varphi, \mu) + F_3(\varphi, \mu),
 \end{aligned}
 \tag{25}$$

where

$$\begin{aligned}
 L(0)\varphi &= \tau \begin{pmatrix} -\frac{\alpha_{20}b_0}{\sigma} \varphi_1(0) - \frac{2\alpha_{20}b_0}{B_0} \varphi_1(-1) - b_0 d \varphi_4(0) - \alpha_4 b_0 \varphi_5(0) \\ -\alpha_1 d_0 \varphi_1(0) - (\mu_{X_U} + \alpha_1 b_0) \varphi_2(0) \\ \alpha_1 d_0 \varphi_1(0) + \alpha_1 b_0 \varphi_2(0) - (\alpha_2 c_0 + \mu_{X_I}) \varphi_3(0) - \alpha_1 k_0 \varphi_5(0) \\ \beta_1 \varphi_1(0) - \mu_{I_R} \varphi_4(0) \\ \beta_2 \varphi_1(0) - \mu_{I_A} \varphi_5(0) \end{pmatrix}, \\
 L_1(\mu)\varphi &= \tau \mu \begin{pmatrix} -\frac{\alpha_{20}b_1}{\sigma} \varphi_1(0) + \frac{4\alpha_{20}b_0 b_1}{B_0} \varphi_1(-1) - (b_0 + d b_1) \varphi_4(0) - \alpha_4 b_1 \varphi_5(0) \\ -\alpha_1 d_1 \varphi_1(0) - \alpha_1 b_1 \varphi_2(0) \\ \alpha_1 d_1 \varphi_1(0) + \alpha_1 b_1 \varphi_2(0) + \alpha_2 c_1 \varphi_3(0) - \alpha_2 k \varphi_5(0) \\ 0 \\ 0 \end{pmatrix},
 \end{aligned}$$

$$\begin{aligned}
 F_2(\varphi, \mu) &= \tau \begin{pmatrix} \varphi_1(0) \left( \frac{2\alpha_{20}b_0}{B_0} \mu \varphi_1(-1) - d\varphi_4(0) - \alpha_4 \varphi_5(0) - \frac{\alpha_{20}}{\sigma} \varphi_1(0) \right) + \frac{\alpha_{20}b_1}{B_0} \varphi_1^2(-1) \\ -\alpha_1 \varphi_1(0) \varphi_2(0) \\ \alpha_1 \varphi_1(0) \varphi_2(0) - \alpha_2 \varphi_3(0) \varphi_5(0) \\ 0 \\ 0 \end{pmatrix}, \\
 F_3(\varphi, \mu) &= \tau \begin{pmatrix} \varphi_1(0) \left( \frac{2b_1\alpha_{20}}{B_0} \mu \varphi_1(-1) - \mu \varphi_4(0) + \frac{\alpha_{20}}{B_0} \varphi_1^2(-1) \right) + \frac{\alpha_{20}b_1}{B_0} \mu \varphi_1^2(-1) \\ 0 \\ 0 \\ 0 \\ 0 \end{pmatrix}.
 \end{aligned}
 \tag{26}$$

From the Riesz representation theorem the linear map  $L$  can be expressed in integral form as follows:

$$L(\mu)(\varphi) = \int_{-1}^0 d\eta_\mu(\theta) \varphi(\theta), \tag{27}$$

where  $\eta_\mu$  is a bounded variation matrix-valued function on  $[-1, 0]$ . In fact, we can define  $\eta_\mu$  as

$$\eta_\mu(\theta) = A(\mu) \delta(\theta) + B(\mu) \delta(\theta + 1), \tag{28}$$

where  $\delta(\cdot)$  is the Dirac delta function.

Let  $\mathbb{R}^{5*}$  be the 5-dimensional vector space of row vectors, and denote  $\mathcal{E}^* = \mathcal{C}([-1, 0], \mathbb{R}^{5*})$ . Then, the adjoint bilinear form on  $\mathcal{E}^* \times \mathcal{E}$  can be defined as

$$\begin{aligned}
 \langle \psi(s), \varphi(\theta) \rangle &= \psi(0) \varphi(0) \\
 &- \int_{-1}^0 \int_0^\theta \psi(\xi - \theta) d\eta_\mu(\theta) \varphi(\xi) d\xi,
 \end{aligned}
 \tag{29}$$

where  $\varphi \in \mathcal{E}$  and  $\psi \in \mathcal{E}^*$ .

Let  $A(\mu)$  be the infinitesimal generator of the flow for the linear system

$$\dot{z}(t) = L(\mu) z_t \tag{30}$$

with spectrum  $\sigma[A(\mu)]$ . The adjoint operator  $A^*(\mu)$  is defined as the infinitesimal generator for the solution operator of the adjoint equation in  $\mathcal{E}^*$ :

$$\dot{w}(t) = - \int_{-1}^0 w(t - \theta) d\eta_\mu(\theta). \tag{31}$$

It is well known that the eigenvalues of  $A(\mu)$  with zero real parts play an important role in the bifurcation theory of RFDES. Denote  $A(0)$  by  $A_0$ , and let  $\Lambda_0 = \{\lambda \in \sigma(A_0) \mid \Re \lambda = 0\}$ . We have  $\Lambda_0 = \{0\}$ .

Using the formal adjoint theory for FDEs in [18], the phase space  $\mathcal{E}$  can be decomposed by  $\Lambda_0$  as  $\mathcal{E} = P \oplus Q$ ,

where  $P$  is the generalized eigenspace associated with the eigenvalues in  $\Lambda_0$ ,  $Q = \{\varphi \in \mathcal{E} \mid \langle \psi, \varphi \rangle = 0 \text{ for all } \psi \in P^*\}$ , and the dual space  $P^*$  is the generalized eigenspace for  $A^*(0)$  associated with the eigenvalues in  $\Lambda_0$ . Assume that  $\Phi$  and  $\Psi$  are the respective dual bases of  $P$  and  $P^*$  and satisfy  $\langle \Psi(s), \Phi(\theta) \rangle = 1$ . We might as well choose  $\Phi$  and  $\Psi$  as follows:

$$\begin{aligned}
 \Phi(\theta) &= \left( 1, -\frac{\alpha_1 d_0}{\mu_{X_U} + \alpha_1 b_0}, \frac{\alpha_1}{\mu_{X_I} + \alpha_2 c_0} \right. \\
 &\cdot \left. \left( d_0 - \frac{b_0 d_0}{\mu_{X_U} + \alpha_1 b_0} - \frac{k_0 \beta_2}{\mu_{I_A}} \right), \frac{\beta_1}{\mu_{I_R}}, \frac{\beta_2}{\mu_{I_A}} \right)^T, \\
 &-1 \leq \theta \leq 0,
 \end{aligned}
 \tag{32}$$

$$\Psi(s) = \left( 1, 0, 0, -\frac{b_0 d}{\mu_{I_R}}, -\frac{\alpha_4 b_0}{\mu_{I_A}} \right) l, \quad 0 \leq s \leq 1,$$

where  $\cdot^T$  is the transpose of  $\cdot$  and  $l = 1/((1 - b_0 d \beta_1 / \mu_{I_R}^2 - \alpha_4 b_0 \beta_2 / \mu_{I_A}^2) + \tau(2\alpha_{20}b_0/B_0))$ .

Let  $B = 0$ . Then, the following equations hold simultaneously:

$$\dot{\Phi} = \Phi B, \quad \dot{\Psi} = -B\Psi. \tag{33}$$

As shown in [16, 17], an appropriate phase space for considering normal forms of (24) is the Banach space  $\mathbf{BC}$  of functions from  $[-1, 0]$  into  $\mathbb{R}^5$ , which are uniformly continuous on  $[-1, 0)$  and with a jump discontinuity at 0. Then, the elements of  $\mathbf{BC}$  have the form  $\varphi + X_0 \rho$ , where  $\varphi \in \mathcal{E}$ ,  $\rho \in \mathbb{R}^5$ , and

$$X_0(\theta) = \begin{cases} I, & \theta = 0, \\ 0, & -1 \leq \theta < 0, \end{cases} \tag{34}$$

so that  $\mathbf{BC}$  is identified with  $\mathcal{E} \times \mathbb{R}^5$  with the norm  $|\varphi + X_0 \rho| = |\varphi|_{\mathcal{C}} + |\rho|_{\mathbb{R}^5}$ .

Let  $\pi : \mathbf{BC} \rightarrow P$  denote the projection

$$\pi(\varphi + X_0\rho) = \Phi(\langle \Psi, \varphi \rangle + \Psi(0)\rho), \quad \varphi \in \mathcal{E}, \rho \in \mathbb{R}^5, \tag{35}$$

and then the decomposition  $\mathcal{E} = P \oplus Q$  yields a decomposition of  $\mathbf{BC}$  by  $\Lambda_0$  as the topological direct sum  $\mathbf{BC} = P \oplus \text{Ker } \pi$  with the property  $Q \subset \text{Ker } \pi$ , where  $Q$  is an infinite dimensional complementary subspace of  $P$  and  $\mathcal{E}$  as shown above. Now, we decompose  $z_t \in \mathcal{E}^1$  in (24) as  $z_t = \Phi x(t) + y$ , where  $x(t) \in \mathbb{R}$  and  $y \in Q^1 = Q \cap \mathcal{E}^1$ ,  $\mathcal{E}^1$  is the subset of  $\mathcal{E}$  consisting of continuously differentiable functions.

Next, we rewrite (24) as follows:

$$\dot{z}(t) = L_0 z_t + L_1(\mu) z_t + F(z_t, \mu), \tag{36}$$

And, then, under the composition  $z_t = \Phi x(t) + y$ , (24) can be decomposed as a system of ODEs in  $\mathbb{R} \times \text{Ker } \pi$  as follows:

$$\begin{aligned} \dot{x} &= Bx + f_2^1(x, y, \mu) + f_3^1(x, y, \mu), \\ \dot{y} &= A_{Q^1} y + f_2^2(x, y, \mu) + f_3^2(x, y, \mu), \end{aligned} \tag{37}$$

where  $A_{Q^1} = \dot{y} + X_0(L_0(y) - \dot{y}(0))$  is the restriction of  $A$  as an operator from  $Q^1$  into  $\text{Ker } \pi$ , and

$$\begin{aligned} f_2^1(x, y, \mu) &= \Psi(0) [L_1(\mu) (\Phi x + y) + F_2(\Phi x + y, \mu)], \\ f_2^2(x, y, \mu) &= (I - \pi) X_0 [L_1(\mu) (\Phi x + y) + F_2(\Phi x + y, \mu)], \\ f_3^1(x, y, \mu) &= \Psi(0) F_3(\Phi x + y, \mu), \\ f_3^2(x, y, \mu) &= (I - \pi) X_0 F_3(\Phi x + y, \mu). \end{aligned} \tag{38}$$

As for autonomous ODEs in  $\mathbb{R}^5$ , the normal forms are obtained by a recursive process of changes of variables. At a step  $j$ , the terms of order  $j = 2$  are computed from the terms of the same order and from the terms of lower orders already computed in previous steps. Assuming that steps of orders  $2, 3, \dots, j - 1$ , have already been performed leads to

$$\begin{aligned} \dot{x} &= Bx + \sum_{l \geq 2}^{j-1} g_l^1(x, y, \mu) + \tilde{f}_j^1(x, y, \mu) + \text{h.o.t.}, \\ \dot{y} &= A_{Q^1} y + \sum_{l \geq 2}^{j-1} g_l^2(x, y, \mu) + \tilde{f}_j^2(x, y, \mu) + \text{h.o.t.}, \end{aligned} \tag{39}$$

where  $\tilde{f}_j \in (\tilde{f}_j^1, \tilde{f}_j^2)$  is the terms of order  $j$  in  $(x, y, \mu)$  after the previous transformations of variables and h.o.t stands for the higher order terms. Following the algorithm of [16, 17] at step  $j$ , using a change of variables of the form

$$(x, y) = (\hat{x}, \hat{y}) + U_j(\hat{x}, \mu) \equiv (\hat{x}, \hat{y}) + [U_j^1(\hat{x}, \mu), U_j^2(\hat{x}, \mu)], \tag{40}$$

where  $x, \hat{x} \in \mathbb{R}$ ,  $y, \hat{y} \in Q^1$ , and  $U_j^1 : \mathbb{R}^2 \rightarrow \mathbb{R}, U_j^2 : \mathbb{R}^2 \rightarrow Q^1$  are homogeneous polynomials of degree  $j$  in  $\hat{x}$  and  $\epsilon$ , after

dropping the hats for simplification of notations, (37) can be put into the normal form

$$\begin{aligned} \dot{x} &= Bx + \sum_{j \geq 2} g_j^1(x, y, \mu), \\ \dot{y} &= A_{Q^1} y + \sum_{j \geq 2} g_j^2(x, y, \mu), \end{aligned} \tag{41}$$

where

$$\begin{aligned} g_j^1(x, y, \mu) &= \tilde{f}_j^1(x, y, \mu) - [D_x U_j^1(x, \mu) Bx - B U_j^1(x)], \\ g_j^2(x, y, \mu) &= \tilde{f}_j^2(x, y, \mu) - [D_x U_j^2(x, \mu) Bx - A_{Q^1} U_j^2(x)]. \end{aligned} \tag{42}$$

It can be verified that (24) satisfies nonresonance conditions; see [16, 17]. Then, the locally invariant manifold for (24) tangent to  $P$  at zero must be  $y = 0$  and the flow on this manifold is given by 1-dimensional ODE

$$\dot{x} = Bx + g_2^1(x, 0, \mu) + g_3^1(x, 0, \mu) + \text{h.o.t.} \tag{43}$$

The nonlinear terms in (41) are in normal form in the classical sense with respect to matrix  $B$ . In application,  $g_j^1(x, 0, \mu)$  usually can be determined by the following procedure.

**Theorem 8.** For  $j \geq 2$ , let  $M_j$  denote the operator defined in  $V_j(\mathbb{R}^2 \times \text{Ker } \pi)$ , with values in the same place, by

$$\begin{aligned} M_j(h_1, h_2) &= (M_j^1 h_1, M_j^2 h_2), \\ (M_j^1 h_1)(x, \mu) &= D_x h_1(x, \mu) Bx - B h_1(x, \mu), \end{aligned} \tag{44}$$

$$(M_j^2)(x, \mu) = D_x h_2(x, \mu) Bx - A_{Q^1} [h_2(x, \mu)]$$

with domain  $D(M_j) = V_j^2(\mathbb{R}^2) \times V_j^2(Q^1)$ . Here, we use the notation  $V_j^2(Y)$  to denote the space of homogeneous polynomials of degree  $j$  in 2 variables  $(x, \epsilon) \in \mathbb{R}^2$ , with coefficients in a Banach space  $Y$ .

According to [16, 17], we derive

$$U_j(x) = M_j^{-1} P_{1,j} \tilde{f}_j(x, 0, \mu) \in \text{Ker}(M_j)^c, \tag{45}$$

and then

$$g_j^1(x, 0, \mu) = (I - P_{1,j}) \tilde{f}_j^1(x, 0, \mu) \in \text{Im}(M_j^1)^c, \tag{46}$$

where  $P_{1,j} = (P_{1,j}^1, P_{1,j}^2)$  is the projection of  $V_j^2(\mathbb{R}^2) \times V_j^2(\text{Ker } \pi)$  on  $\text{Im}(M_j^1) \times \text{Im}(M_j^2)$ .

Since  $B = 0$ , it can be checked that  $[\text{Im}(M_j^1)]^c = \text{span}\{x^2, x\mu, \mu^2\}$ . Then, we obtain

$$\begin{aligned} g_2^1(x, 0, \mu) &= l\mu\tau \left( \frac{4\alpha_{20} b_0 b_1}{B_0} - \frac{d\beta_1}{\mu_{1R}} - \frac{(b_0 + db_1)\beta_1}{\mu_{1R}} - \frac{\alpha_4 b_1 \beta_2}{\mu_{1A}} \right) x \\ &+ l\tau \left( \frac{3b_0 \alpha_{20}}{B_0} - \frac{d\beta_1}{\mu_{1R}} - \frac{\alpha_4 \beta_2}{\mu_{1A}} - \frac{\alpha_{20}}{\sigma} + \frac{\alpha_{20} b_0}{B_0} \right) x^2, \end{aligned} \tag{47}$$



and further then the normal form of (24) on the invariant local center manifold  $y = 0$  is given by

$$\begin{aligned} \dot{x} = & l\mu\tau \left( \frac{4\alpha_{20}b_0b_1}{B_0} - \frac{d\beta_1}{\mu_{I_R}} - \frac{(b_0 + db_1)\beta_1}{\mu_{I_R}} - \frac{\alpha_4b_1\beta_2}{\mu_{I_A}} \right) x \\ & + l\tau \left( \frac{3b_0\alpha_{20}}{B_0} - \frac{d\beta_1}{\mu_{I_R}} - \frac{\alpha_4\beta_2}{\mu_{I_A}} - \frac{\alpha_{20}}{\sigma} \right) x^2 \\ & + f_3(x, 0, \mu) + \text{h.o.t.} \end{aligned} \quad (48)$$

If  $R_0 < 1$ , by using the definitions, we can verify that

$$\frac{b_0\alpha_{20}}{B_0} > \frac{d\beta_1}{\mu_{I_R}} + \frac{\alpha_4\beta_2}{\mu_{I_A}} + \frac{\alpha_{20}}{\sigma}. \quad (49)$$

**Theorem 9.** *If  $R_1 = 1$ , the positive equilibrium  $E_4$  is unstable for any  $\tau > 0$ .*

For  $R_0 = 1$ ,  $E_0$  is also an equilibrium with simple zero singularity. To discuss its stability, we employ the following perturbation form:

$$\alpha_{20} = \bar{d} + \bar{\mu}, \quad (50)$$

where  $\bar{d} = (\alpha_3 S_{I_R} / \mu_{I_R}) + (\alpha_4 S_{I_A} / \mu_{I_A})$  and  $\bar{\mu}$  is a small parameter. Then, the normal form of (24) near the bacteria-free equilibrium is as follows:

$$\dot{x} = \tau \bar{\mu} x - \tau \left( \frac{\bar{d}}{\sigma} + \frac{\alpha_3 \beta_1}{\mu_{I_R}} + \frac{\alpha_4 \beta_2}{\mu_{I_A}} \right) x^2 + \text{h.o.t.} \quad (51)$$

**Theorem 10.** *If  $R_0 = 1$ , the bacteria-free equilibrium is unstable for any  $\tau > 0$ .*

### 5. Sensitive Analysis

Sensitivity indices allow us to measure the relative change in a variable when a parameter changes. The normalized forward sensitivity index of a variable to a parameter is the ratio of the relative change in the variable to the relative change in the parameter. When the variable is a differentiable function of the parameter, the sensitivity index may be alternatively defined using partial derivatives. Here, one adopts the following definition as described by Chitnis et al. [19].

*Definition 11.* The normalized forward sensitivity index of a variable,  $u$ , that depends differentially on a parameter,  $p$ , is defined as  $\gamma_p^u = (\partial u / \partial p) \times (p/u)$ .

To clear bacteria in the body, we must take measures to make  $R_0 > 1$  hold. For this end, it is important to determine how crucial each parameter is to  $R_0$ . Table 1 exhibits the analytical sensitivity indices of  $R_0$  to the parameters  $\alpha_{20}$ ,  $\alpha_3$ ,  $\alpha_4$ ,  $\mu_{I_R}$ ,  $\mu_{I_A}$ ,  $S_{I_R}$ , and  $S_{I_A}$ .

The negative sign of the sensitivity index for  $R_0$  implies that increase in the relevant parameter leads to the decrease

TABLE 1: Sensitivity indices of  $R_0$  to its parameters.

Parameter $j$	$\alpha_{20}$	$\alpha_3$	$\alpha_4$	$\mu_{I_R}$	$\mu_{I_A}$	$S_{I_R}$	$S_{I_A}$
$\gamma_j^{R_0}$	-1	$l_1$	$l_2$	$-l_1$	$-l_2$	$l_1$	$l_2$

$l_1 = S_{I_R} \mu_{I_A} \alpha_3 / (\alpha_3 S_{I_R} \mu_{I_A} + \alpha_4 S_{I_A} \mu_{I_R})$  and  $l_2 = S_{I_A} \mu_{I_R} \alpha_4 / (\alpha_3 S_{I_R} \mu_{I_A} + \alpha_4 S_{I_A} \mu_{I_R})$ .

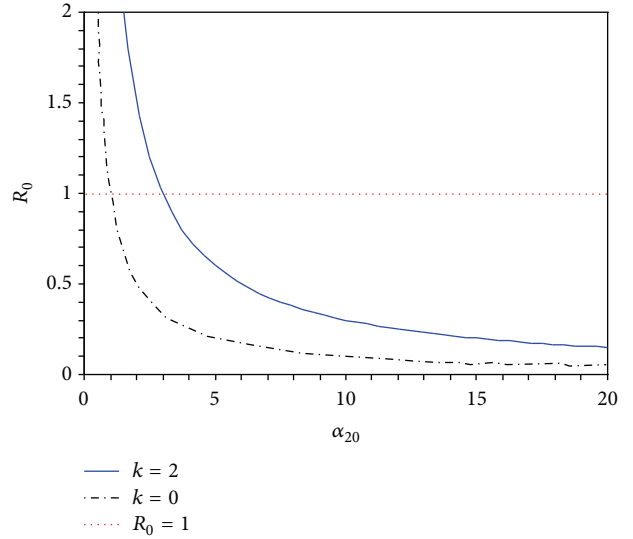


FIGURE 1: Plot of  $R_0$  against the the effective reproductive rate  $\alpha_{20} \in (0, 20]$  for  $k = 0, 2$ .

in  $R_0$ . Note that  $0 < l_1, l_2 < 1$ . The most sensitive parameter is the growth rate  $\alpha_{20}$ , which has a high impact on  $R_0$  and should be targeted by intervention strategies. To reduce  $\alpha_{20}$  to ensure  $R_0 > 1$ , we may develop an inhibitor to control the reproduction of bacteria or to kill bacteria individual, which agrees with an intuitive expectation.

To understand how  $\alpha_{20}$  affects  $R_0$ , we might as well assume  $\alpha_4 = k\alpha_3$ ,  $S_{I_A} = kS_{I_R}$ , and  $\mu_{I_A} = k\mu_{I_R}$ , and then gets  $R_0 = (\alpha_3 S_{I_R} / \alpha_{20} \mu_{I_R})(1 + k)$ . Clearly, for  $k = 0$ , only the innate immune cell competes with bacteria. Figure 1 exhibits the relationship between  $R_0$  and  $\alpha_{20}$  with  $\alpha_3 = \mu_{I_R} = 0.02$ ,  $S_{I_R} = 0.04$ , and  $k = 0, 2$ . It can be seen from the plot that as  $\alpha_{20}$  decreases  $R_0$  increases faster when  $k = 2$  than the case of  $k = 0$ , which reveals that vaccination or other strategies adopted to stimulate immunity of the body are beneficial to the clearance of bacteria.

Figure 2 shows the solutions of system (2) starting from different initial values chosen arbitrarily together with  $\alpha_{20} = 0.01, 0.5, 2$ ,  $\alpha_3 = 0.02$ ,  $\alpha_4 = 0.04$ ,  $S_{I_R} = 1$ ,  $S_{I_A} = 2$ ,  $\mu_{I_R} = 0.02$ ,  $\mu_{I_A} = 0.04$ ,  $B_0 = 1$ ,  $\sigma = 1/3$ ,  $\beta_1 = 0.01$ ,  $\beta_2 = 0.02$ ,  $\alpha_1 = 0.01$ ,  $\alpha_2 = 0.01$ ,  $\mu_{X_U} = 0.02$ , and  $\mu_{X_I} = 0.04$ . It is easy to see that the bacteria concentration  $B(t)$  tends to zero, and the smaller growth rate leads to the faster convergence.

### 6. Conclusions

This paper formulates the competition between bacteria and immune system by DDEs. Then, the qualitative properties of the model are analyzed. Specially, by virtue of the center manifold reduction and normal form theory due to Faria

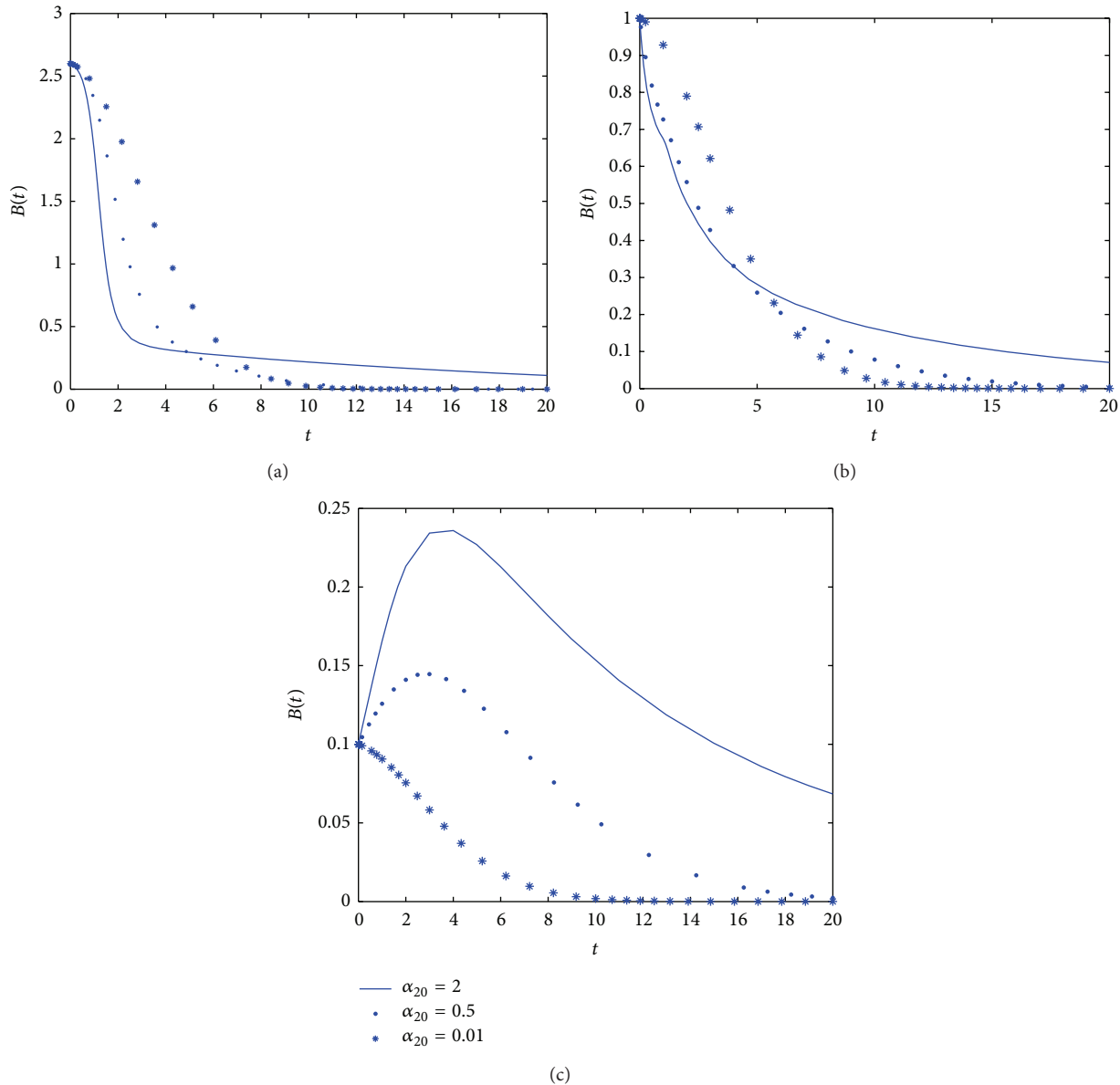


FIGURE 2: Simulations of  $B(t)$  for  $\alpha_{20} = 2, 0.5, 0.01$  in each figure.

and Magalhaes [16, 17], the normal form of system (2) associated with zero eigenvalue is computed, from which one deduces that the bacteria-free equilibrium  $E_0$  and the positive equilibrium  $E_4$  are unstable under the conditions of  $R_0 = 1$  and  $R_1 = 1$ , respectively. Next, sensitivity analysis and numerical simulations indicate that the effective reproductive rate  $\alpha_{20}$  is the most sensitive parameter to  $R_0$ . Theorem 3 suggests the strategies target the decrease of the growth rate which can be successful in disease elimination.

On the biological viewpoint, the terms  $\alpha_3 S_{I_R} / \mu_{I_R}$  and  $\alpha_4 S_{I_A} / \mu_{I_A}$  measure the respective strengths of the innate and adaptive immune system defense against the bacterial challenge, while the factor  $\alpha_{20}$  measures the bacteria's offensive strength. So with  $(\alpha_3 S_{I_R}) / (\alpha_{20} \mu_{I_R}) + (\alpha_4 S_{I_A}) / (\alpha_{20} \mu_{I_A})$ , we can compare the strength of the immune system against the bacterial offensive. Thus, Theorems 2 and 3 have the biological

explanations: in the domain of attraction of  $E_0$ , bacteria will be cleared if  $R_0 > 1$ ; that is, the strength of the immune system defense against the bacteria challenge is not weaker than the bacteria's offensive strength; in the domain of attraction of  $E_3$ , bacteria coexist with immune cells when  $R_0 < 1$  and the bacterial challenge is weaker than bacteria's offensive strength.

### Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

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