Research Article

The Effect of Impulsive Diffusion on Dynamics of a Stage-Structured Predator-Prey System

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We investigate a predator-prey model with impulsive diffusion on predator and stage structure on prey. The globally attractive condition of prey-extinction periodic solution of the system is obtained by the stroboscopic map of the discrete dynamical system. The permanent condition of the system is also obtained by the theory of impulsive delay differential equation. The results indicate that the discrete time delay has influence on the dynamical behaviors of the system. Finally, some numerical simulations are carried out to support the analytic results.

1. Introduction

The dispersal is a ubiquitous phenomenon in the natural world. It is well recognized that the spatial distribution of populations and population dynamics are much affected by spatial heterogeneity and population mobility [1]. The fragmented landscapes are common because the populations of most species occupy mosaic habitats and because of rapid destruction of natural habitats. Briggs and Hoopes [2] identify three mechanisms whereby limited dispersal of hosts and parasitoids combined with other features, such as spatial and temporal heterogeneity, can promote persistence and stability of populations mirrored by the large number of mathematical models devoting to it in the scientific literatures [3–6]. In recent years, the analysis of these models focuses on the coexistence of population and local (or global) stability of the population, although the complete results have not yet been obtained even

in the simplest one-species case. Most previous papers focused on the population dynamical system modeled by the ordinary differential equations; if the population dynamics with the effects of spatial heterogeneity are modeled by a diffusion process, it will be very interesting. While in practice, it is often the case that diffusion occurs in regular pulse. For example, when winter comes, birds will migrate between patches in search for a better environment, whereas they do not diffuse in other seasons, and the excursion of foliage seeds occurs at fixed period of time every year. Thus, impulsive diffusion provides a more natural description. Lately theories of impulsive differential equations [14] have been introduced into population dynamics. Impulsive differential equations are found in almost a domain of applied science [15–24]. Newly, persistence and stability of population dynamical system involving time delay have been discussed by some authors; see, for example, [8, 25, 26] and references cited therein. They obtained some sufficient conditions that guarantee permanence of population or stability of positive equilibria or positive periodic solutions.

The organization of this paper is as follows. In the next section, we introduce the model and background concepts. In Section 3, some important lemmas are presented. In Section 4, we give the conditions of global attractivity and permanence for system (2.3). In Section 5, A brief discussion is given in the last section to conclude this work.

2. The Model

Stage-structured models were analyzed in many literatures [7, 27–33]. The following stagestructured Holling mass defence predator-prey model with impulsive perturbations on predators was introduced by Jiao et al. [31]:

$$\begin{aligned} x_1'(t) &= rx_2(t) - re^{-\omega\tau_1}x_2(t - \tau_1) - \omega x_1(t), \\ x_2'(t) &= re^{-\omega\tau_1}x_2(t - \tau_1) - \frac{\beta x_2(t)}{1 + ax_2 + bx_2^2}x_3(t) - d_3x_2(t) - d_4x_2^2(t), \quad t \neq n\tau, \\ x_3'(t) &= \frac{k\beta x_2(t)}{1 + ax_2 + bx_2^2}x_3(t) - dx_3(t), \\ \Delta x_1(t) &= 0, \\ \Delta x_2(t) &= 0, \quad t = n\tau, \ n = 1, 2, \dots, \\ \Delta x_3(t) &= \mu, \end{aligned}$$

$$(\varphi_1(\zeta), \varphi_2(\zeta), \varphi_3(\zeta)) \in C_+ = C\Big([-\tau_1, 0], R_+^3\Big), \quad \varphi_i(0) > 0, \ i = 1, 2, 3, \end{aligned}$$

where $x_1(t)$, $x_2(t)$ represent the immature and mature pest densities, respectively, $x_3(t)$ denotes the density of nature enemy, τ_1 represents a constant time to maturity, and $r, w, d_3, d_4, d, k, a, b$ and β are positive constants. This model is derived as follows. We assume that at any time t > 0, birth into the immature population is proportional to the existing mature population with proportionality constant r. We then assume that the death rate of immature population is proportional to the existing immature population with proportional to the existing immature population with proportionality constant w. w (w > d), d_3 , and d ($d > d_3$) are called the death coefficient of $x_1(t), x_2(t)$, and $x_3(t)$, respectively. We assume that the death rate of mature populations are

of a logistic nature, that is, proportional to the square of the population with proportionality constant d_4 . k > 0 is the rate of conversing prey into predator. $\Delta x_3(t) = x_3(t^+) - x_3(t), \mu \ge 0$ is the releasing amount of natural enemies at $t = n\tau$, $n \in Z_+$ and $Z_+ = \{1, 2, ...\}, \tau$ and is the period of the impulsive immigration of the predator.

Diffusive predator-prey models were analyzed in literatures [16, 34, 35]. In [16], Hui and Chen considered the single species model with impulsive diffusion as follows:

$$\frac{dx_{1}(t)}{dt} = x_{1}(t)(a_{1} - b_{1}x_{1}(t)),$$

$$\frac{dx_{2}(t)}{dt} = x_{2}(t)(a_{2} - b_{2}x_{2}(t)),$$

$$\Delta x_{1}(t) = d_{1}(x_{2}(t) - x_{1}(t)),$$

$$\Delta x_{2}(t) = d_{2}(x_{1}(t) - x_{2}(t)),$$

$$t = n\tau, \ n = 1, 2, \dots,$$
(2.2)

where we suppose that the system is composed of two patches connected by diffusion and x_i (i = 1, 2) is the density of species in the *i*th patch. Intrinsic rate of natural increase and density dependence rate of prey population in the first habitat are denoted by a_i (i = 1, 2), and d_i (i = 1, 2) is the dispersal rate in the *i*th patch. It is assumed here that the net exchange from the *j*th patch to *i*th patch is proportional to the difference $x_j - x_i$ of population densities. The pulse diffusion occurs every τ period ($\tau > 0$). The system evolves from its initial state without being further affected by diffusion until the next pulse appears; $\Delta x_i = x_i(n\tau^+) - x_i(n\tau)$, and $x_i(n\tau^+)$ represents the density of population in the *i*th patch immediately after the *n*th diffusion pulse at time $t = n\tau$, while $x_i(n\tau)$ represents the density of population in the *i*th patch before the *n*th diffusion pulse at time $t = n\tau$, $n = 0, 1, 2, ...; r_i, k_i$, and d_i (i = 1, 2) are positive constants.

Motivated by these ideals from all of the above, we assume that the predator population diffuses between the two patches, and the prey population exists only in one patch. Then, we consider a delayed predator-prey model with impulsive diffusion on predator and stage structure on prey as follows:

$$\frac{dx_{1}(t)}{dt} = rx_{2}(t) - re^{-w\tau_{1}}x_{2}(t-\tau_{1}) - wx_{1}(t),$$

$$\frac{dx_{2}(t)}{dt} = re^{-w\tau_{1}}x_{2}(t-\tau_{1}) - dx_{2}(t) - \beta x_{2}(t)y_{1}(t),$$

$$\frac{dy_{1}(t)}{dt} = -a_{1}y_{1}(t) + k\beta x_{2}(t)y_{1}(t),$$

$$\frac{dy_{2}(t)}{dt} = y_{2}(t)(a_{2} - b_{2}y_{2}(t)),$$

$$\Delta x_{1}(t) = 0,$$

$$\Delta x_{1}(t) = 0,$$

$$\Delta y_{1}(t) = D(y_{2}(t) - y_{1}(t)),$$

$$dx_{1}(t) = D(y_{1}(t) - y_{2}(t)),$$
(2.3)

with initial condition

$$\left(\varphi_1(\zeta),\varphi_2(\zeta),\varphi_3(\zeta),\varphi_4(\zeta)\right) \in C_+ = C\left([-\tau_1,0],R_+^4\right), \quad \varphi_i(0) > 0, \ i = 1,2,3,4,$$
(2.4)

where $x_1(t)$, $x_2(t)$ represent the immature and mature prey population densities, respectively. τ_1 represents a constant time to maturity, that is, immature individuals and mature individuals are divided by age τ_1 . It is assumed that the system is composed of two patches connected by diffusion and occupied by a single species and $y_i(t)$ (i = 1, 2) is the density of predator species in the *i*th patch. Death rate of the predator population in the first patch is denoted by $a_1 > 0$. Intrinsic rate of natural increase and density dependence rate of predator population in the second patch are denoted by $a_2 > 0$ and $b_2 > 0$; a_2/b_2 denotes the carrying capacity in the second patch. 0 < D < 1 is dispersal rate of the predator population between two patches. It is assumed here that the net exchange from the *j*th patch to *i*th patch is proportional to the difference $y_i - y_i$ of predator population densities. β is the prey capture rate by mature predator. k is the rate of conversion of nutrients into the reproduction rate of the mature predator. w is the death rate of immature prey population in the first patch. d is the death rate of mature prey population in first patch. The pulse diffusion occurs every $\tau > 0$ period. The system evolves from its initial state without being further affected by diffusion until the next pulse appears. $\Delta y_i(n\tau) = y_i(n\tau^+) - y_i(n\tau)$, where $y_i(n\tau^+)$ represents the density of population in the *i*th patch immediately after the *n*th diffusion pulse at time $t = n\tau$, while $y_i(n\tau)$ represents the density of population in the *i*th patch before the *n*th diffusion pulse at time $t = n\tau$ (n = 0, 1, 2, ...).

Obviously, (2.3) can be simplified as follows:

$$\frac{dx_{2}(t)}{dt} = re^{-t\omega\tau_{1}}x_{2}(t-\tau_{1}) - dx_{2}(t) - \beta x_{2}(t)y_{1}(t),
\frac{dy_{1}(t)}{dt} = -a_{1}y_{1}(t) + k\beta x_{2}(t)y_{1}(t), \qquad t \neq n\tau,
\frac{dy_{2}(t)}{dt} = y_{2}(t)(a_{2} - b_{2}y_{2}(t)), \qquad (2.5)
\Delta x_{2}(t) = 0,$$

$$\Delta y_1(t) = D(y_2(t) - y_1(t)), \quad t = n\tau, \ n = 1, 2, \dots,$$

$$\Delta y_2(t) = D(y_1(t) - y_2(t)),$$

with initial condition

$$\left(\varphi_2(\zeta),\varphi_3(\zeta),\varphi_4(\zeta)\right) \in C'_+ = C\left([-\tau_1,0],R^3_+\right), \quad \varphi_i(0) > 0, \ i = 2,3,4.$$
(2.6)

3. The Lemmas

The solution of (2.3), denoted by $x(t) = (x_1(t), x_2(t), y_1(t), y_2(t))^T$, is a piecewise continuous function $x : R_+ \to R_+^4$. x(t) is continuous on $(n\tau, (n+1)\tau]$, $n \in Z_+$, and $x(n\tau^+) = \lim_{t \to n\tau^+} x(t)$ exists. Obviously, the global existence and uniqueness of solutions of (2.3) is guaranteed by

the smoothness of properties of f, which denotes the mapping defined by right-side of system (2.3) (see Lakshmikantham et al. [36]). Before having the main results, we need some lemmas which will be used next.

According to the biological meanings, it is assumed that $x_1(t) \ge 0$, $x_2(t) \ge 0$, $y_1(t) \ge 0$, and $y_2(t) \ge 0$. Now, we will show that all solutions of (2.3) are uniformly ultimately bounded.

Lemma 3.1. There exists a constant M > 0 such that $x_1(t) \le M/k$, $x_2(t) \le M/k$, $y_1(t) \le M$, and $y_2(t) \le M$, for each solution $(x_1(t), x_2(t), y_1(t), y_2(t))$ of (2.3) with all large enough t.

Proof. Define $V(t) = kx_1(t) + kx_2(t) + y_1(t) + y_2(t)$. Denote $\overline{d} = \min\{a_1, r - d, w\}$. When $t \neq n\tau$, we have

$$D^{+}V(t) + \overline{d}V(t) = -k\left(w - \overline{d}\right)x_{1}(t) - k\left[(r - d) - \overline{d}\right]x_{2}(t)$$
$$-\left(a_{1} - \overline{d}\right)y_{1}(t) + y_{2}(t)\left[\left(a_{2} + \overline{d}\right) - b_{2}y_{2}(t)\right]$$
$$\leq y_{2}(t)\left(\left(a_{2} + \overline{d}\right) - b_{2}y_{2}(t)\right) \leq M_{0},$$
(3.1)

where $M_0 = (a_2 + \overline{d})^2 / 4b_2$. When $t = n\tau$, we also have $V(n\tau^+) = V(n\tau)$. By Lemma 2.3 in [14], for $t \in (n\tau, (n+1)\tau]$, we obtain

$$V(t) \leq V(0) \exp\left(-\overline{d}t\right) + \int_{0}^{t} M_{0} \exp\left(-\overline{d}(t-s)\right) ds$$

$$= V(0) \exp\left(-\overline{d}t\right) + \frac{M_{0}}{\overline{d}} \left(1 - \exp\left(-\overline{d}t\right)\right)$$

$$< V(0) \exp\left(-\overline{d}t\right) + \frac{M_{0}}{\overline{d}} \left(1 - \exp\left(-\overline{d}t\right)\right)$$

$$\longrightarrow \frac{M_{0}}{\overline{d}}, \quad \text{as } t \longrightarrow \infty.$$

(3.2)

So, V(t) is uniformly ultimately bounded. Hence, by the definition of V(t), there exists a constant M > 0 such that $x_1(t) \le M/k$, $x_2(t) \le M/k$, $y_1(t) \le M$, $y_2(t) \le M$ for all t large enough. The proof is complete.

If $x_2(t) = 0$, the subsystem of (2.3) is obtained as follows:

$$\frac{dy_{1}(t)}{dt} = -a_{1}y_{1}(t),$$

$$\frac{dy_{2}(t)}{dt} = y_{2}(t)(a_{2} - b_{2}y_{2}(t)),$$

$$\Delta y_{1}(t) = D(y_{2}(t) - y_{1}(t)),$$

$$\Delta y_{2}(t) = D(y_{1}(t) - y_{2}(t)),$$

$$t = n\tau, \ n = 1, 2, \dots$$
(3.3)

It is easy to solve the first two equations of system (3.3) between pulses

$$y_{1}(t) = y_{1}(n\tau^{+})e^{-a_{1}(t-n\tau)}, \quad n\tau < t \le (n+1)\tau,$$

$$y_{2}(t) = \frac{a_{2}e^{a_{2}(t-n\tau)}y_{1}(n\tau^{+})}{a_{2} + b_{2}(e^{a_{2}(t-n\tau)} - 1)y_{1}(n\tau^{+})}, \quad n\tau < t \le (n+1)\tau.$$
(3.4)

By considering the last two equations of System (3.3), we obtain the following stroboscopic map of system (3.3):

$$y_{1}((n+1)\tau^{+}) = (1-D)e^{-a_{1}\tau}y_{1}(n\tau^{+}) + D\frac{a_{2}e^{a_{2}\tau}y_{2}(n\tau^{+})}{a_{2} + b_{2}(e^{a_{2}\tau} - 1)y_{2}(n\tau^{+})},$$

$$y_{2}((n+1)\tau^{+}) = (1-D)\frac{a_{2}e^{a_{2}\tau}y_{2}(n\tau^{+})}{a_{2} + b_{2}(e^{a_{2}\tau} - 1)y_{2}(n\tau^{+})} + De^{-a_{1}\tau}y_{1}(n\tau^{+}).$$
(3.5)

Then, two lemmas are obtained as follows.

Lemma 3.2 (see [37, Theorem 3.1]). If $a_2\tau < \ln((1-(1-D)e^{-a_1\tau})/(1-(1-2D)e^{-a_1\tau}))$, the trivial equilibrium (0,0) of (3.5) is globally asymptotically stable. If $a_2\tau > \ln((1-(1-D)e^{-a_1\tau}))/(1-(1-2D)e^{-a_1\tau}))$, there exists a unique positive equilibrium (y_1^*, y_2^*) , which is globally asymptotically stable, where $y_1^* = De^{a_2\tau}y_2^*/((1-D)e^{a_2\tau} - (1-2D)e^{(a_2-a_1)\tau})$ and $y_2^* = a_2 [(1-D)(e^{a_2\tau} + e^{-a_1\tau}) - (1-2D)e^{(a_2-a_1)\tau} - 1]/b_2(e^{a_2\tau} - 1)(1-(1-D)e^{(a_2-a_1)\tau})$.

Lemma 3.3 (see [37, Theorem 3.2]). If $a_2\tau > \ln((1 - (1 - D)e^{-a_1\tau})/(1 - (1 - 2D)e^{-a_1\tau}))$, then (3.3) has a τ -periodic positive solution $(\widetilde{y_1(t)}, \widetilde{y_2(t)})$, which is globally asymptotically stable. Here $(\widetilde{y_1(t)}, \widetilde{y_2(t)})$ can be expressed as

$$\widetilde{y_1(t)} = y_1^* e^{-a_1(t-n\tau)}, \quad n\tau < t \le (n+1)\tau,$$

$$\widetilde{y_2(t)} = \frac{a_2 e^{a_2(t-n\tau)} y_2^*}{a_1 + b_2 (e^{a_2(t-n\tau)} - 1) y_2^*}, \quad n\tau < t \le (n+1)\tau,$$
(3.6)

where y_1^* and y_2^* are defined as in Lemma 3.2.

Lemma 3.4 (see [38]). Consider the following delay equation:

$$\frac{dx(t)}{dt} = c_1 x(t-\tau) - c_2 x(t).$$
(3.7)

one assumes that $c_1, c_2, \tau > 0$, x(t) > 0 for $-\tau \le t \le 0$. Assuming that $c_1 < c_2$, then

$$\lim_{t \to \infty} x(t) = 0. \tag{3.8}$$

4. The Dynamics

From the above discussion, we can easily know that there exists a prey-extinction boundary periodic solution $(0, 0, \widetilde{y_1(t)}, \widetilde{y_2(t)})$ of system (2.3). In this section, we will prove that the prey-extinction boundary periodic solution $(0, 0, \widetilde{y_1(t)}, \widetilde{y_2(t)})$ of system (2.3) is globally attractive.

Theorem 4.1. If

$$a_{2}\tau > \ln \frac{1 - (1 - D)e^{-a_{1}\tau}}{1 - (1 - 2D)e^{-a_{1}\tau}},$$

$$re^{-w\tau_{1}} < d + \frac{D\beta e^{a_{2}\tau}y_{2}^{*}}{(1 - D)e^{a_{2}\tau} - (1 - 2D)e^{(a_{2} - a_{1})\tau}},$$
(4.1)

hold, the prey-extinction boundary periodic solution $(0, 0, \widetilde{y_1(t)}, \widetilde{y_2(t)})$ of (2.3) is globally attractive. Here, $y_2^* = a_2 [(1-D)(e^{a_2\tau} + e^{-a_1\tau}) - (1-2D)e^{(a_2-a_1)\tau} - 1]/b_2(e^{a_2\tau} - 1)[1-(1-D)e^{(a_2-a_1)\tau}].$

Proof. It is clear that the global attractivity of the predator-extinction boundary periodic solution $(0, 0, \widetilde{y_1(t)}, \widetilde{y_2(t)})$ of system (2.3) is equivalent to the global attractivity of predator-extinction boundary periodic solution $(0, \widetilde{y_1(t)}, \widetilde{y_2(t)})$ of system (2.5). So, we devote ourselves to investigate system (2.5). Since $Dk\beta y_2^* e^{a_2\tau - w\tau_1} < d[(1 - D)e^{a_2\tau} - (1 - 2D)e^{(a_2 - a_1)\tau}]$, we can choose ε_0 sufficiently small such that

$$k\beta e^{-\omega\tau_1} \left[\frac{De^{a_2\tau}y_2^*}{(1-D)e^{a_2\tau} - (1-2D)e^{(a_2-a_1)\tau}} + \varepsilon_0 \right] < d.$$
(4.2)

Since we know from the second equation of system (2.5) that $dy_1(t)/dt \le -a_1y_1(t)$, we consider the following comparison an impulsive differential system:

$$\frac{dy'_{1}(t)}{dt} = -a_{1}y'_{1}(t),$$

$$\frac{dy'_{2}(t)}{dt} = y'_{2}(t)(a_{2} - b_{2}y'_{2}(t)),$$

$$\Delta y'_{1}(t) = D(y'_{2}(t) - y'_{1}(t)),$$

$$\Delta y'_{2}(t) = D(y'_{1}(t) - y'_{2}(t)),$$

$$t = n\tau, \ n = 1, 2,$$

(4.3)

In view of Lemma 3.3 and (3.6), we obtain the τ -periodic solution of system (4.3)

$$\widetilde{y'_{1}(t)} = y'^{*}_{1} e^{-a_{1}(t-n\tau)}, \quad n\tau < t \le (n+1)\tau,$$

$$\widetilde{y'_{2}(t)} = \frac{a_{2} e^{a_{2}(t-n\tau)} y'^{*}_{2}}{a_{2} + b_{2} (e^{a_{2}(t-n\tau)} - 1) y'^{*}_{2}}, \quad n\tau < t \le (n+1)\tau,$$
(4.4)

which is globally asymptotically stable, where $y_1^{\prime *} = De^{a_2 \tau} y_2^{\prime *} / ((1 - D)e^{a_2 \tau} - (1 - 2D)e^{(a_2 - a_1)\tau})$ and $y_2^{\prime *} = a_2((1 - D)(e^{a_2 \tau} + e^{-a_1 \tau}) - (1 - 2D)e^{(a_2 - a_1)\tau} - 1)/b_2(e^{a_2 \tau} - 1)(1 - (1 - D)e^{(a_2 - a_1)\tau}).$ From Lemma 3.3 and comparison theorem of impulsive equation [36], we have $y_1(t) \le y'_1(t)$ and $y'_1(t) \to y'_1(t)$ as $t \to \infty$. Then there exists an integer $k_2 > k_1$, $t > k_2$ such that

$$y_1(t) \le y'_1(t) \le \widetilde{y_1(t)} + \varepsilon_0, \quad n\tau < t \le (n+1)\tau, \ n > k_2,$$
(4.5)

that is,

$$y_1(t) < \widetilde{y_1(t)} + \varepsilon_0 \le y_1^* + \varepsilon_0 \stackrel{\Delta}{=} \varrho, \quad n\tau < t \le (n+1)\tau, \ n > k_2.$$

$$(4.6)$$

From (2.5), we get

$$\frac{dx_2(t)}{dt} \le r e^{-w\tau_1} x_2(t-\tau_1) - (d+\beta \varrho) x_2(t), \quad t > n\tau + \tau_1, \ n > k_2.$$
(4.7)

Consider the following comparison differential system:

$$\frac{dx_2'(t)}{dt} = re^{-\omega\tau_1}x_2'(t-\tau_1) - (d+\beta\varrho)x_2'(t), \quad t > n\tau + \tau_1, \ n > k_2.$$
(4.8)

So we have $re^{-w\tau_1} < d + \beta q$. According to Lemma 3.4, we have $\lim_{t\to\infty} x'_2(t) = 0$.

Let $(x_2(t), y_1(t), y_2(t))$ be the solution of system (2.5) with its initial conditions and $x_2(\zeta) = \varphi_2(\zeta)$ ($\zeta \in [-\tau_1, 0]$), $y'_2(t)$ is the solution of system (4.8) with initial condition $x'_2(\zeta) = \varphi_2(\zeta)$ ($\zeta \in [-\tau_1, 0]$). By the comparison theorem, we have

$$\lim_{t \to \infty} x_2(t) < \lim_{t \to \infty} x_2'(t) = 0.$$
(4.9)

Incorporating into the positivity of $x_2(t)$, we know that $\lim_{t\to\infty} x_2(t) = 0$. Therefore, for any $\varepsilon_1 > 0$ (sufficiently small), there exists an integer k_3 ($k_3\tau > k_2\tau + \tau_1$) such that $x_2(t) < \varepsilon_1$ for all $t > k_3\tau$.

For system (2.5), we have

$$-a_2 y_1(t) \le \frac{dy_1(t)}{dt} \le -(a_2 - k\beta\varepsilon_1)y_1(t).$$

$$(4.10)$$

Then, $z_2(t) \le y_1(t) \le z'_2(t)$ and $z_2(t) \to \widetilde{y_1(t)}, z'_2(t) \to \widetilde{y_1(t)}$ as $t \to \infty$, where $(z_1(t), z_2(t))$ and $(z'_1(t), z'_2(t))$ are the solutions of

$$\frac{dz_1(t)}{dt} = -a_1 z_1(t),$$

$$\frac{dz_2(t)}{dt} = z_2(t) [a_2 - b_2 z_2(t)],$$

(4.11)

$$\begin{split} \Delta z_1(t) &= D(z_2(t) - z_1(t)), \\ \Delta z_2(t) &= D(z_1(t) - z_2(t)), \end{split} \quad t = n\tau, \ n = 1, 2, \dots, \end{split}$$

$$\frac{dz'_{1}(t)}{dt} = -(a_{1} - \beta \varepsilon_{1})z'_{1}(t), \qquad t \neq n\tau,
\frac{dz'_{2}(t)}{dt} = z'_{2}(t)(a_{2} - b_{2}z'_{2}(t)), \qquad (4.12)$$

$$\begin{split} \Delta z_1'(t) &= D\big(z_2'(t) - z_1'(t)\big), \\ \Delta z_2'(t) &= D\big(z_1'(t) - z_2'(t)\big), \end{split} \quad t = n\tau, \ n = 1, 2, \dots, \end{split}$$

respectively. Here

$$\widetilde{z'_{1}(t)} = z_{1}^{\prime*} e^{-(a_{1}-k\beta\varepsilon_{1})(t-n\tau)}, \quad n\tau < t \le (n+1)\tau,$$

$$\widetilde{z'_{2}(t)} = \frac{a_{2}e^{a_{2}(t-n\tau)}z_{2}^{\prime*}}{a_{2}+b_{2}(e^{a_{2}(t-n\tau)}-1)z_{2}^{\prime*}}, \quad n\tau < t \le (n+1)\tau,$$
(4.13)

where

$$z_{1}^{\prime*} = \frac{De^{a_{2}\tau} z_{2}^{\prime*}}{(1-D)e^{a_{2}\tau} - (1-2D)e^{(a_{2}-(a_{1}-k\beta\epsilon_{1}))\tau}},$$

$$z_{2}^{\prime*} = \frac{a_{2}((1-D)(e^{a_{2}\tau} + e^{-(a_{1}-k\beta\epsilon_{1})\tau}) - (1-2D)e^{(a_{2}-(a_{1}-k\beta\epsilon_{1}))\tau} - 1)}{b_{2}(e^{a_{2}\tau} - 1)(1-(1-D)e^{(a_{2}-(a_{1}-k\beta\epsilon_{1}))\tau})}.$$
(4.14)

Therefore, for any $\varepsilon_2 > 0$, there exists an integer k_4 , $n > k_4$ such that $\widetilde{z_2(t)} - \varepsilon_2 < y_1(t) < \widetilde{z'_2(t)} + \varepsilon_2, \widetilde{z_1(t)} - \varepsilon_2 < y_2(t) < \widetilde{z'_1(t)} + \varepsilon_2$. Let $\varepsilon_1 \to 0$. So we have $\widetilde{y_1(t)} - \varepsilon_2 < y_1(t) < \widetilde{y_1(t)} + \varepsilon_2, \widetilde{y_2(t)} - \varepsilon_2 < y_2(t) < \widetilde{y_2(t)} + \varepsilon_2$, for *t* large enough, which implies $y_1(t) \to \widetilde{y_1(t)}$ and $y_2(t) \to \widetilde{y_2(t)}$ as $t \to \infty$. This completes the proof.

The next work is to investigate the permanence of the system (2.3). Before starting the following theorem, we give the following definition.

Definition 4.2. System (2.3) is said to be permanent if there are constants m, M > 0 (independent of initial value) and a finite time T_0 , such that for all solutions $(x_1(t), x_2(t), y_1(t), y_2(t))$ with all initial values $x_1(0^+) > 0$, $x_2(0^+) > 0$, $y_1(0^+) > 0$, $y_2(0^+) > 0$, $m \le x_1(t) \le M/k$, $m \le x_2(t) \le M/k$, $m \le y_1(t) \le M$, $m \le y_2(t) \le M$, holds for all $t \ge T_0$. Here T_0 may depend on the initial values $(x_1(0^+), x_2(0^+), y_1(0^+), y_2(0^+))$.

Theorem 4.3. If

$$a_{2}\tau > \ln \frac{1 - (1 - D)e^{-(a_{1} - \beta x_{2}^{*})\tau}}{1 - (1 - 2D)e^{-(a_{1} - \beta x_{2}^{*})\tau}},$$

$$re^{-w\tau_{1}} > d + \frac{D\beta v_{1}^{*}e^{(a_{1}\tau - w\tau_{1})}}{(1 - D)e^{a_{1}\tau} - (1 - 2D)e^{(a_{1} - (a_{2} + \beta x_{2}^{*}))\tau}},$$
(4.15)

hold, there is a positive constant g such that each positive solution $(x_2(t), y_1(t), y_2(t))$ of (2.5) satisfies $x_2(t) \ge g$ for large enough t. x_2^* can be confirmed by the equation

$$\left(re^{-\omega\tau_{1}}-d\right)\left[(1-D)e^{a_{2}\tau}-(1-2D)e^{(a_{2}-(a_{1}+\beta x_{2}^{*}))\tau}\right]=D\beta v_{1}^{*}e^{a_{2}\tau-\omega\tau_{1}}.$$
(4.16)

Here

$$v_1^* = \frac{a_2((1-D)(e^{a_2\tau} + e^{-(a_1 + \beta x_2^*)\tau}) - (1-2D)e^{(a_2 - (a_1 + \beta x_2^*))\tau} - 1)}{b_2(e^{a_2\tau} - 1)(1 - (1-D)e^{(a_2 - (a_1 + \beta x_2^*))\tau})}.$$
(4.17)

Proof. The fourth equation of (2.5) can be rewritten as

$$\frac{dx_2(t)}{dt} = \left[re^{-\omega\tau_1} - \left(d + \beta y_1(t) \right) \right] x_2(t) - re^{-\omega\tau_1} \frac{d}{dt} \int_{t-\tau_1}^t x_2(u) du.$$
(4.18)

Let us consider any positive solution $(x_2(t), y_1(t), y_2(t))$ of system (2.5). According to (4.18), Q(t) is defined as

$$Q(t) = x_2(t) + re^{-\omega\tau_1} \int_{t-\tau_1}^t x_2(u) du.$$
(4.19)

We calculate the derivative of Q(t) along the solution of (2.5)

$$\frac{dQ(t)}{dt} = \left[re^{-w\tau_1} - \left(d + \beta y_1(t) \right) \right] x_2(t).$$
(4.20)

Since $Dk\beta v_1^* e^{(a_1\tau - w\tau_1)} / ((1 - D)e^{a_1\tau} - (1 - 2D)e^{(a_1 - (a_2 + \beta y_2^*))\tau}) > d$, we can easily know that there exists sufficiently small $\varepsilon > 0$ such that

$$k\beta e^{-w\tau_1}\left(\frac{Dv_1^*e^{a_1\tau}}{(1-D)e^{a_1\tau}-(1-2D)e^{(a_1-(a_2+\beta y_2^*))\tau}}-\varepsilon\right) > d.$$
(4.21)

We claim that for any $t_0 > 0$, it is impossible that $x_2(t) < x_2^*$ for all $t > t_0$. Suppose that the claim is not valid. Then there is a $t_0 > 0$ such that $x_2(t) < x_2^*$ for all $t > t_0$. It follows from the first equation of (2.5) that for all $t > t_0$,

$$\frac{dy_1(t)}{dt} < -(a_2 - k\beta x_2^*)y_1(t).$$
(4.22)

Consider the following comparison an impulsive system for all $t > t_0$

$$\frac{dv_{1}(t)}{dt} = -(a_{1} - k\beta x_{2}^{*})v_{1}(t), \qquad t \neq n\tau,
\frac{dv_{2}(t)}{dt} = v_{2}(t)(a_{2} - b_{2}v_{2}(t)), \qquad (4.23)$$

$$\Delta v_{1}(t) = D(v_{2}(t) - v_{1}(t)), \qquad t = n\tau, \ n = 1, 2, \dots$$

By Lemma 3.2 and the condition $a_2 \tau > \ln((1 - (1 - D)e^{-(a_1 + \beta x_2^*)\tau})/(1 - (1 - 2D)e^{-(a_1 + \beta x_2^*)\tau}))$, we obtain

$$\widetilde{v_{1}(t)} = v_{1}^{*} e^{-(a_{1} + \beta x_{2}^{*})(t - n\tau)}, \quad n\tau < t \le (n+1)\tau,$$

$$\widetilde{v_{2}(t)} = \frac{a_{2} e^{a_{2}(t - n\tau)} v_{2}^{*}}{a_{2} + b_{2} (e^{a_{2}(t - n\tau)} - 1) v_{2}^{*}}, \quad n\tau < t \le (n+1)\tau,$$
(4.24)

where

$$v_{1}^{*} = \frac{De^{a_{2}\tau}v_{2}^{*}}{(1-D)e^{a_{2}\tau} - (1-2D)e^{(a_{2}-(a_{1}-\beta x_{2}^{*}))\tau}},$$

$$v_{2}^{*} = \frac{a_{2}((1-D)(e^{a_{2}\tau} + e^{-(a_{1}-\beta x_{2}^{*})\tau}) - (1-2D)e^{(a_{2}-(a_{1}-\beta x_{2}^{*}))\tau} - 1)}{b_{2}(e^{a_{2}\tau} - 1)(1-(1-D)e^{(a_{2}-(a_{1}-\beta x_{2}^{*}))\tau})},$$
(4.25)

is the unique positive periodic solution of (4.23), which is globally asymptotically stable. By the comparison theorem for impulsive differential equation [36], we know that there exists $t_1(> t_0 + \tau_1)$ such that the inequalities $y_1(t) \le \widetilde{v_1(t)} + \varepsilon$ and $y_2(t) \le \widetilde{v_2(t)} + \varepsilon$ hold for $t \ge t_1$. Thus $y_1(t) \le v_1^* + \varepsilon$ for all $t \ge t_1$. We make notation as $\sigma \stackrel{\Delta}{=} v_1^* + \varepsilon$ for convenience. Also

$$re^{-\omega\tau_1} > d + \beta\sigma. \tag{4.26}$$

Then

$$\frac{dQ(t)}{dt} > x_2(t) \left[re^{-\omega\tau_1} - \left(d + \beta\sigma\right) \right]$$
(4.27)

for all $t > t_1$. Setting $x_2^m = \min_{t \in [t_1, t_1 + \tau_1]} x_2(t)$, we will show that $x_2(t) \ge x_2^m$ for all $t \ge t_1$. Supposing the contrary, then there is a $T_0 > 0$ such that $x_2(t) \ge x_2^m$ for $t_1 \le t \le t_1 + \tau_1 + T_0$, $x_2(t_1 + \tau_1 + T_0) = x_2^m$ and $x'_2(t_1 + \tau_1 + T_0) < 0$. Hence, the first equation of system (2.5) implies that

$$\begin{aligned} x_{2}'(t_{1}+\tau_{1}+T_{0}) &= re^{-w\tau_{1}}x_{2}(t_{1}+T_{0}) - dx_{2}(t_{1}+\tau_{1}+T_{0}) - \beta x_{2}(t_{1}+\tau_{1}+T_{0})y_{1}(t_{1}+\tau_{1}+T_{0})\\ &\geq \left[re^{-w\tau_{1}} - \left(d+\beta\sigma\right)\right]x_{2}^{m} > 0. \end{aligned}$$

$$(4.28)$$

This is a contradiction. Thus, $x_2(t) \ge x_2^m$ for all $t > t_1$. As a consequence, $Q'(t) > x_2^m [re^{-w\tau_1} - (d + \beta\sigma)] > 0$ for all $t > t_1$. This implies that as $t \to \infty$, $Q(t) \to \infty$. It is a contradiction to $Q(t) \le M(1 + \tau_1 re^{-w\tau_1})$. Hence, the claim is complete.

By the claim, we are left to consider two case. First, $x_2(t) \ge x_2^*$ for all t large enough. Second, $x_2(t)$ oscillates about x_2^* for large enough t.

Define

$$g = \min\left\{\frac{x_2^*}{2}, g_1\right\},\tag{4.29}$$

where $g_1 = x_2^* e^{-(d+dM)\tau_1}$. We hope to show that $x_2(t) \ge g$ for all *t* large enough. The conclusion is evident in the first case. For the second case, let $t^* > 0$ and $\xi > 0$ satisfy $x_2(t^*) = x_2(t^*+\xi) = x_2^*$ and $x_2(t) < x_2^*$ for all $t^* < t < t^* + \xi$, where t^* is sufficiently large such that $x_2(t) > \sigma$ for $t^* < t < t^* + \xi$, $x_2(t)$ is uniformly continuous. The positive solutions of (2.5) are ultimately bounded, and $x_2(t)$ is not affected by impulses. Hence, there is a T ($0 < t < \tau_1$ and T is dependent on the choice of t^*) such that $x_2(t^*) > x_2^*/3$ for $t^* < t < t^* + T$. If $\xi < T$, there is nothing to prove. Let us consider the case $T < \xi < \tau_1$. Since $x'_2(t) > -(d + \beta M)x_2(t)$ and $x_2(t^*) = x_2^*$, it is clear that $x_2(t) \ge g_1$ for $t \in [t^*, t^* + \tau_1]$. Then, proceeding exactly as the proof for the above claim, we see that $x_2(t) \ge g_1$ for $t \in [t^* + \tau_1, t^* + \xi]$. Because the kind of interval, $t \in [t^*, t^* + \xi]$ is chosen in an arbitrary way (we only need t^* to be large). We concluded that $x_2(t) \ge g$ for all large t. In the second case, in view of our above discussion, the choice of g is independent of the positive solution, and we proved that any positive solution of (2.5) satisfies $x_2(t) \ge g$ for all sufficiently large t. This completes the proof of this theorem.

Theorem 4.4. If

$$a_{2}\tau > \ln \frac{1 - (1 - D)e^{-(a_{1} - \beta x_{2}^{*})\tau}}{1 - (1 - 2D)e^{-(a_{1} - \beta x_{2}^{*})\tau}},$$

$$re^{-w\tau_{1}} > d + \frac{D\beta v_{1}^{*}e^{(a_{1}\tau - w\tau_{1})}}{(1 - D)e^{a_{1}\tau} - (1 - 2D)e^{(a_{1} - (a_{2} + \beta x_{2}^{*}))\tau}},$$
(4.30)

hold, system (2.3) is permanent. Here

$$v_{1}^{*} = \frac{a_{2}((1-D)(e^{a_{2}\tau} + e^{-(a_{1}+\beta x_{2}^{*})\tau}) - (1-2D)e^{(a_{2}-(a_{1}+\beta x_{2}^{*}))\tau} - 1)}{b_{2}(e^{a_{2}\tau} - 1)(1-(1-D)e^{(a_{2}-(a_{1}+\beta x_{2}^{*}))\tau})}.$$
(4.31)



Figure 1: Dynamical behavior of system (2.5) on prey-extinction periodic solution with $x_2(0) = 5.2$, $y_1(0) = 5.0$, $y_2(0) = 5.6$, r = 1, $\tau_1 = 3$, d = 3, $a_1 = 1$, $a_2 = 1$, $b_2 = 1$, $\beta = 4$, k = 0.9, D = 0.5, w = 1, and $\tau = 1$. (a) Time-series of $x_2(t)$, (b) time-series of $y_1(t)$, (c) time-series of $y_2(t)$.

Proof. Let $(x_1(t), x_2(t), y_1(t), y_2(t))$ be any solution of system (2.3). From system (2.3) and Theorem 4.3, we have

$$\frac{dx_{1}(t)}{dt} = x_{1}(t)(a_{1} - b_{1}x_{1}(t)),$$

$$t \neq n\tau,$$

$$\frac{dx_{2}(t)}{dt} \ge -a_{2}x_{2}(t),$$

$$\Delta x_{1}(t) = D(x_{2}(t) - x_{1}(t)),$$

$$\Delta x_{2}(t) = D(x_{1}(t) - x_{2}(t)),$$

$$t = n\tau, \ n = 1, 2, \dots$$
(4.32)



Figure 2: Dynamical behavior of the permanence of system (2.5) with $x_2(0) = 5.2$, $y_1(0) = 5.0$, $y_2(0) = 5.6$, $\tau_1 = 1$, d = 3, $a_1 = 1$, $a_2 = 1$, $b_2 = 1$, $\beta = 4$, k = 0.9, D = 0.5, w = 1, and $\tau = 1$. (a) Time-series of $x_2(t)$. (b) Time-series of $y_1(t)$. (c) Time-series of $y_2(t)$.

By the same argument as those in the proof of Theorem 4.1, we have that $x_1(t) \ge x_1''* - \varepsilon = p^1$, and $x_2(t) \ge x_1''*e^{-a2\tau} - \varepsilon = p^2$, and $\varepsilon > 0$ is small enough. Here

$$x_1^{\prime\prime*} = \frac{a_2 \big((1-D)(e^{a_2\tau} + e^{-a_1\tau}) - (1-2D)e^{(a_2-a_1)\tau} - 1 \big)}{b_2 (e^{a_2\tau} - 1) \big(1 - (1-D)e^{(a_2-a_1)\tau} \big)},$$
(4.33)

$$x_2^{\prime\prime*} = \frac{De^{a_2\tau}x_1^{\prime\prime*}}{(1-D)e^{a_2\tau} - (1-2D)e^{(a_2-a_1)\tau}}.$$
(4.34)

In view of Theorem 4.1, the third equation of system (2.3) becomes

$$\frac{dy_1(t)}{dt} \ge r(p_2g - Me^{-w\tau_1}) - wy_1(t).$$
(4.35)

It is easy to obtain $y_1(t) \ge \delta - \varepsilon$, and $\varepsilon > 0$ is small enough, where $\delta = r(p_2g - Me^{-w\tau_1})/w - \varepsilon$. By Theorem 4.3 and the above discussion, system (2.3) is permanent. The proof of Theorem 4.4 is complete.

5. Discussion

In this paper, we investigate a delayed predator-prey model with impulsive diffusion on predator and stage structure on prey. We analyze that the prey-extinction periodic solution of system (2.3) is globally attractive, and we also obtain the permanent condition of system (2.3). It is assumed that $x_2(0) = 5.2$, $y_1(0) = 5.0$, $y_2(0) = 5.6$, r = 1, $\tau_1 = 3$, d = 3, $a_1 = 1$, $a_2 = 1$, $b_2 = 1$, $\beta = 4$, k = 0.9, D = 0.5, w = 1, and $\tau = 1$. Then the prey-extinction periodic solution is a global attractivity (see Figure 1). We also assume that $x_2(0) = 5.2$, $y_1(0) = 5.0$, $y_2(0) = 5.6$, r = 1, $\tau_1 = 1$, d = 3, $a_1 = 1$, $b_1 = 1$, $a_2 = 1$, $b_2 = 1$, $\beta = 4$, k = 0.9, D = 0.5, w = 1, then, system (2.5) is permanent (see Figure 2). From Theorems 4.1 and 4.4, we can easily guess that there must exist a threshold τ_1^* . If $\tau_1 > \tau_1^*$, the prey-extinction periodic solution ($\widetilde{x_1(t)}, \widetilde{x_2(t)}, 0, 0$) of (2.3) is globally attractive. If $\tau_1 < \tau_1^*$, system (2.3) is permanent. The results show that the diffusion and the mature time of the prey play important roles for the permanence of system (2.3) and provide tactical basis for the biological resource protection.

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