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Research article

Modelling and analysis of a stochastic nonautonomous predator-prey model with impulsive effects and nonlinear functional response

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Abstract: In this paper, a new stochastic predator-prey model with impulsive perturbation and Crowley-Martin functional response is proposed. The dynamical properties of the model are systematically investigated. The existence and stochastically ultimate boundedness of a global positive solution are derived using the theory of impulsive stochastic differential equations. Some sufficient criteria are obtained to guarantee the extinction and a series of persistence in the mean of the system. Moreover, we provide conditions for the stochastic permanence and global attractivity of the model. Numerical simulations are performed to support our qualitative results.

Keywords: extinction; global attractivity; impulsive effects; persistence; stochastic predator-prey model

1. Introduction

Predator-prey models are highly important in general and mathematical ecology [1]. In the past decades, many factors have been considered to describe the ecological predator-prey system more correctly and reasonably [2,3]. Notably, population models in the real world is inevitably influenced by numerous unpredictable environmental noise, and deterministic systems are fairly challenged in describing the fluctuation accurately [4,5]. Hence, an increasing number of researchers have paid attention to stochastic models and proposed various population models with stochastic perturbations, such as in [6–10]. Liu and Wang [10] introduced a stochastic non-autonomous predator-prey model for one species with white noise as follows:

$$dx(t) = r(t)[x(t) - a(t)x(t)]dt + \sigma(t)x(t)dB(t).$$
(1.1)

The group analyzed the conditions for extinction and species persistence in Eq (1.1). On the basis of the theoretical and practical significance of this stochastic model, many results have been presented, particularly, in [11–13]. However, the influence of the functional response to systems has been rarely considered in previous stochastic population models.

Generally, two types of functional response exist, that is, prey- and predator-dependent responses. The first functional response considers only the prey density, whereas the other accounts for both prey and predator densities [2]. When investigating biological phenomena, one must not ignore the predator's functional response to prey because of such response's effect on dynamical system properties [14-18]. Among many different forms of predator-dependent functional responses, the three classical ones include the Beddington-DeAngelis, Hassell-Varley and Crowley-Martin types. We let $x_1(t)$ and $x_2(t)$ denote the prev and predator population densities, respectively, at time t. Then, $\frac{\omega(t)x_1(t)}{1+a(t)x_1(t)+b(t)x_2(t)+a(t)b(t)x_1(t)x_2(t)}$ becomes the Crowley-Martin functional response, where a(t), b(t) and $\omega(t)$ represent the effects of handling time, the magnitude of interference among predators, and capture rate, respectively. Interestingly, if a(t) = 0 and b(t) = 0, then the Crowley-Martin functional response becomes a linear mass-action functional response. If a(t) = 0 and b(t) > 0, the response represents a saturation response; if a(t) > 0 and b(t) = 0, then the response becomes a Michaelis-Menten functional response (or Holling type-II functional response) [1,19,20]. Given its importance and appeal, some scholars have studied stochastic predator-prey models incorporating Crowley-Martin functional response [21–23] and in this paper, we consider the Crowley-Martin functional response to embody interference among predators and provide insight into the dynamics of the predator-prey population model.

Meanwhile, the theory of impulsive differential equation was well developed recently, and impulsive differential equations were found as a more effective method for describing species and the ecological systems more realistically. Many important and peculiar results have been obtained regarding the dynamical behavior of these systems, including the permanence, extinction of positive solution and dynamical complexity. However, few studies have addressed the population dynamics of two species both with stochastic and impulsive perturbations, except in [24–26]. In [25], Zhang and Tan considered a stochastic autonomous predator-prey model in a polluted environment with impulsive perturbations and analyzed the extinction and persistence of the system. By contrast, the model proposed is autonomous, that is, the parameters are assumed as constants and independent of time. In [26], Wu, Zou, and Wang proposed a stochastic Lotka-Volterra model with impulsive perturbations. The asymptotic properties of the model were examined. However, their model was based on the prey-dependent functional response and did not consider the predator's functional response to prey. In addition, there are four approaches to introduce stochastic perturbations to the model as usual, through time Markov chain model, parameter perturbation, being proportional to the variables, and robusting the positive equilibria of deterministic models [27]. In this paper, we adopt the third approach to include stochastic effects to Eq (1.2).

Inspired by the above discussion and [11,25,26], we consider the possible effects of impulsive and stochastic perturbations on the system and propose the following non-autonomous stochastic differen-

tial equation:

$$\begin{cases} dx_1(t) = x_1 \left(r(t) - k(t)x_1 - \frac{\omega(t)x_2}{1 + a(t)x_1 + b(t)x_2 + a(t)b(t)x_1x_2} \right) dt + x_1\sigma_1(t)dB_1(t), \\ dx_2(t) = x_2 \left(-g(t) - h(t)x_2 + \frac{f(t)x_1}{1 + a(t)x_1 + b(t)x_2 + a(t)b(t)x_1x_2} \right) dt + x_2\sigma_2(t)dB_2(t), \quad t \neq \tau_k, \quad (1.2) \\ x_1(\tau_k^+) = (1 + \rho_{1k})x_1(\tau_k), \\ x_2(\tau_k^+) = (1 + \rho_{2k})x_2(\tau_k), \quad t = \tau_k, k = 1, 2, 3, \dots \end{cases}$$

The parameters are defined as follows: r(t) and g(t) denote the intrinsic growth rate of the prey and predator population at time *t*; k(t) and h(t) are the density-dependent coefficients of prey and predator populations, respectively; f(t) represents the conversion rate of nutrients into the reproductive predator population; $\sigma_i^2(t)$ (i = 1, 2) refers to the intensities of the white noises at time *t*; $\dot{B}_1(t)$ and $\dot{B}_2(t)$ are standard white noises, in particular, $B_1(t)$, $B_2(t)$ are Brownian motions defined on a complete probability space ($\Omega, \mathcal{F}, \mathbb{P}$) [6].

Throughout this paper, all the coefficients are assumed to be positive and continuously bounded on $\mathbb{R}_+ = [0, +\infty)$. The impulsive points satisfy $0 < \tau_1 < \tau_2 < \ldots < \tau_k < \ldots$ and $\lim_{k \to +\infty} \tau_k = +\infty$. According to biological meanings, $\rho_{1k} > -1$, $\rho_{2k} > -1$. Moreover, we assume that there are some positive constants m, M, \tilde{m} and \tilde{M} that satisfy $0 < m \leq \prod_{0 < \tau_k < t} (1 + \rho_{1k}) \leq M$ and $0 < \tilde{m} \leq \prod_{0 < \tau_k < t} (1 + \rho_{2k}) \leq m$

 \widetilde{M} , for all t > 0.

The remaining portion of this paper is arranged as follows. In the next section, some preliminaries are introduced. We analyze the impulsive stochastic differential model and obtain the existence, uniqueness and stochastically ultimate boundness of the positive solution in Section 3. In Section 4, sufficient conditions for extinction and a set of persistence in the mean, including non-persistence, weak persistence, and strong persistence in the mean, are presented. Additionally, we provide conditions to guarantee the stochastic permanence of the system. In Section 5, the global attractiveness of Eq (1.2) is studied. Finally, some numerical simulations, which verify our theoretical results, are given in Section 6. We compare the results of stochastic models under positive or negative impulsive perturbations with those without such disturbances, as well as, the figures with different stochastic perturbations and same impulse. By doing so, we clearly show that the impulsive and stochastic perturbations are of great importance to species permanence and extinction.

2. Preliminaries

To proceed, we list some appropriate definitions, notations, and lemmas as follows. For convenience, we denote

$$f^{u} = \sup_{t \ge 0} f(t), \quad f^{l} = \inf_{t \ge 0} f(t), \quad \langle f(t) \rangle = \frac{1}{t} \int_{0}^{t} f(s) ds, \quad f^{*} = \limsup_{t \to +\infty} f(t), \quad f_{*} = \liminf_{t \to +\infty} f(t),$$

where f(t) is a continuous and bound function defined on $[0, +\infty)$. X(t) represents $(x_1(t), x_2(t))$ and $|X(t)| = (x_1^2(t) + x_2^2(t))^{\frac{1}{2}}$. \mathbb{N} is the set of positive integers and $\mathbb{R}^n_+ = \{x \in \mathbb{R}^n : x_i > 0, i = 1, 2, 3...n\}$. **Definition 2.1.**

- (a) If $\lim x(t) = 0$ a.s., then the species x(t) is said to go to extinction.
- (b) If $\langle x \rangle^* = 0$ a.s., then the population x(t) is said to be non-persistent in the mean.
- (c) If $\langle x \rangle^* > 0$ a.s., then the population x(t) is said to be weakly persistent in the mean.
- (d) If $\langle x \rangle_* > 0$ a.s., then the population x(t) is said to be strongly persistent in the mean.
- (e) If $x^* > 0$ a.s., then the population x(t) is said to be weakly persistent.

Definition 2.2 ([28]) Solution $X(t) = (x_1(t), x_2(t))$ of Eq (1.2) is said to be stochastically ultimately bound, if for arbitrary $\varepsilon \in (0, 1)$, a positive constant $\delta = \delta(\varepsilon)$ exists, such that for any given initial value $X_0 = (x_1(0), x_2(0)) \in \mathbb{R}^2_+$, the solution X(t) to Eq (1.2) satisfies $\limsup P\{|X(t)| > \delta\} < \varepsilon$.

Definition 2.3 ([28]) Solution $X(t) = (x_1(t), x_2(t))$ of Eq (1.2) is said to be stochastically permanent, if for any $\varepsilon \in (0, 1)$, there is a pair of positive constants $\delta = \delta(\varepsilon)$ and $\chi = \chi(\varepsilon)$ such that for any initial value $X_0 = (x_1(0), x_2(0)) \in \mathbb{R}^2_+$, the solution X(t) to Eq (1.2) satisfies $\liminf_{t \to \infty} P\{|X(t)| \ge \delta\} \ge 1 - \varepsilon$, $\liminf_{t \to \infty} P\{|X(t)| \le \chi\} \ge 1 - \varepsilon$.

Definition 2.4. Eq (1.2) is said to be globally attractive if

$$\lim_{t \to +\infty} |x_1(t) - \overline{x}_1(t)| = 0, \lim_{t \to +\infty} |x_2(t) - \overline{x}_2(t)| = 0,$$

for any two solutions $(x_1(t), x_2(t))$, $(\overline{x}_1(t), \overline{x}_2(t))$ of Eq (1.2). **Definition 2.5.** ([29]) Consider the impulsive stochastic equation

$$dx(t) = F(t, x(t))dt + G(t, x(t))dB(t), \quad t \neq t_k, t > 0,$$

$$x(t_k^+) - x(t_k) = \alpha_k x(t_k), \quad t = t_k, k = 1, 2, \cdots$$
(2.1)

with the initial value $x(0) = x_0 \in \mathbb{R}^n$. A stochastic process $x(t) = (x_1(t), x_2(t), \dots, x_n(t))^T$, $t \in [0, +\infty)$ is the solution of Eq (2.1) if

(a) x(t) is \mathcal{F}_t adapted and continuous on $(0, t_1)$ and each interval $(t_k, t_{k+1}), k \in \mathbb{N}$ and $F(t, x(t)) \in L^1(\mathbb{R}^+, \mathbb{R}^n), G(t, x(t)) \in L^2(\mathbb{R}^+, \mathbb{R}^n)$.

(b) For each t_k , $x(t_k^+) = \lim_{t \to t_k^+} x(t)$ and $x(t_k^-) = \lim_{t \to t_k^-} x(t)$ and $x(t_k) = x(t_k^-)$ a.s..

(c) x(t) obeys the equivalent integral Eq (2.1) for almost every $t \in \mathbb{R}_+ \setminus t_k$ and satisfies the impulsive conditions at $t = t_k$ a.s..

Lemma 2.1. ([5]) Suppose that $x(t) \in \mathbb{C}[\Omega \times \mathbb{R}_+, \mathbb{R}^0_+]$, where $\mathbb{R}^0_+ = (0, +\infty)$ and $B_i(t)$ (i = 1, 2, 3, ..., n) are independent Brownian motions defined on a complete probability space $(\Omega, \mathcal{F}, \mathbb{P})$, then

(a) If there are positive constants λ_0 , T and $\lambda \ge 0$ satisfying

$$lnx(t) \leq \lambda t - \lambda_0 \int_0^t x(s) ds + \sum_{i=1}^n \beta_i B_i(t),$$

for all $t \ge T$, where β_i is a constant, $1 \le i \le n$, then, $\langle x \rangle^* \le \lambda / \lambda_0$ a.s.

(b) If there are positive constants λ_0 , T and $\lambda \ge 0$ satisfying

$$lnx(t) \geq \lambda t - \lambda_0 \int_0^t x(s) ds + \sum_{i=1}^n \beta_i B_i(t),$$

for all $t \ge T$, where β_i is a constant, $1 \le i \le n$, then, $\langle x \rangle_* \ge \lambda/\lambda_0$ a.s. **Lemma 2.2.** ([30]) Let f be a non-negative function defined on \mathbb{R}_+ such that f is integrate and is uniformly continuous. Then $\lim_{t \to +\infty} f(t) = 0$.

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3. Properties of the solution

In this section, the existence, uniqueness, and stochastically ultimate boundedness of the global positive solution are obtained.

Firstly, we denote

$$x_1(t) = \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1(t), \quad x_2(t) = \prod_{0 < \tau_k < t} (1 + \rho_{2k}) y_2(t),$$

$$\lambda(t) = 1 + a(t) \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 + b(t) \prod_{0 < \tau_k < t} (1 + \rho_{2k}) y_2 + a(t) b(t) \prod_{0 < \tau_k < t} (1 + \rho_{1k}) \prod_{0 < \tau_k < t} (1 + \rho_{2k}) y_1 y_2,$$

then by virtue of Lemma 2.1 in [30], the following lemma can be obtained. **Lemma 3.1.** *For the stochastic equations without impulses*

$$dy_{1}(t) = y_{1}\left(r(t) - k(t)\prod_{0 < \tau_{k} < t} (1 + \rho_{1k})y_{1} - \frac{\omega(t)}{\lambda(t)}\prod_{0 < \tau_{k} < t} (1 + \rho_{2k})y_{2}\right)dt + y_{1}\sigma_{1}(t)dB_{1}(t),$$

$$dy_{2}(t) = y_{2}\left(-g(t) - h(t)\prod_{0 < \rho_{2k} < t} (1 + \rho_{1k})y_{2} + \frac{f(t)}{\lambda(t)}\prod_{0 < \tau_{k} < t} (1 + \rho_{1k})y_{1}\right)dt + y_{2}\sigma_{2}(t)dB_{2}(t),$$
(3.1)

 $(y_1(t), y_2(t))$ is a solution of Eq (3.1) if and only if $(x_1(t), x_2(t))$ is a solution of Eq (1.2) with initial value $(x_1(0), x_2(0)) = (y_1(0), y_2(0)).$

The proof can be given easily as in [31], but such approach is not applied herein. **Theorem 3.1** For any given value $(x_1(0), x_2(0)) = X_0 \in \mathbb{R}^2_+$, a unique solution $(x_1(t), x_2(t))$ exists for Equation (1.2) on $t \ge 0$ and the solution will remain in \mathbb{R}^2_+ with probability one.

The proof of Theorem 3.1 is standard and is presented in Appendix A. **Theorem 3.2** *The solutions of Eq (1.2) are stochastically ultimately bounded for any initial value* $X_0 = (x_1(0), x_2(0)) \in \mathbb{R}^2_+$.

The proof of Theorem 3.2 is presented in Appendix A.

4. Long time behavior of Eq (1.2)

In this section, sufficient conditions for extinction and a series of persistence in the mean, such as non-persistence, weak persistence and strong persistence in the mean, are established. Furthermore, we obtain conditions to guarantee the stochastic permanence of the system. Before giving the main theorems, we introduce a lemma essential to our proofs.

Lemma 4.1. If $\limsup_{t\to\infty} \frac{\prod\limits_{t\to\infty}^{(1+\rho_{1k})}}{\ln t} < \infty$, $\limsup_{t\to\infty} \frac{\prod\limits_{t\to\infty}^{(1+\rho_{2k})}}{\ln t} < \infty$ hold, then for any initial value $(x_1(0), x_2(0)) \in \mathbb{R}^2_+$, the solution $X_1(t) = (x_1(t), x_2(t))$ of Eq (1.2) satisfies

$$\limsup_{t \to \infty} \frac{\ln x_1(t)}{t} \le 0, \quad \limsup_{t \to \infty} \frac{\ln x_2(t)}{t} \le 0, \qquad a.s$$

The proof of Lemma 4.1 is given in Appendix A.

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4.1. Extinction and persistence of the system

The results about persistence in the mean and extinction of the prey and predator populations are presented in Theorems 4.1.1 and 4.1.2.

- **Theorem 4.1.1** For the prey population x_1 of Eq (1.2), we have (a) If $\hat{r}_1 < 0$, then the prey population x_1 is extinct with probability 1, where $r_1(t) = r(t) 0.5\sigma_1^2(t)$, $\widehat{r}_{1} = \limsup_{t \to +\infty} \frac{1}{t} \left[\sum_{0 < \tau_{k} < t} ln(1 + \rho_{1k}) + \int_{0}^{t} r_{1}(s) ds \right].$ (b) If $\widehat{r}_{1} = 0$, then the prey population x_{1} is non-persistent in the mean with probability 1.
- (c) If $\hat{r}_1 > 0$ and $\hat{r}_2 < 0$, then the prey population x_1 is weakly persistent in the mean with probability *I*, where $\widehat{r}_2 = \limsup_{t \to +\infty} \frac{1}{t} [\sum_{0 < \tau_k < t} ln(1 + \rho_{2k}) + \int_0^t r_2(s) ds].$ (d) If $\check{r}_1 - \langle \frac{\omega}{b} \rangle^* > 0$, then the prey population x_1 is strongly persistent in the mean with probability 1,
- where $\check{r}_1 = \liminf_{t \to +\infty} \frac{1}{t} [\sum_{0 < \tau_k < t} ln(1 + \rho_{1k}) + \int_0^t r_1(s) ds].$
- (e) If $\widehat{r_1} > 0$, the prey population $x_1(t)$ holds a superior bound in time average, that is, $\langle x_1(t) \rangle^* \leq \frac{\widehat{r_1}}{k^l} \triangleq$ M_{x} .

Proof. (a) According to $It\hat{o}$'s formula and Eq (3.1), the function can be expressed as

$$dlny_{1} = \left[r(t) - k(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{1k}) y_{1} - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_{k} < t} (1 + \rho_{2k}) y_{2} \right] dt - 0.5\sigma_{1}^{2}(t) dt + \sigma_{1}(t) dB_{1}(t)$$

$$= \left[r_{1}(t) - k(t) x_{1} - \frac{\omega(t) x_{2}}{1 + a(t) x_{1} + b(t) x_{2} + a(t) b(t) x_{1} x_{2}} \right] dt + \sigma_{1}(t) dB_{1}(t),$$

$$dlny_{2} = \left[-g(t) - h(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{2k}) y_{2} + \frac{f(t)}{\lambda(t)} \prod_{0 < \tau_{k} < t} (1 + \rho_{1k}) y_{1} \right] dt - 0.5\sigma_{2}^{2}(t) dt + \sigma_{2}(t) dB_{2}(t)$$

$$= \left[r_{2}(t) - h(t) x_{2} + \frac{f(t) x_{1}}{1 + a(t) x_{1} + b(t) x_{2} + a(t) b(t) x_{1} x_{2}} \right] dt + \sigma_{2}(t) dB_{2}(t).$$

$$(4.1)$$

Taking integral on both sides of Eq (4.1) results in

$$\begin{aligned} \ln y_1(t) - \ln y_1(0) &= \int_0^t r_1(s) ds - \int_0^t k(s) x_1(s) ds - \int_0^t \frac{\omega(s) x_2(s)}{1 + a(s) x_1(s) + b(s) x_2(s) + a(s) b(s) x_1(s) x_2(s)} ds + N_1(t), \\ \ln y_2(t) - \ln y_2(0) &= \int_0^t r_2(s) ds - \int_0^t h(s) x_2(s) ds + \int_0^t \frac{f(s) x_1(s)}{1 + a(s) x_1(s) + b(s) x_2(s) + a(s) b(s) x_1(s) x_2(s)} ds + N_2(t). \end{aligned}$$

We let $N_1(t) = \int_0^t \sigma_1(s) dB_1(s)$, $N_2(t) = \int_0^t \sigma_2(s) dB_2(s)$, where $N_i(t)(i = 1, 2)$ is a local martingale with a quadratic variation satisfying $\langle N_1, N_1 \rangle_t = \int_0^t \sigma_1^2(s) ds \le (\sigma_1^u)^2 t$, $\langle N_2, N_2 \rangle_t = \int_0^t \sigma_2^2(s) ds \le (\sigma_2^u)^2 t$. Using the strong law of large numbers for martingales, we show that

$$\limsup_{t \to \infty} \frac{N_i(t)}{t} = 0, \quad a.s. \quad i = 1, 2.$$
(4.2)

Thus

$$\frac{\ln x_1(t) - \ln x_1(0)}{t} = \frac{\sum\limits_{0 < \tau_k < t} \ln(1 + \rho_{1k})}{-\frac{1}{t} \int_0^t k(s) x_1(s) ds} + \frac{\ln y_1(t) - \ln y_1(0)}{t} = \frac{\sum\limits_{0 < \tau_k < t} \ln(1 + \rho_{1k}) + \int_0^t r_1(s) ds}{\frac{\omega(s) x_2(s)}{t} + \frac{\omega(s) x_2(s)}{t} ds}$$

$$\frac{\ln x_2(t) - \ln x_2(0)}{t} = \frac{\sum_{0 < \tau_k < t} \ln(1 + \rho_{2k})}{-\frac{1}{t} \int_0^t h(s) x_2(s) ds} + \frac{\ln y_2(t) - \ln y_2(0)}{t} = \frac{\sum_{0 < \tau_k < t} \ln(1 + \rho_{2k}) + \int_0^t r_2(s) ds}{\frac{f(s) x_1(s)}{t} + \frac{1}{t} \int_0^t \frac{f(s) x_1(s)}{1 + a(s) x_1(s) + b(s) x_2(s) + a(s) b(s) x_1(s) x_2(s)}} ds.$$
(4.3)

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Therefore,

$$\frac{\ln x_1(t) - \ln x_1(0)}{t} \le \frac{\sum\limits_{0 < \tau_k < t} \ln(1 + \rho_{1k}) + \int_0^t r_1(s) \mathrm{d}s}{t} + \frac{N_1(t)}{t}.$$
(4.4)

Making use of Eq (4.2) and the superior limit as $t \to \infty$ in Eq (4.4), we get $\left[\frac{\ln x_1(t) - \ln x_1(0)}{t}\right]^* \le \hat{r}_1 < 0$ which results in $\lim x_1(t) = 0$.

(b). According to the definition of superior limit and Eq (4.2), for an arbitrary $\varepsilon > 0$, there is a T > 0 satisfying $\frac{1}{t} \left[\sum_{0 < \tau_k < t} \ln(1 + \rho_{1k}) + \int_0^t r_1(s) ds \right] \le \hat{r}_1 + \frac{\varepsilon}{2}$, $\frac{N_1(t)}{t} \le \frac{\varepsilon}{2}$ for all t > T. Substituting the above inequalities into the first equation of (4.4), we easily show that

$$\frac{\ln x_1(t) - \ln x_1(0)}{t} \le \widehat{r_1} - k_* \langle x_1 \rangle + \varepsilon \le \varepsilon - k_* \langle x_1 \rangle.$$
(4.5)

By virtue of Lemma 2.1, we obtain $\langle x_1(t) \rangle^* \leq \frac{\varepsilon}{k_*}$. In accordance with the arbitrariness of ε , we achieve the result.

(c). By virtue of Eq (4.2), superior limit and Lemma 4.1, we show that

$$k^{u}\langle x_{1}\rangle^{*} + \omega^{u}\langle x_{2}\rangle^{*} \ge \left[\frac{\ln x_{1}(t) - \ln x_{1}(0)}{t}\right]^{*} + \langle k(t)x_{1}\rangle^{*} + \langle \frac{\omega(t)x_{2}}{1 + a(t)x_{1} + b(t)x_{2} + a(t)b(t)x_{1}x_{2}}\rangle^{*}$$

$$\ge \widehat{r_{1}} > 0.$$
(4.6)

Thus, $\langle x_1(t) \rangle^* > 0$ a.s. By reduction to absurdity, we can assume that for any $v \in \{\langle x_1(t, v) \rangle^* = 0\}$, by Equation (4.6), we obtain $\langle x_2(t, v) \rangle^* > 0$. Meanwhile, using the superior limit for the second equation of (4.3) and $\langle x_1(t, v) \rangle^* = 0$ leads to

$$\left[\frac{\ln x_2(t,\upsilon) - \ln x_2(0)}{t}\right]^* \le \widehat{r}_2 + f^u \langle x_1(t,\upsilon) \rangle^* + h^l \langle -x_2(t,\upsilon) \rangle^* < 0.$$

Therefore, $\lim x_2(t, v) = 0$. This expression is a contradiction. Then, $\langle x_1(t) \rangle^* > 0$ a.s.

(d). Under the condition $\check{r}_1 - \langle \frac{\omega}{b} \rangle^* > 0$, an arbitrary $\varepsilon > 0$ satisfying $\check{r}_1 - \langle \frac{\omega}{b} \rangle^* - \varepsilon > 0$ exists. According to the definition of superior limit, interior limit and Eq (4.2), for the above-mentioned positive constant ε , there is a T > 0 satisfying $\frac{1}{t} (\sum_{0 < \tau_k < t} \ln(1 + \rho_{1k}) + \int_0^t r_1(s) ds) > \check{r}_1 - \frac{\varepsilon}{3}, \langle \frac{\omega}{b} \rangle < \langle \frac{\omega}{b} \rangle^* + \frac{\varepsilon}{3}, \frac{N_1(t)}{t} > -\frac{\varepsilon}{3}$, for all t > T. Then, from Eq (4.3),

$$\frac{\ln x_1(t) - \ln x_1(0)}{t} \ge \check{r}_1 - \langle \frac{\omega}{b} \rangle^* - \varepsilon - k^{\mu} \langle x_1 \rangle.$$

Using Lemma 2.1 and the arbitrariness of ε , we have that

$$\langle x_1(t) \rangle_* \ge \frac{\check{r}_1 - \langle \frac{\omega}{b} \rangle^*}{k^u} \triangleq m_x > 0.$$
(4.7)

(e). Passing to the first equation of (4.3), we yield

$$\frac{\ln x_1(t) - \ln x_1(0)}{t} \le \frac{1}{t} \left(\sum_{0 < \tau_k < t} \ln(1 + \rho_{1k}) + \int_0^t r_1(s) \mathrm{d}s \right) - k^l \langle x_1(t) \rangle + \frac{N_1(t)}{t}.$$
(4.8)

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Thus, $\langle x_1(t) \rangle^* \leq \frac{\widehat{r}_1}{k^l} \triangleq M_x$, which is obtained by a similar process in the proof of conclusion (2) and is omitted.

Let $(\bar{x}_1(t), \bar{x}_2(t))$ be the solution of the following comparison equation

$$d\bar{x}_{1} = \bar{x}_{1} \left(r(t) - k(t)\bar{x}_{1} \right) dt + \bar{x}_{1}\sigma_{1}(t) dB_{1}(t),$$

$$d\bar{x}_{2} = \bar{x}_{2} \left(-h(t)\bar{x}_{2} + \frac{f(t)}{a(t)} \right) dt + \bar{x}_{2}\sigma_{2}(t) dB_{2}(t).$$
(4.9)

with initial value $(x_1(0), x_2(0)) \in \mathbb{R}^2_+$, then we hold the following theorem.

Theorem 4.1.2. For the predator population x_2 of Eq (1.2),

- (a) if $k_* \hat{r}_2 + f^* \hat{r}_1 < 0$, then the predator population x_2 is extinct with probability 1;
- (b) if $k_* \hat{r}_2 + f^* \hat{r}_1 = 0$, then the predator population x_2 is non-persistent in the mean with probability 1:
- (c) if $\hat{r}_2 + \langle \frac{f\bar{x}_1}{1+a\bar{x}_1+b\bar{x}_2+ab\bar{x}_1\bar{x}_2} \rangle^* > 0$, then the predator population x_2 is weakly persistent in the mean with probability 1;
- (d) if $\widehat{r_2} + \langle \frac{f}{a} \rangle^* > 0$, then the predator population $x_2(t)$ has a superior bound in time average, that is, $\langle x_2(t) \rangle^* \leq \frac{\widehat{r}_2 + \langle \frac{f}{a} \rangle^*}{h^l} \triangleq M_y;$ (e) if $\widehat{r}_2 > 0, \widehat{r}_1 \leq 0$, then the predator population x_2 is weakly persistent.

Proof. (a). Case I. If $\hat{r}_1 \leq 0$, then by virtue of Theorem 4.1.1, we obtain $\langle x_1(t) \rangle^* = 0$. According to the definition of superior limit, for an arbitrary $\varepsilon > 0$, there is a T > 0 satisfying $\frac{1}{t} [\sum \ln(1 + \rho_{2k}) + \rho_{2k}]$

 $\int_{0}^{t} r_{2}(s) ds] < \hat{r}_{2} + \frac{\varepsilon}{2}, \frac{N_{2}(t)}{t} < \frac{\varepsilon}{2} \text{ for all } t > T. \text{ By the second equation of (4.3), we noted}$

$$\left[\frac{\ln x_2(t) - \ln x_2(0)}{t}\right]^* \le \widehat{r}_2 + f^* \langle x_1(t) \rangle^* + \varepsilon = \widehat{r}_2 + \varepsilon < 0,$$

then $\lim_{t\to\infty} x_2(t) = 0.$

Case II. If $\hat{r}_1 > 0$, by Eq (4.3), for the above constant $\varepsilon > 0$, there is a $T_1 > 0$ such that $\frac{\ln x_1(t) - \ln x_1(0)}{t} \le t$ $\widehat{r}_1 - k_* \langle x_1(t) \rangle + \varepsilon$, for all $t > T_1$. Applying Lemma 2.1, we show that

$$\langle x_1(t) \rangle^* \le \frac{\widehat{r_1} + \varepsilon}{k_*}.$$
 (4.10)

Substituting the above inequality into the second equation of (4.3) and using the arbitrariness of ε , we yield

$$\left[\frac{\ln x_2(t) - \ln x_2(0)}{t}\right]^* \le \widehat{r}_2 + f^* \langle x_1(t) \rangle^* \le \frac{k_* \widehat{r}_2 + f^* (\widehat{r}_1 + \varepsilon)}{k_*} < 0.$$
(4.11)

Thus, $\lim_{t \to \infty} x_2(t) = 0$.

(b). In (1), we prove that if $\hat{r}_1 \leq 0$, then $\lim_{t \to \infty} x_2(t) = 0$, consequently, $\langle x_2(t) \rangle^* = 0$. At this point, we only need to show that if $\widehat{r}_1 > 0$, then $\langle x_2(t) \rangle^* = 0$ is also valid. Otherwise, $\langle x_2(t) \rangle^* > 0$, and by Lemma 4.1, we obtain that $\left[\frac{\ln x_2(t)}{t}\right]^* = 0$. From Eq (4.11), we note that

$$0 = \left[\frac{\ln x_2(t) - \ln x_2(0)}{t}\right]^* \le \widehat{r}_2 + f^* \langle x_1(t) \rangle^*$$

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Meanwhile, for any constant $\varepsilon > 0$, there is a T > 0 satisfying $\frac{1}{t} \left[\sum_{0 < \tau_k < t} \ln(1 + \rho_{2k}) + \int_0^t r_2(s) ds \right] < \widehat{r}_2 + \frac{\varepsilon}{3}, \langle f(t)x_1(t) \rangle \le f^* \langle x_1(t) \rangle^* + \frac{\varepsilon}{3}, \frac{N_2(t)}{t} \le \frac{\varepsilon}{3}$ for all t > T. By the second equation of (4.3),

$$\frac{\ln x_2(t) - \ln x_2(0)}{t} \leq \frac{1}{t} \left[\sum_{0 < \tau_k < t} \ln(1 + \rho_{2k}) + \int_0^t r_2(s) \mathrm{d}s \right] + \langle f(t)x_1(t) \rangle - \langle h(t)x_2(t) \rangle + \frac{N_2(t)}{t} \\ \leq \widehat{r_2} + f^* \langle x_1(t) \rangle^* + \varepsilon - h_* \langle x_2(t) \rangle.$$

Then, making use of Lemma 2.1, we achieve $\langle x_2(t) \rangle^* \leq \frac{\widehat{r}_2 + f^* \langle x_1(t) \rangle^* + \varepsilon}{h_*}$, which indicates that $\langle x_2(t) \rangle^* \leq \frac{\widehat{r}_2 + f^* \langle x_1(t) \rangle^*}{h_*}$. By virtue of Eq (4.10) and the arbitrariness of ε , we obtain $\langle x_2(t) \rangle^* \leq \frac{k_* \widehat{r}_2 + f^* \widehat{r}_1}{h_* k_*} = 0$. This is a contradiction. Therefore, $\langle x_2(t) \rangle^* = 0$ a.s.

(c). In the following, we show that $\langle x_2(t) \rangle^* > 0$ a.s.. By reduction to absurdity, for arbitrary $\varepsilon_1 > 0$ and initial value $(x_1(0), x_2(0)) \in \mathbb{R}^2_+$, there is a solution $(\check{x}_1(t), \check{x}_2(t))$ of Eq (1.2) satisfying $P\{\langle \check{x}_2(t) \rangle^* < \varepsilon_1\} > 0$. Let ε_1 be sufficiently small that

$$\widehat{r_2} + \left\langle \frac{f\bar{x}_1}{1 + a\bar{x}_1 + b\bar{x}_2 + ab\bar{x}_1\bar{x}_2} \right\rangle^* > 2\left(\frac{f^u\omega^u}{k^l} + h^u + 1\right)\varepsilon_1.$$

From the second equation of (4.3), it can be shown that

$$\frac{\ln\check{x}_{2}(t) - \ln x_{2}(0)}{t} = \frac{1}{t} \Big[\sum_{0 < \tau_{k} < t} \ln(1 + \rho_{2k}) + \int_{0}^{t} r_{2}(s) ds \Big] + \left\langle \frac{f(t)\bar{x}_{1}}{1 + a(t)\bar{x}_{1} + b(t)\bar{x}_{2} + a(t)b(t)\bar{x}_{1}\bar{x}_{2}} \right\rangle + \frac{N_{2}(t)}{t} + \left\langle \frac{f(t)\check{x}_{1}}{1 + a(t)\check{x}_{1} + b(t)\bar{x}_{2} + a(t)b(t)\bar{x}_{1}\bar{x}_{2}} - \frac{f(t)\bar{x}_{1}}{1 + a(t)\bar{x}_{1} + b(t)\bar{x}_{2} + a(t)b(t)\bar{x}_{1}\bar{x}_{2}} \right\rangle - \left\langle h(t)\check{x}_{2}(t) \right\rangle.$$

Herein, $\check{x}_1(t) \leq \bar{x}_1(t)$, $\check{x}_2(t) \leq \bar{x}_2(t)$, a.s. for $t \in [0, +\infty)$. Note that

$$\frac{f(t)\check{x}_{1}}{1+a(t)\check{x}_{1}+b(t)\check{x}_{2}+a(t)b(t)\check{x}_{1}\check{x}_{2}} - \frac{f(t)\bar{x}_{1}}{1+a(t)\bar{x}_{1}+b(t)\bar{x}_{2}+a(t)b(t)\bar{x}_{1}\bar{x}_{2}} = f(t)\frac{-(\bar{x}_{1}-\check{x}_{1})+a(t)b(t)\bar{x}_{1}\check{x}_{1}(\bar{x}_{2}-\check{x}_{2})+b(t)\check{x}_{1}(\bar{x}_{2}-\check{x}_{2})-b(t)\check{x}_{2}(\bar{x}_{1}-\check{x}_{1})}{(1+a(t)\check{x}_{1}+b(t)\check{x}_{2}+a(t)b(t)\check{x}_{1}\check{x}_{2})(1+a(t)\bar{x}_{1}+b(t)\bar{x}_{2}+a(t)b(t)\bar{x}_{1}\bar{x}_{2})} \ge -2f(t)(\bar{x}_{1}-\check{x}_{1}),$$

then

$$\frac{\ln\check{x}_{2}(t) - \ln x_{2}(0)}{t} \geq \frac{1}{t} \left[\sum_{0 < \tau_{k} < t} \ln(1 + \rho_{2k}) + \int_{0}^{t} r_{2}(s) ds \right] - \langle h(t)\check{x}_{2}(t) \rangle + \frac{N_{2}(t)}{t} + \left\langle \frac{f(t)\bar{x}_{1}}{1 + a(t)\bar{x}_{1} + b(t)\bar{x}_{2} + a(t)b(t)\bar{x}_{1}\bar{x}_{2}} \right\rangle - 2\langle f(t)(\bar{x}_{1} - \check{x}_{1}) \rangle.$$
(4.12)

Construct the Lyapunov function $V_3(t) = |\ln \bar{x}_1(t) - \ln \check{x}_1(t)|$, where $V_3(t)$ is a positive function on \mathbb{R}_+ . By virtue of Itô's formula and Eq (4.9), we achieve the following expression:

$$D^{+}V_{3}(t) \leq \operatorname{sgn}(\bar{x}_{1} - \check{x}_{1}) \Big\{ -k(\bar{x}_{1} - \check{x}_{1}) + \frac{\omega(t)\check{x}_{2}}{1 + a(t)\check{x}_{1} + b(t)\check{x}_{2} + a(t)b(t)\check{x}_{1}\check{x}_{2}} \Big\}.$$
(4.13)

Moreover, integrating the above inequality from 0 to t and dividing by t on both sides of the above inequality result in $\frac{V_3(t)-V_3(0)}{t} \le \omega^u \langle \check{x}_2(t) \rangle - k^l \langle |\bar{x}_1 - \check{x}_1| \rangle$. Then we achieve

$$\langle \bar{x}_1 - \check{x}_1 \rangle \leq \frac{\omega^u}{k^l} \langle \check{x}_2(t) \rangle.$$

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By substituting the above inequality into Eq (4.12) and taking the superior limit of the inequality, we obtain

$$\left[\frac{\ln\check{x}_{2}(t) - \ln x_{2}(0)}{t}\right]^{*} \geq \widehat{r}_{2} - h^{u}\langle\check{x}_{2}(t)\rangle_{*} + \frac{N_{2}(t)}{t} + \left\langle\frac{f(t)\bar{x}_{1}}{1 + a(t)\bar{x}_{1} + b(t)\bar{x}_{2} + a(t)b(t)\bar{x}_{1}\bar{x}_{2}}\right\rangle^{*} - 2\frac{f^{u}\omega^{u}}{k^{l}}\langle\check{x}_{2}(t)\rangle^{*} \\
\geq \widehat{r}_{2} + \left\langle\frac{f(t)\bar{x}_{1}}{1 + a(t)\bar{x}_{1} + b(t)\bar{x}_{2} + a(t)b(t)\bar{x}_{1}\bar{x}_{2}}\right\rangle^{*} - 2\left(\frac{f^{u}\omega^{u}}{k^{l}} + h^{u} + 1\right)\varepsilon_{1} > 0,$$
(4.14)

Equation (4.14) contradicts Lemma 4.1, therefore $\langle x_2(t) \rangle^* > 0$ a.s. The proof is hence completed.

(d). By the second equation of (4.3), we obtain the following equation

$$\frac{\ln x_2(t) - \ln x_2(0)}{t} \le \widehat{r}_2 + \langle \frac{f}{a} \rangle^* - h^l \langle x_2(t) \rangle + \frac{N_2(t)}{t}.$$
(4.15)

Moreover, from the definition of superior limit and Eq (4.2), for the given positive number ε , there is a $T_2 > 0$ satisfying $\frac{1}{t} \left[\sum_{0 < \tau_k < t} \ln(1 + \rho_{2k}) + \int_0^t r_2(s) ds \right] < \hat{r}_2 + \frac{\varepsilon}{3}, \langle \frac{f}{a} \rangle < \langle \frac{f}{a} \rangle^* + \frac{\varepsilon}{3}, \frac{N_2(t)}{t} < \frac{\varepsilon}{3}$, for all $t > T_2$. In accordance with Lemma 2.1 and the arbitrariness of ε , we easily achieve

$$\langle x_2(t) \rangle^* \leq \frac{\widehat{r}_2 + \langle \frac{f}{a} \rangle^*}{h^l} \triangleq M_y.$$

The desired result is obtained.

(e). If $x_2^* > 0$ a.s is false, let $\Omega = \{x_2^* = 0\}$, then $P(\Omega) > 0$. For an arbitrary $\nu \in \Omega$, we have $\lim_{t \to \infty} x_2(t, \nu) = 0$. From the second equation of (4.3) and by virtue of Eq (4.2), we show that $\left[\frac{\ln x_2(t,\nu)}{t}\right]^* = \widehat{r_2} > 0$ a.s. Then we follow that $P\{\left[\frac{\ln x_2(t,\nu)}{t}\right]^* > 0\} > 0$, which contradicts with Lemma 4.1. The result is then concluded.

Remark 1. By the proof of Theorem 4.1.2, we observe that if $\hat{r}_1 < 0$, then $k_*\hat{r}_2 + f^*\hat{r}_1 < 0$. Thus, if the prey species is extinct, then the predator species will also be extinct. This notion is consistent with the reality. Moreover, if $\hat{r}_1 > 0$ and $k_*\hat{r}_2 + f^*\hat{r}_1 < 0$, then even if the prey population is persistent, the predators end in extinction because of an excessively large diffusion coefficient σ_2^2 .

Remark 2. According to conclusion (5) of Theorem 4.1.2, with the effect of impulsive perturbations despite the regression of the prey population to extinction, the predator may remain weakly persistent.

4.2. Stochastic permanence

Theorem 4.2.1 If $(\max\{\sigma_1^u, \sigma_2^u\})^2 + 2(\frac{\omega^u}{b^l} + g^u) < 2\min\{r^l, \frac{f^l}{a^u}\}$ holds, then Eq (1.2) is stochastically permanent.

Proof. The whole proof is divided into two parts. First, we must prove that for arbitrary $\varepsilon > 0$, there is a constant $\delta > 0$ satisfying $P_*\{|X_1(t)| \ge \delta\} \ge 1 - \varepsilon$, where $X_1(t) = (x_1(t), x_2(t))$.

At this point, we show that for any initial value $X(0) = (y_1(0), y_2(0)) \in \mathbb{R}^2_+$, the solution $X(t) = (y_1(t), y_2(t))$ holds the property that

$$\limsup_{t\to\infty} \mathrm{E}(\frac{1}{|\overline{X}(t)|^{\theta}}) \leq M_0,$$

where $\theta > 0$ is a sufficiently small constant, such that

$$\min\{r^{l}, \frac{f^{l}}{a^{u}}\} > (\frac{\omega^{u}}{b^{l}} + g^{u}) + \frac{(\theta + 1)}{2}(\max\{\sigma_{1}^{u}, \sigma_{2}^{u}\})^{2}.$$
(4.16)

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By virtue of Eq (4.16), there is an arbitrary constant p > 0 such that

$$\min\{r^{l}, \frac{f^{l}}{a^{u}}\} - \frac{(\theta+1)}{2}(\max\{\sigma_{1}^{u}, \sigma_{2}^{u}\})^{2} - (\frac{\omega^{u}}{b^{l}} + g^{u}) - p > 0.$$
(4.17)

Define $V(y_1, y_2) = y_1 + y_2$, then

$$dV(y_1, y_2) = y_1 \Big(r(t) - k(t) \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1 + \rho_{2k}) y_2 \Big) dt + y_1 \sigma_1(t) dB_1(t) + y_2 \Big(-g(t) - h(t) \prod_{0 < \rho_{2k} < t} (1 + \rho_{1k}) y_2 + \frac{f(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 \Big) dt + y_2 \sigma_2(t) dB_2(t).$$

Let $U(y_1, y_2) = \frac{1}{V(y_1, y_2)}$, according to the *Itô*'s formula, we obtain

$$\begin{split} dU(\overline{X}) &= -U^2(\overline{X}) \Big[y_1 \Big(r(t) - k(t) \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1 + \rho_{2k}) y_2 \Big) dt \Big] \\ &- U^2(\overline{X}) y_2 \Big(-g(t) - h(t) \prod_{0 < \rho_{2k} < t} (1 + \rho_{1k}) y_2 + \frac{f(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 \Big) dt \\ &+ U^3(\overline{X}) [y_1^2 \sigma_1^2(t) + y_2^2 \sigma_2^2(t)] dt - U^2(\overline{X}) \sigma_1(t) y_1 dB_1(t) + \sigma_2(t) y_2 dB_2(t) \\ &= \mathrm{L}U(\overline{X}) dt - U^2(\overline{X}) [\sigma_1(t) y_1 dB_1(t) + \sigma_2(t) y_2 dB_2(t)]. \end{split}$$

Under the condition of this theorem, a positive constant θ can be chosen to satisfy Eq (4.16). By the Itô formula,

$$\mathcal{L}(1+U(\overline{X}))^{\theta} = \theta(1+U(\overline{X}))^{\theta-1}\mathcal{L}U(\overline{X}) + \frac{1}{2}\theta(\theta-1)(1+U(\overline{X}))^{\theta-2}U^{4}(\overline{X}) + \left(y_{1}^{2}\sigma_{1}^{2}(t) + y_{2}^{2}\sigma_{2}^{2}(t)\right).$$

Then we choose p > 0 to be sufficiently small such that the term satisfies Eq (4.17). We define $W(\overline{X}) = e^{pt}(1 + U(\overline{X}))^{\theta}$ and consequently achieve

$$\begin{split} \mathsf{L}W(\overline{X}) &= pe^{pt}(1+U(\overline{X}))^{\theta} + e^{pt}\mathsf{L}(1+U(\overline{X}))^{\theta} \\ &= e^{pt}(1+U(\overline{X}))^{\theta-2} \Big\{ p(1+U(\overline{X}))^2 - \theta U^2(\overline{X})y_1\Big(r(t) - k(t) \prod_{0 < \tau_k < t} (1+\rho_{1k})y_1 \\ &\quad - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1+\rho_{2k})y_2 \Big) - \theta U^2(\overline{X})y_2\Big(- g(t) - h(t) \prod_{0 < \tau_k < t} (1+\rho_{2k})y_2 \\ &\quad + \frac{f(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1+\rho_{1k})y_1 \Big) - \theta U^3(\overline{X})y_2(-g(t) - h(t) \prod_{0 < \tau_k < t} (1+\rho_{2k})y_2) \\ &\quad - \theta U^3(\overline{X}) \Big[y_1\Big(r(t) - k(t) \prod_{0 < \tau_k < t} (1+\rho_{1k})y_1 - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1+\rho_{2k})y_2 \Big) \\ &\quad + \frac{f(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1+\rho_{1k})y_1 \Big] + \theta U^3(\overline{X}) \Big[y_1^2 \sigma_1^2(t) + y_2^2 \sigma_2^2(t) \Big] + \frac{\theta(\theta+1)}{2} U^4(\overline{X}) \Big[y_1^2 \sigma_1^2(t) + y_2^2 \sigma_2^2(t) \Big] \Big\} \\ &\leq e^{pt}(1+U(\overline{X}))^{\theta-2} \Big\{ (p+\theta\max\{k^{\mu}M,h^{\mu}\widetilde{M}\}) + \Big[2p-\theta\min\{r^l,\frac{f^l}{a^{\mu}}\} + \theta\max\{k^{\mu}M,h^{\mu}\widetilde{M}\} \Big] U(\overline{X}) \\ &\quad + \Big[p+\theta(\frac{\omega^{\mu}}{b^l}+g^{\mu}) - \theta\min\{r^l,\frac{f^l}{a^{\mu}}\} + \frac{\theta(\theta+1)}{2} (\max\{\sigma_1^{\mu},\delta_1^{\mu}\})^2 \Big] U^2(\overline{X}) \Big\}. \end{split}$$

By Eq (4.17), a positive constant *S* satisfying $LW(\overline{X}) \leq Se^{pt}$ is easily noted. Consequently, $E[e^{pt}(1+U(\overline{X}))^{\theta}] \leq (1+U(0))^{\theta} + \frac{S(e^{pt}-1)}{p}$ and then

$$\limsup_{t\to\infty} \mathbb{E}[U^{\theta}(\overline{X}(t))] \le \limsup_{t\to\infty} \mathbb{E}[(1+U(\overline{X}(t))^{\theta}] \le \frac{S}{p}.$$

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Notably, $(y_1 + y_2)^{\theta} \le 2^{\theta} (y_1^2 + y_2^2)^{\frac{\theta}{2}} = 2^{\theta} |\overline{X}|^{\theta}$, where $\overline{X} = (y_1, y_2) \in \mathbb{R}^2_+$. Then, we obtain

$$\limsup_{t\to\infty} \mathbb{E}\left[\frac{1}{|\overline{X}(t)|^{\theta}}\right] \le 2^{-\theta} \limsup_{t\to\infty} \mathbb{E}U^{\theta}(\overline{X}) \le 2^{-\theta}\frac{S}{p} := M_0,$$

and

$$\limsup_{t\to\infty} \mathbb{E}\left[\frac{1}{|X_1(t)|^{\theta}}\right] \le \overline{m}^{-\theta} \limsup_{t\to\infty} \mathbb{E}U^{\theta}(\overline{X}) \le \overline{m}^{-\theta}M_0 := \overline{M_0},$$

where $\overline{m} = \min\{m, \widetilde{m}\}$. Therefore, for arbitrary $\varepsilon > 0$, we let $\delta = (\varepsilon \setminus \overline{M_0})^{\frac{1}{\theta}}$ in accordance with Chebyshev's inequality, thereby yielding

$$P\{|X_1(t)| < \delta\} = P\{|X_1(t)|^{-\theta} > \delta^{-\theta}\} \le E[|X_1(t)|^{-\theta}]/\delta^{-\theta} = \delta^{\theta} E[|X_1(t)|^{-\theta}],$$

thus, $P_*\{|X_1(t)| \ge \delta\} \ge 1 - \varepsilon$.

In the following relations, we prove that for any $\varepsilon > 0$, there exists $\chi > 0$ satisfying $P_*\{|X_1(t)| \le \chi\} \ge 1 - \varepsilon$. Define $V_4(\overline{X}) = y_1^q + y_2^q$, herein 0 < q < 1 and $\overline{X} = (y_1, y_2) \in \mathbb{R}^2_+$, then by virtue of *Itô*'s formula, we obtain the expression

$$\begin{aligned} dV_4(\overline{X}(t)) &= q y_1^q \Big[r(t) - k(t) \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 + \frac{q-1}{2} \sigma_1^2(t) - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1 + \rho_{2k}) y_2 \Big] dt \\ &+ q y_2^q \Big[-g(t) - h(t) \prod_{0 < \tau_k < t} (1 + \rho_{2k}) + \frac{q-1}{2} \sigma_2^2(t) y_2 + \frac{f(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 \Big] dt \\ &+ q y_1^q \sigma_1(t) dB_1(t) + q y_2^q \sigma_2(t) dB_2(t). \end{aligned}$$

Let n_0 be a sufficiently large constant, such that $y_1(0)$, $y_2(0)$ remain within the internal $[\frac{1}{n_0}, n_0]$. For each integer $n \ge n_0$, we define the stopping time $t_n = \inf\{t \ge 0 : y_1(t) \notin (1/n, n) \text{ or } y_2(t) \notin (1/n, n)\}$. Obviously, t_n increases as $n \to +\infty$. Using $It\hat{o}$'s formula again for $\exp\{t\}V_4(\overline{X})$ and accounting for the expectations on both sides, we show that

$$\begin{split} & \operatorname{E}[\exp\{t \wedge t_{n}\}\overline{X}^{q}(t \wedge t_{n})] - \overline{X}^{q}(0) \\ & \leq q \operatorname{E} \int_{0}^{t \wedge t_{n}} \exp\{s\} y_{1}^{q}(s) \Big[1 + q(r(s) - k(s) \prod_{0 < \tau_{k} < s} (1 + \rho_{1k}) y_{1}(s) - \frac{1 - q}{2} \sigma_{1}^{2}(s)) \Big] \mathrm{d}s \\ & + q \operatorname{E} \int_{0}^{t \wedge t_{n}} \exp\{s\} \varphi^{q}(s) \Big[1 + q(-g(s) - h(s) \prod_{0 < \tau_{k} < s} (1 + \rho_{2k}) y_{2}(s) + \frac{f(s)}{a(s)} - \frac{1 - q}{2} \sigma_{2}^{2}(s)) \Big] \mathrm{d}s \\ & \leq \operatorname{E} \int_{0}^{t \wedge t_{n}} (K_{1} + K_{2}) \exp\{s\} \mathrm{d}s \\ & \leq (K_{1} + K_{2}) (\exp\{t\} - 1), \end{split}$$

where K_1, K_2 are positive constants. Letting $n \to +\infty$ yields

$$\exp\{t\} \mathbb{E}[\overline{X}^{q}(t)] \le \overline{X}^{q}(0) + (K_{1} + K_{2})(\exp\{t\} - 1).$$

Then, we achieve $\limsup_{t \to +\infty} \mathbb{E}[\overline{X}^{q}(t)] \leq K_{1} + K_{2}$ and $\limsup_{t \to +\infty} \mathbb{E}[X_{1}^{q}(t)] \leq \overline{M}^{q}(K_{1} + K_{2})$, where $\overline{M} = \max\{M, \widetilde{M}\}$. Therefore, at any given $\varepsilon > 0$, we let $\chi = \frac{\overline{M}(K_{1}+K_{2})^{1/q}}{\varepsilon^{1/q}}$, by virtue of the Chebyshev inequality, we easily show that

$$\mathbf{P}\{|X_1(t)| > \chi\} = \mathbf{P}\{|X_1(t)|^q > \chi^q\} \le \mathbf{E}[|X_1(t)|^q]/\chi^q.$$

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Consequently, $P_*\{|X_1(t)| \le \chi\} \ge 1 - \varepsilon$. Theorem 4.2.1 is proven.

Remark 3. From the conditions of Theorem 4.2.1, we find that although the stochastic disturbance greatly influence the dynamical property of the system, the bounded impulsive perturbations do not affect the stochastic permanence of the model.

Remark 4. We should point out that the definition of stochastically permanent which requires that all species have positive upper bounds and at least one species has a positive lower bound, cannot demonstrate the permanence of all species. It has some limitations and deficiency. If there is only one species having a positive lower bound and all the other species go extinction, the system is still permanent. A new definition of stochastic permanence [31] may be more appropriate.

5. Global attractiveness

In this section, we provide some sufficient criteria to ensure the global attractiveness of the Equation (1.2).

Theorem 5.1. For any initial value $(y_1(0), y_2(0)) \in \mathbb{R}^+_2$, $(y_1(t), y_2(t))$ is a solution of Eq (3.1) on $[0, +\infty)$. Then almost every sample path of $(y_1(t), y_2(t))$ is uniformly continuous.

The proof of Theorem 5.1 is given in Appendix A. **Theorem 5.2.** Suppose that constants $\mu_i > 0$ (i = 1, 2) satisfying $\liminf A_i(t) > 0$ exist, where

$$A_{1}(t) = \mu_{1}[mk(t) - 2\frac{\omega(t)a(t)}{b(t)}] - 2\mu_{2}f(t),$$

$$A_{2}(t) = \mu_{2}[\widetilde{m}h(t) - 2\frac{f(t)b(t)}{a(t)}] - 2\mu_{1}\omega(t),$$
(5.1)

then Eq (1.2) is globally attractive.

Proof. Let $(y_1(t), y_2(t))$, $(\tilde{y}_1(t), \tilde{y}_2(t))$ be two arbitrary solutions of Eq (3.1) with initial value $(y_1(0), y_2(0)), (\tilde{y}_1(0), \tilde{y}_2(0)) \in \mathbb{R}^2_+$. Denote $\tilde{\lambda}(t) = 1 + a(t) \prod (1 + \rho_{1k})\tilde{y}_1 + b(t) \prod (1 + \rho_{2k})\tilde{y}_2 + b(t) \prod ($ $0 < \tau_k < t$ $a(t)b(t)\prod_{0<\tau_k< t}(1+\rho_{1k})\prod_{0<\tau_k< t}(1+\rho_{2k})\tilde{y}_1\tilde{y}_2$ and define a Lyapunov function as follows:

$$V(t) = \mu_1 |\ln y_1(t) - \ln \tilde{y}_1(t)| + \mu_2 |\ln y_2(t) - \ln \tilde{y}_2(t)|$$

Then,

$$\begin{split} D^{+}(V(t)) &= \mu_{1} \mathrm{sgn}(y_{1} - \tilde{y}_{1}) \Big([r(t) - k(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{1k})y_{1} - 0.5\sigma_{1}^{2}(t) - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_{k} < t} (1 + \rho_{2k})y_{2}] dt \\ &- [r(t) - k(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{1k})\tilde{y}_{1} - 0.5\sigma_{1}^{2}(t) - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_{k} < t} (1 + \rho_{2k})\tilde{y}_{2}] dt \\ &+ \mu_{2} \mathrm{sgn}(y_{2} - \tilde{y}_{2}) \Big([-g(t) - h(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{2k})y_{2} - 0.5\sigma_{2}^{2}(t) + \frac{f(t)}{\lambda(t)} \prod_{0 < \tau_{k} < t} (1 + \rho_{1k})y_{1}] dt \Big) \\ &- [-g(t) - h(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{2k})\tilde{y}_{2} - 0.5\sigma_{2}^{2}(t) + \frac{f(t)}{\lambda(t)} \prod_{0 < \tau_{k} < t} (1 + \rho_{1k})\tilde{y}_{1}] dt \Big) \\ &\leq \mu_{1} \mathrm{sgn}(y_{1} - \tilde{y}_{1}) \Big(-k(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{1k})(y_{1} - \tilde{y}_{1}) + \omega(t) [\frac{0 < \tau_{k} < t}{\lambda(t)} - \frac{0 < \tau_{k} < t}{\lambda(t)} - \frac{0 < \tau_{k} < t}{\lambda(t)}] \Big) dt \\ &+ \mu_{2} \mathrm{sgn}(y_{2} - \tilde{y}_{2}) \Big(-h(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{2k})(y_{2} - \tilde{y}_{2}) + f(t) [\frac{0 < \tau_{k} < t}{\lambda(t)} - \frac{0 < \tau_{k} < t}{\lambda(t)} - \frac{0 < \tau_{k} < t}{\lambda(t)}] \Big) dt \\ &\leq - \Big[\mu_{1}(mk(t) - 2\frac{\omega(t)a(t)}{b(t)}) - 2\mu_{2}f(t) \Big] |y_{1} - \tilde{y}_{1}| - \Big[\mu_{2}(\tilde{m}h(t) - 2\frac{f(t)b(t)}{a(t)}) - 2\mu_{1}\omega(t) \Big] |y_{2} - \tilde{y}_{2}|. \end{split}$$

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By the condition $\liminf_{t\to\infty} A_i(t) > 0$ (i = 1, 2), constants $\alpha > 0$ and $T_0 > 0$ satisfying $A_i(t) \ge \alpha$ (i = 1, 2) exist for all $t \ge T_0$. Moreover, we obtain the following relation

$$D^{+}(V(t)) \leq -\alpha(|y_{1} - \tilde{y}_{1}| + |y_{2} - \tilde{y}_{2}|),$$
(5.2)

for all $t \ge T_0$. By integrating Eq (5.2) from T_0 to t, we achieve

$$V(t) - V(T_0) \le -\alpha \int_{T_0}^t (|y_1(s) - \tilde{y}_1(s)| + |y_2(s) - \tilde{y}_2(s)|) \, ds.$$

Consequently,

$$V(t) + \alpha \int_{T_0}^t \left(|y_1(s) - \tilde{y}_1(s)| + |y_2(s) - \tilde{y}_2(s)| \right) ds \le V(T_0) < +\infty.$$
(5.3)

Then by $V(t) \ge 0$, we note that $|y_1(t) - \tilde{y}_1(t)| \in L^1[0, +\infty)$, $|y_2(t) - \tilde{y}_2(t)| \in L^1[0, +\infty)$. Thus,

$$\lim_{t \to +\infty} |y_1(t) - \tilde{y}_1(t)| = 0, \lim_{t \to +\infty} |y_2(t) - \tilde{y}_2(t)| = 0.$$

Next,

$$\begin{split} \lim_{t \to +\infty} |x_1(t) - \tilde{x}_1(t)| &= \lim_{t \to +\infty} \prod_{0 < \tau_k < t} (1 + \rho_{1k}) |y_1(t) - \tilde{y}_1(t)| \le M \lim_{t \to +\infty} |y_1(t) - \tilde{y}_1(t)| = 0, \quad a.s.\\ \lim_{t \to +\infty} |x_2(t) - \tilde{x}_2(t)| &= \lim_{t \to +\infty} \prod_{0 < \tau_k < t} (1 + \rho_{2k}) |y_2(t) - \tilde{y}_2(t)| \le \widetilde{M} \lim_{t \to +\infty} |y_2(t) - \tilde{y}_2(t)| = 0. \quad a.s. \end{split}$$

Therefore, the desired assertion is obtained by Theorem 5.1 and Lemma 2.2.

6. Numerical simulations

In this section, some numerical simulations and examples are given to illustrate and augment our theoretical findings of Eq (1.2) by means of the Milstein method mentioned in Higham [35]. Moreover, the effects of impulsive and stochastic perturbations on population dynamics are discussed.

Example 1.

$$\begin{cases} dx_1(t) = x_1 \left[r(t) - (0.7 + 0.01 \sin t) x_1 - \frac{(0.1 + 0.02 \sin t) x_2}{1 + a(t) x_1 + b(t) x_2 + a(t) b(t) x_1 x_2} \right] dt \\ + x_1 \sigma_1(t) dB_1(t), \\ dx_2(t) = x_2 \left[-(0.2 + 0.05 \sin t) - (0.2 + 0.01 \sin t) x_2 + \frac{(0.2 + 0.02 \sin t) x_1}{1 + a(t) x_1 + b(t) x_2 + a(t) b(t) x_1 x_2} \right] dt \quad (6.1) \\ + x_2 \sigma_2(t) dB_2(t), \qquad t \neq \tau_k, \\ x_1(\tau_k^+) = (1 + \rho_{1k}) x_1(\tau_k), \\ x_2(\tau_k^+) = (1 + \rho_{2k}) x_2(\tau_k), \qquad t = \tau_k, k = 1, 2, 3, \dots \end{cases}$$

We set $r(t) = 0.2 + 0.01 \sin t$, $a(t) = 0.1 + 0.04 \sin t$, $b(t) = 0.5 + 0.05 \sin t$, $\frac{\sigma_1^2(t)}{2} = \frac{\sigma_2^2(t)}{2} = 0.3 + 0.02 \sin t$, $\rho_{1k} = \rho_{2k} = e^{(-1)^{k+1}\frac{1}{k}} - 1$ and $\tau_k = k$, then it can be obtained that $\hat{r}_1 = -0.1 < 0$. By Theorems 4.1.1 and 4.1.2, both prey and predator populations (x_1 and x_2 , respectively) regress to extinction, which is also further confirmed by Figure 1.

We then choose $\frac{\sigma_1^{2(t)}}{2} = 0.3 + 0.02 \sin t$, $\frac{\sigma_2^{2(t)}}{2} = 0.4 + 0.02 \sin t$, $\rho_{1k} = \rho_{2k} = e^{(-1)^{k+1}\frac{1}{k}} - 1$, and $r(t) = 0.4 + 0.01 \sin t$. The other parameters are the same as that in example 1, then $\hat{r}_1 = 0.1 > 0$, $\hat{r}_2 = -0.6 < 0$, and $k_* \hat{r}_2 + f^* \hat{r}_1 = -0.116 < 0$. In Figure 2, although the prey population x_1 is weakly persistent in the mean, the predator population x_2 end in extinction because of the effects of the white noises, which are of great importance in maintaining the coexistence of populations.

Example 2.

$$\begin{cases} dx_1(t) = x_1 \left[(0.6 + 0.1 \sin t)) - (0.7 + 0.01 \sin t) x_1 - \frac{(0.1 + 0.02 \sin t) x_2}{1 + a(t) x_1 + b(t) x_2 + a(t) b(t) x_1 x_2} \right] dt \\ + x_1 \sigma_1(t) dB_1(t), \\ dx_2(t) = x_2 \left[-(0.2 + 0.05 \sin t) - (0.2 + 0.01 \sin t) x_2 + \frac{(0.62 + 0.02 \sin t) x_1}{1 + a(t) x_1 + b(t) x_2 + a(t) b(t) x_1 x_2} \right] dt \quad (6.2) \\ + x_2 \sigma_2(t) dB_2(t), \qquad t \neq \tau_k, \\ x_1(\tau_k^+) = (1 + \rho_{1k}) x_1(\tau_k), \\ x_2(\tau_k^+) = (1 + \rho_{2k}) x_2(\tau_k), \qquad t = \tau_k, k = 1, 2, 3, \dots \end{cases}$$

We let $\frac{\sigma_1^2(t)}{2} = 0.1 + 0.05 \sin t$, $\frac{\sigma_2^2(t)}{2} = 0.1 + 0.02 \sin t$, $a(t) = 0.1 + 0.04 \sin t$, $b(t) = 0.5 + 0.05 \sin t$ $\rho_{1k} = \rho_{2k} = e^{-\frac{1}{k^2}} - 1$, and $\tau_k = k$, then $\hat{r}_1 = 0.5 > 0$, $\hat{r}_2 = -0.3 < 0$. Both the prey and predator populations (x_1 and x_2 , respectively) are weakly persistent in the mean of Figure 3.

Example 3.

$$\begin{cases} dx_{1}(t) = x_{1} \left[(1.21 + 0.01 \sin t)) - (0.7 + 0.01 \sin t)x_{1} - \frac{(0.1 + 0.02 \sin t)x_{2}}{1 + a(t)x_{1} + b(t)x_{2} + a(t)b(t)x_{1}x_{2}} \right] dt \\ + x_{1}\sigma_{1}(t)dB_{1}(t), \\ dx_{2}(t) = x_{2} \left[-(0.2 + 0.05 \sin t) - (0.2 + 0.01 \sin t)x_{2} + \frac{(0.62 + 0.02 \sin t)x_{1}}{1 + a(t)x_{1} + b(t)x_{2} + a(t)b(t)x_{1}x_{2}} \right] dt \quad (6.3) \\ + x_{2}\sigma_{2}(t)dB_{2}(t), \qquad t \neq \tau_{k}, \\ x_{1}(\tau_{k}^{+}) = (1 + \rho_{1k})x_{1}(\tau_{k}), \\ x_{2}(\tau_{k}^{+}) = (1 + \rho_{2k})x_{2}(\tau_{k}), \qquad t = \tau_{k}, k = 1, 2, 3, \dots \end{cases}$$

In Figure 4, we choose $\sigma_1(t) = 0.2 + 0.02 \sin t$, $\sigma_2(t) = 0.3 + 0.02 \sin t$, $a(t) = 0.1 + 0.04 \sin t$, $b(t) = 0.5 + 0.05 \sin t$, and the impulsive perturbations are (a) $\rho_{1k} = \rho_{2k} = e^{\frac{(-1)^{k+1}}{k}} - 1$, (b) $\rho_{1k} = \rho_{2k} = e^{\frac{1}{k^2}} - 1$, (c) $\rho_{1k} = \rho_{2k} = 0$. The conditions of Theorem 4.2.1 are satisfied in all of those cases and the Eq (1.2) displays stochastic permanence in Figure 4. Moreover, in Figure 4(a)–4(c), the bounded impulsive perturbations do not affect the stochastic permanence of the model.



Figure 1. Solutions of Eq (1.2) with $\frac{\sigma_1^2(t)}{2} = \frac{\sigma_2^2(t)}{2} = 0.3 + 0.02 \sin t$, $\rho_{1k} = \rho_{2k} = e^{(-1)^{k+1} \frac{1}{k}} - 1$, $\tau_k = k$. Both prey population x_1 and predator population x_2 go to extinction.



Figure 2. $\frac{\sigma_1^{2}(t)}{2} = 0.3 + 0.02 \sin t$, $\frac{\sigma_2^{2}(t)}{2} = 0.4 + 0.02 \sin t$, $\rho_{1k} = \rho_{2k} = e^{(-1)^{k+1}\frac{1}{k}} - 1$, $\tau_k = k$. The prey population is weakly persistent in the mean and predator population is extinct.



Figure 3. Both populations are weakly persistent in the mean for Eq (1.2).

Example 4.

$$\begin{cases} dx_{1}(t) = x_{1} \left[(1.8 + 0.01 \sin t)) - (0.7 + 0.01 \sin t)x_{1} - \frac{(0.1 + 0.02 \sin t)x_{2}}{1 + a(t)x_{1} + b(t)x_{2} + a(t)b(t)x_{1}x_{2}} \right] dt \\ + x_{1}\sigma_{1}(t)dB_{1}(t), \\ dx_{2}(t) = x_{2} \left[-(0.2 + 0.05 \sin t) - (0.8 + 0.01 \sin t)x_{2} + \frac{(1.12 + 0.02 \sin t)x_{1}}{1 + a(t)x_{1} + b(t)x_{2} + a(t)b(t)x_{1}x_{2}} \right] dt \quad (6.4) \\ + x_{2}\sigma_{2}(t)dB_{2}(t), \qquad t \neq \tau_{k}, \\ x_{1}(\tau_{k}^{+}) = (1 + \rho_{1k})x_{1}(\tau_{k}), \\ x_{2}(\tau_{k}^{+}) = (1 + \rho_{2k})x_{2}(\tau_{k}), \qquad t = \tau_{k}, k = 1, 2, 3, \dots \end{cases}$$

Set $\frac{\sigma_1^2(t)}{2} = 0.1 + 0.05 \sin t$, $\frac{\sigma_2^2(t)}{2} = 0.6 + 0.02 \sin t$, $x_1(0) = 2$, $x_2(0) = 3$, $\tilde{x}_1(0) = 0.5$, and $\tilde{x}_2(0) = 0.3$. We then observe from Figure 5 that Eq (1.2) is globally attractive.

In the following instance, the effects of negative and positive impulses on the species are investigated. We let $\rho_{1k} = e^{-1.92} - 1$, $\rho_{2k} = e^{-0.02} - 1$, and $\tau_k = k$. The other parameters are the same as those in Example 3, then we obtain that $\hat{r}_1 = -0.03 < 0$ and $\hat{r}_2 = -0.36 < 0$. On the basis of Theorems 4.1.1 and 4.1.2, both the prey and predator populations end in extinction, which is further confirmed by Figure 6(b). By comparing Figure 6(a) with Figure 6(b), we observe that the negative impulses do not benefit species coexistence. Moreover, considering the impulsive perturbations are (a) $\rho_{1k} = \rho_{2k} = 0$, (b) $\rho_{1k} = e^{\frac{1}{k^2}} - 1$, $\rho_{2k} = e^{0.8} - 1$, (c) $\rho_{1k} = \rho_{2k} = e^{0.8} - 1$ and the other parameters are the same as those in Example 1. Hence, the system can be altered from extinction to persistence with the effects of positive impulsive perturbations (Figure 7(a)-7(c)). Herein, persistence can be divided into two cases as follows: first, the predator population $x_2(t)$ is weakly persistent and the prey population $x_1(t)$ pro-



Figure 4. $\sigma_1(t) = 0.2 + 0.02 \sin t$, $\sigma_2(t) = 0.3 + 0.02 \sin t$, the only difference between these graphs is the impulses: (a) $\rho_{1k} = \rho_{2k} = e^{\frac{(-1)^{k+1}}{k}} - 1$, (b) $\rho_{1k} = \rho_{2k} = e^{\frac{1}{k^2}} - 1$, (c) $\rho_{1k} = \rho_{2k} = 0$. The system is stochastic permanence.

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Figure 5. The figures show the attractivity of Eq (1.2).



Figure 6. $\sigma_1(t) = 0.2 + 0.02 \sin t$, $\sigma_2(t) = 0.3 + 0.02 \sin t$, and the different impulses chosen for the graphs are : (a) $\rho_{1k} = \rho_{2k} = 0$, (b) $\rho_{1k} = e^{-1.92} - 1$, $\rho_{2k} = e^{-0.02} - 1$. The negative impulses do not benefit species coexistence.



Figure 7. $\frac{\sigma_1^2(t)}{2} = \frac{\sigma_2^2(t)}{2} = 0.3 + 0.02 \sin t$, and the impulsive perturbations are: (a) $\rho_{1k} = \rho_{2k} = 0$, (b) $\rho_{1k} = e^{\frac{1}{k^2}} - 1$, $\rho_{2k} = e^{0.8} - 1$, (c) $\rho_{1k} = \rho_{2k} = e^{0.8} - 1$. Positive impulses are advantageous for the coexistence of ecosystems.

ceeds to extinction and second, both of the populations are persistent. Therefore, positive impulses are advantageous for the coexistence of ecosystems. Moreover, comparing figures 3 and 6, figures 1 and 7, we can derive that if the impulsive perturbations are unbounded, some properties may be changed significantly.

7. Conclusions

In this paper, we propose a stochastic non-autonomous predator-prey system with impulsive perturbations and investigate the qualitative dynamic properties of the model. Under some sufficient conditions, we present the extinction and a series of persistence in the mean of the system, including non-persistence, weak persistence and strong persistence in the mean. Furthermore, we obtain the global attractivity of the model. From the assumptions of Theorems 4.1.1, 4.1.2 and 4.2.1, we demonstrate that the stochastic and impulsive disturbances greatly influence the extinction and persistence of the system. Positive impulses are advantageous for the coexistence of ecosystem, whereas negative impulses are not beneficial for species coexistence. Moreover, the results show that the bounded impulsive perturbations do not affect all the properties, such as the stochastic permanence of the model. However, if the impulsive perturbations are unbounded, some properties may be changed significantly.

Some interesting topics require further investigations. If we also consider the effects of time delays and telephone noise [36-38] on Eq (1.2) to propose a more realistic model, then how will the properties change? We leave it for future investigation.

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Conflict of interest

The authors declare that they have no competing of interests regarding the publication of this paper.

References

- 1. H. I. Freedman, Deterministic Mathematical Models in Population Ecology, Marcel Dekker, New York, 1980.
- 2. Z. Yao, S. Xie, N. Yu, Dynamics of cooperative predator-prey system with impulsive effects and Beddington-DeAngelis functional response, *J. Egypt. Math. Soc.*, **21** (2013), 213–223.
- 3. Z. Shen, J. Wei, Hopf bifurcation analysis in a diffusive predator-prey system with delay and surplus killing effect, *Math. Biosci. Eng.*, **15** (2018), 693–715.
- 4. M. Bandyopadhyay, J. Chattopadhyay, Ratio-dependent predator-prey model: Effect of environmental fluctuation and stability, *Nonlinearity*, **18** (2005), 913–936.
- 5. M. Liu, K. Wang, Q. Wu, Survival analysis of stochastic competitive models in a polluted environment and stochastic competitive exclusion principle, *B. Math. Biol.*, **73** (2011), 1969–2012.

- 6. M. Liu, C. Bai, K. Wang, Asymptotic stability of a two-group stochastic SEIR model with infinite delays, *Commun. Nonlinear. Sci. Numer. Simulat.*, **19** (2014), 3444–3453.
- X. Mao, G. Marion, E. Renshaw, Environmental Brownian noise suppresses explosions in populations dynamics, *Stoch. Proc. Appl.*, 97 (2002), 95–110.
- 8. D. Jana, R. Agrawal, R. K. Upadhyay, Dynamics of generalist predator in a stochastic environment: Effect of delayed growth and prey refuge, *Appl. Math. Comput.*, **268** (2015), 1072–1094.
- 9. Y. Lin, D. Jiang, S. Wang, Stationary distribution of a stochastic SIS epidemic model with vaccination, *Physica A*, **394** (2014), 718–727.
- M. Liu, K. Wang, Persistence and extinction in stochastic non-autonomous logistic systems, J. Math. Anal. Appl., 375 (2011), 443–457.
- 11. M. Liu, K. Wang, Persistence, extinction and global asymptotical stability of a non-autonomous predator-prey model with random perturbation, *Appl. Math. Model.*, **36** (2012), 5344–5353.
- 12. Y. Zhao, S. Yuan, J. Ma, Survival and stationary distribution analysis of a stochastic competitive model of three species in a polluted environment, *B. Math. Biol.*, **77** (2015), 1285–1326.
- 13. L. Zu, D. Jiang, D. O'Regan, B. Ge, Periodic solution for a non-autonomous Lotka-Volterra predator-prey model with random perturbation, *J. Math. Anal. Appl.*, **430** (2015), 428–437.
- 14. X. Liu, S. Zhong, B. Tian, F. Zheng, Asymptotic properties of a stochastic predator-prey model with Crowley-Martin functional response, *J. Appl. Math. Comput.*, **43** (2013), 479–490.
- 15. M. Hassell, G. Varley, New inductive population model for insect parasites and its bearing on biological control, *Nature*, **223** (1969), 1133–1137.
- 16. X. Yan, C. Zhang, Stability and turing instability in a diffusive predator-prey system with Beddington-DeAngelis functional response, *Nonlinear Anal-Real.*, **20** (2014), 1–13.
- 17. C. Xu, P. Li, 20 Oscillations for a delayed predator-prey model with Hassell-Varley-type functional response, *C. R. Biol.*, **338** (2015), 227–240.
- 18. Y. Zhang, S. Gao, Y. Liu, Analysis of a nonautonomous model for migratory birds with saturation incidence rate, *Commun. Nonlinear Sci. Numer. Simulat.*, **17** (2012), 1659–1672.
- 19. R. May, Stability and Complexity in Model Ecosystems, Princeton: Princeton University Press, 1974.
- 20. X. Shi, X. Zhou, X. Song, Analysis of a stage-structured predator-prey model with Crowley-Martin function, *J. Appl. Math. Comput.*, **36** (2011), 459–472.
- 21. T. Zhang, J. Zhang, X. Meng, Geometric analysis of a pest management model with Holling's type III functional response and nonlinear state feedback control, *Nonlinear Dyn.*, **84** (2016), 1529–1539.
- 22. S. Chen, J. Wei, J. Yu, Stationary patterns of a diffusive predator-prey model with Crowley-Martin functional response, *Nonlinear Anal. Real*, **39** (2018), 33–57.
- J. Tripathi, S. Tyagi, S. Abbas, Global analysis of a delayed density dependent predator Cprey model with Crowley-Martin functional response, *Commun. Nonlinear. Sci. Numer. Simulat.*, 30 (2016), 45–69.

- 24. R. Tan, Z. Liu, S. Guo, H. Xiang, On a nonautonomous competitive system subject to stochastic and impulsive perturbations, *Appl. Math. Comput.*, **256** (2015), 702–714.
- 25. S. Zhang, D. Tan, Dynamics of a stochastic predator-prey system in a polluted environment with pulse toxicant input and impulsive perturbations, *Appl. Math. Model.*, **39** (2015), 6319–6331.
- R. Wu, X. Zou, K. Wang, Asymptotic behavior of a stochastic non-autonomous predator-prey model with impulsive perturbations, *Commun. Nonlinear Sci. Numer. Simulat.*, **20** (2015), 965– 974.
- 27. Y. Zhang, K. Fan, S. Gao, S. Chen, A remark on stationary distribution of a stochastic SIR epidemic model with double saturated rates, *Appl. Math. lett.*, **76** (2018), 46–52.
- 28. X. Li, X. Mao, Population dynamical behavior of non-autonomous Lotka-Volterra competitive system with random perturbation, *Discret Contin. Dyn. Syst.*, **24** (2009), 523–593.
- 29. M. Liu and K. Wang, On a stochastic logistic equation with impulsive perturbations, *Comput. Math. Appl.*, **63** (2012), 871–886.
- 30. X. Mao, Stochastic versions of the Lassalle Theorem, T. Differ. Equ., 153 (1999), 175–195.
- 31. R. Tan, Z. Liu, R.A. Cheke, Periodicity and stability in a single-species model governed by impulsive differential equation, *Appl. Math. Comput.*, **36** (2012), 1085–1094.
- 32. M. Liu, M. Fan, Permanence of stochastic Lotka-Volterra systems, J. Nonlinear Sci., 27 (2017), 425–452.
- 33. K. Wang, Stochastic Models in Mathematical Biology, Beijing: Science Press, 2010.
- 34. S. Cheng, Stochastic population systems, Stoch. Proc. Appl., 27 (2009), 854-874.
- 35. D. Higham, An algorithmic introduction to numerical simulation of stochastic differential equations, *SIAM Rev.*, **43** (2001), 525–546.
- 36. M. Liu, C. Bai, Optimal harvesting of a stochastic mutualism model with regime-switching, *Appl. Math. Comput.*, **373** (2020).
- 37. W. Ji, Z. Wang, G. Hu, Stationary distribution of a stochastic hybrid phytoplankton model with allelopathy, *Adv. Differ. Equ. NY*, 2020.
- 38. Z. Wang, M. Deng, M. Liu, Stationary distribution of a stochastic ratio-dependent predator-prey system with regime-switching, *Chaos Soliton Fract.*, 2020, 110462.

Appendix A

Proof of Theorem 3.1. From Lemma 3.1, it suffices to show that Eq (3.1) has a unique solution, $(y_1(t), y_2(t))$, for all $t \ge 0$ and will remain in \mathbb{R}^2_+ with probability one. The proof of this theorem is standard.

Let $n_0 > 0$ be sufficiently large such that $y_1(0)$, $y_2(0)$ lie within the interval $[1/n_0, n_0]$. For each integer $n > n_0$, define the stopping times $t_n = \inf\{t \in [0, t_e] : y_1(t) \notin (1/n, n) \text{ or } y_2(t) \notin (1/n, n)\}$. Obviously, t_n is increasing as $n \to +\infty$. Denote $t_{+\infty} = \lim_{n \to +\infty} t_n$, thus $t_{+\infty} \le t_e$. To complete the proof, it only needs to show $t_{+\infty} = +\infty$ a.s. If the statement is not true, there exist two constants T > 0 and $\varepsilon \in (0, 1)$, such that $P\{t_{+\infty} < +\infty\} > \varepsilon$. Therefore, there is an integer $n_1 \ge n_0$ satisfying $P\{t_n \le T\} \ge \varepsilon$, for all $n > n_1$.

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Define a \mathbb{C}^2 -function $V : \mathbb{R}^2_+ \to \mathbb{R}_+$ by $V(y_1, y_2) = (y_1 - 1 - \ln y_1) + (y_2 - 1 - \ln y_2)$, then we obtain that $V(y_1, y_2)$ is nonnegative. By virtue of *Itô*'s formula, we achieve that

$$\begin{aligned} dV(y_1, y_2) &= V_{y_1} dy_1 + 0.5 V_{y_1 y_1} (dy_1)^2 + V_{y_2} dy_2 + 0.5 V_{y_2 y_2} (dy_2)^2 \\ &= \left[(1 - 1/y_1) y_1 \Big(r(t) - k(t) \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1 + \rho_{2k}) y_2 \Big) \right. \\ &+ (1 - 1/y_2) y_2 \Big(-g(t) - h(t) \prod_{0 < \tau_k < t} (1 + \rho_{2k}) y_2 + \frac{f(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 \Big) \Big] dt \\ &+ 0.5 (\sigma_1^2(t) + \sigma_2^2(t)) dt + (1 - 1/y_1) y_1 \sigma_1(t) dB_1(t) + (1 - 1/y_2) y_2 \sigma_2(t) dB_2(t) \\ &\leq \left[(r^u + k^u M) y_1 - k^l m y_1^2 - r^l + \frac{\omega^u}{b^l} + g^u + (\widetilde{M}h^u + \frac{f^u}{a^l} - g^l) y_2 - \widetilde{m}h^l y_2^2 \Big] dt \\ &+ 0.5 [(\sigma_1^u)^2 + (\sigma_2^u)^2] dt + (y_1 - 1) \sigma_1(t) dB_1(t) + (y_2 - 1) \sigma_2(t) dB_2(t). \end{aligned}$$

According to the negative coefficients of the quadratic terms, there is a positive number G satisfying

$$dV(y_1, y_2) \le Gdt + (y_1 - 1)\sigma_1(t)dB_1(t) + (y_2 - 1)\sigma_2(t)dB_2(t).$$

Thus

$$\int_0^{t_n \wedge T} \mathrm{d}V(y_1, y_2) \le \int_0^{t_n \wedge T} G \mathrm{d}t + \int_0^{t_n \wedge T} \left[(y_1(t) - 1)\sigma_1(t) \mathrm{d}B_1(t) + (y_2(t) - 1)\sigma_2(t) \mathrm{d}B_2(t) \right],$$

where $t_n \wedge T = \min\{t_n, T\}$. Taking expectation yields that

$$EV(y_1(t_n \wedge T), y_2(t_n \wedge T)) \le V(y_1(0), y_2(0)) + GE(t_n \wedge T) \le V(y_1(0), y_2(0)) + GT.$$

Let $\Omega_n = \{t_n \leq T\}$, then $P(\Omega_n) \geq \varepsilon$. For any $w \in \Omega_n$, $y_1(t_n, w)$ or $y_2(t_n, w)$ equals either *n* or 1/n, thus

$$V(y_1(t_n, w), y_2(t_n, w)) \ge \min\{n - 1 - \ln n, 1/n - 1 + \ln n\}.$$

Therefore it can be shown that

$$V(y_1(0), y_2(0)) + G_1T \ge \mathbb{E}[1_{\Omega_n}(w)V(y_1(t_n, w), y_2(t_n, w))]$$

$$\ge \varepsilon \min\{n - 1 - \ln n, 1/n - 1 + \ln n\},\$$

where 1_{Ω_n} is the indicator function of Ω_n . Letting $n \to +\infty$ leads to the contradiction. So we obtain that $t_{+\infty} = +\infty$ a.s. This completes the proof.

Proof of Theorem 3.2. Define $V_1(y_1) = y_1^p$ and $V_2(y_2) = y_2^p$, respectively, for $(y_1, y_2) \in \mathbb{R}^2_+$ and p > 1. According to the *Itô*'s formula, we have

$$\begin{aligned} \mathsf{d}(y_1^p) &= p y_1^{p-1} \mathsf{d} y_1 + 0.5 p(p-1) y_1^{p-2} (\mathsf{d} y_1)^2 \\ &= p y_1^p \Big(r(t) - k(t) \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1 + \rho_{2k}) y_2 + 0.5(p-1) \sigma_1^2(t) \Big) \mathsf{d} t \\ &+ p y_1^p \sigma_1(t) \mathsf{d} B_1(t) \\ &\leq p y_1^p [r^u - mk^l y_1 + 0.5(p-1)(\sigma_1^u)^2] \mathsf{d} t + p y_1^p \sigma_1(t) \mathsf{d} B_1(t). \end{aligned}$$

and

$$\begin{aligned} \mathsf{d}(y_2^p) &= p y_2^p \Big(-g(t) - h(t) \prod_{0 < \tau_k < t} (1 + \rho_{2k}) y_2 + \frac{f(t)}{\lambda(t)} \prod_{0 < \tau_k < t} (1 + \rho_{1k}) y_1 + 0.5(p-1)\sigma_2^2(t) \Big) \mathsf{d}t \\ &+ p y_2^p \sigma_2(t) \mathsf{d}B_2(t) \\ &\leq p y_2^p [\frac{f^u}{a^l} + 0.5p(\sigma_2^u)^2 - \widetilde{m}h^l y_2] \mathsf{d}t + p y_2^p \sigma_2(t) \mathsf{d}B_2(t). \end{aligned}$$

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Taking expectation and then

$$\begin{aligned} \frac{dE[y_1^p(t)]}{dt} &\leq p\{[r^u + 0.5(p-1)(\sigma_1^u)^2]E[y_1^p(t)] - mk^l E[y_1^{p+1}(t)]\} \\ &\leq p\{[r^u + 0.5(p-1)(\sigma_1^u)^2]E[y_1^p(t)] - mk^l [E(y_1^p(t))]^{1+\frac{1}{p}}\} \\ &\leq pE[y_1^p(t)]\{[r^u + 0.5p(\sigma_1^u)^2] - mk^l [E(y_1^p(t))]^{\frac{1}{p}}\} \end{aligned}$$
(A.1)

and

$$\frac{\mathrm{dE}[y_2^p(t)]}{\mathrm{d}t} \le p \mathbb{E}[y_2^p(t)] \{ [\frac{f^u}{a^l} + 0.5p(\sigma_2^u)^2] - \widetilde{m}h^l [\mathbb{E}(y_2^p(t))]^{\frac{1}{p}} \}.$$
(A.2)

For Eq (A.1), considering the following equation

$$\frac{\mathrm{d}v(t)}{\mathrm{d}t} = pv(t)\{[r^u + 0.5p(\sigma_1^u)^2] - mk^l v^{\frac{1}{p}}(t)\}$$

with initial value $v(0) = v_0$. Obviously, we can obtain that

$$v(t) = \left(\frac{v_0^{1/p}(r^u + 0.5p(\sigma_1^u)^2)}{[r^u + 0.5p(\sigma_1^u)^2]e^{-(r^u + 0.5p(\sigma_1^u)^2)t} + mk^l v_0^{1/p}(1 - e^{-(r^u + 0.5p(\sigma_1^u)^2)t})}\right)^p.$$

Let $t \to \infty$ and thus $\lim_{t \to +\infty} v(t) = \left(\frac{r^{u}+0.5p(\sigma_1^{u})^2}{mk^l}\right)^p$. Using the comparison theorem yields that $\limsup_{t \to +\infty} E[y_1^p(t)] \leq G_1 < +\infty$, where $G_1 = \left(\frac{r^{u}+0.5p(\sigma_1^{u})^2}{mk^l}\right)^p$. In the same way, we can achieve $\limsup_{t \to +\infty} E[y_2^p(t)] \leq G_2 < +\infty$ and $G_2 = \left(\frac{f^{u}/a^l+0.5p(\sigma_2^{u})^2}{\tilde{m}h^l}\right)^p$.

Consequently, for a given constant $\varepsilon > 0$, there is a T > 0 satisfying $\mathbb{E}[y_1^p(t)] \le G_1 + \varepsilon$, $\mathbb{E}[y_2^p(t)] \le G_2 + \varepsilon$, for all t > T. Considering the continuity of $\mathbb{E}[y_1^p(t)]$ and $\mathbb{E}[y_2^p(t)]$, there exist $\overline{G_1}(p), \overline{G_2}(p) > 0$ such that $\mathbb{E}[y_1^p(t)] \le \overline{G_1}(p)$ and $\mathbb{E}[y_2^p(t)] \le \overline{G_2}(p)$ for $t \le T$. Denote $M_1(p) = \max\{\overline{G_1}(p), G_1 + \varepsilon\}$, $M_2(p) = \max\{\overline{G_2}(p), G_2 + \varepsilon\}$, then for all $t \in \mathbb{R}_+$,

$$E[y_1^p(t)] \le M_1(p), \quad E[y_2^p(t)] \le M_2(p)$$

and

$$\mathbf{E}[x_1^p(t)] = \mathbf{E}\Big[\big(\prod_{0 < \tau_k < t} (1 + \rho_{1k})y_1(t)\big)^p\Big] \le M^p M_1(p), \quad \mathbf{E}[x_2^p(t)] = \mathbf{E}\Big[\big(\prod_{0 < \tau_k < t} (1 + \rho_{2k})y_2(t)\big)^p\Big] \le \widetilde{M}^p M_1(p).$$

Thus, for $X_1(t) = (x_1(t), x_2(t)) \in \mathbb{R}^2_+$, it is obvious that $|X_1(t)|^p \le 2^{\frac{p}{2}} [x_1^p(t) + x_2^p(t)]$. Therefore,

$$E|X_1(t)|^p \le M_p < +\infty,$$

herein, $M_p = 2^{\frac{p}{2}} (M^p M_1(p) + \widetilde{M}^p M_2(p))$. By virtue of the Chebyshev inequality, the proof is completed.

Proof of Theorem 4.2. From Eq (3.1), we have

$$dy_{1} \leq y_{1} \Big(r(t) - k(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{1k}) y_{1} \Big) dt + y_{1} \sigma_{1}(t) dB_{1}(t),$$

$$dy_{2} \leq y_{2} \left(-h(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{2k}) y_{2} + \frac{f(t)}{a(t)} \right) dt + y_{2} \sigma_{2}(t) dB_{2}(t).$$
(A.3)

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Construct the comparison equation

$$d\bar{y}_{1} = \bar{y}_{1} \Big(r(t) - k(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{1k}) \bar{y}_{1} \Big) dt + \bar{y}_{1} \sigma_{1}(t) dB_{1}(t),$$

$$d\bar{y}_{2} = \bar{y}_{2} \left(-h(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{2k}) \bar{y}_{2} + \frac{f(t)}{a(t)} \right) dt + \bar{y}_{2} \sigma_{2}(t) dB_{2}(t),$$
(A.4)

where $(\bar{y}_1(t), \bar{y}_2(t))$ is a solution of Eq (A.4) with initial value $(y_1(0), y_2(0)) \in \mathbb{R}_2^+$. According to the comparison theorem for stochastic differential equations ([33]) and Theorem 4.1 ([34]), we can obtain that $\limsup_{t\to\infty} \frac{\ln y_1(t)}{\ln t} \leq \limsup_{t\to\infty} \frac{\ln y_2(t)}{\ln t} \leq \limsup_{t\to\infty} \frac{\ln y_2(t)}{\ln t} \leq 1$, a.s. Then,

$$\limsup_{t \to \infty} \frac{\ln y_1(t)}{t} \le \limsup_{t \to \infty} \frac{\ln y_1(t)}{\ln t} \cdot \limsup_{t \to \infty} \frac{\ln t}{t} \le \limsup_{t \to \infty} \frac{\ln t}{t} = 0.$$

Similarly, we have that $\limsup_{t\to\infty} \frac{\ln y_2(t)}{t} \le 0$. Therefore,

$$\limsup_{t \to \infty} \frac{\ln x_1(t)}{t} = \limsup_{t \to \infty} \frac{\sum_{0 < \tau_k < t} \ln(1 + \rho_{1k}) + \ln y_1(t)}{t} = \limsup_{t \to \infty} \frac{\ln y_1(t)}{t} + \limsup_{t \to \infty} \frac{\sum_{0 < \tau_k < t} \ln(1 + \rho_{1k})}{\ln t} \cdot \frac{\ln t}{t} \le 0,$$

$$\lim_{t \to \infty} \frac{\ln x_2(t)}{t} = \limsup_{t \to \infty} \frac{\sum_{0 < \tau_k < t} \ln(1 + \rho_{2k}) + \ln y_2(t)}{t} \le 0.$$
(A.5)

The proof is completed.

Proof of Theorem 5.1. The first equation of Eq (3.1) is equivalent to the following stochastic integral equation

$$y_{1}(t) = y_{1}(0) + \int_{0}^{t} y_{1}(s) \Big(r(s) - k(s) \prod_{0 < \tau_{k} < s} (1 + \rho_{1k}) y_{1}(s) - \frac{\omega(s)}{\lambda(s)} \prod_{0 < \tau_{k} < s} (1 + \rho_{2k}) y_{2}(s) \Big) ds + \int_{0}^{t} y_{1}(s) \sigma_{1}(s) dB_{1}(s),$$

Denote $f_1(s) = y_1(s)(r(s) - k(s) \prod_{0 < \tau_k < s} (1 + \rho_{1k})y_1(s) - \frac{\omega(s)}{\lambda(s)} \prod_{0 < \tau_k < s} (1 + \rho_{2k})y_2(s)), f_2(s) = y_1(s)\sigma_1(s)$, then

$$\begin{split} \mathsf{E}|f_{1}(t)|^{p} &= \mathsf{E}\Big|y_{1}(r(t) - k(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{1k})y_{1} - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_{k} < t} (1 + \rho_{2k})y_{2})\Big|^{p} \\ &= \mathsf{E}\Big[|y_{1}|^{p} \cdot |(r(t) - k(t) \prod_{0 < \tau_{k} < t} (1 + \rho_{1k})y_{1} - \frac{\omega(t)}{\lambda(t)} \prod_{0 < \tau_{k} < t} (1 + \rho_{2k})y_{2})|^{p}\Big] \\ &\leq \frac{1}{2}\mathsf{E}|y_{1}|^{2p} + \frac{1}{2}\mathsf{E}|r^{u} + Mk^{u}y_{1} + \widetilde{M}\omega^{u}y_{2}|^{2p} \\ &\leq \frac{1}{2}\mathsf{E}|y_{1}|^{2p} + \frac{1}{2}3^{2p-1}[(r^{u})^{2p} + (Mk^{u})^{2p}\mathsf{E}|y_{1}|^{2p} + (\widetilde{M}\omega^{u})^{2p}\mathsf{E}|y_{2}|^{2p}] \\ &\leq \frac{1}{2}M_{1}(2p) + \frac{3^{2p-1}}{2}[(r^{u})^{2p} + (Mk^{u})^{2p}M_{1}(2p) + (\widetilde{M}\omega^{u})^{2p}M_{2}(2p)] \\ &\triangleq F_{1}(p), \end{split}$$

$$(A.6)$$

$$\mathbf{E}|f_{2}(t)|^{p} = \mathbf{E}|y_{1}\sigma_{1}(t)|^{p} \le \sigma_{1}^{u}\mathbf{E}|y_{1}|^{p} \le \sigma_{1}^{u}M_{1}(p) \triangleq F_{2}(p).$$
(A.7)

By virtue of the moment inequality for stochastic integrals, we can show that for $0 \le t_1 \le t_2$ and p > 2,

$$\mathsf{E}|\int_{t_1}^{t_2} f_2(s) \mathrm{d}B_1(s)|^p \le (\frac{p(p-1)}{2})^{\frac{p}{2}} (t_2 - t_1)^{\frac{p-2}{2}} \int_{t_1}^{t_2} \mathsf{E}|f_2(s)|^p \mathrm{d}s \le (\frac{p(p-1)}{2})^{\frac{p}{2}} (t_2 - t_1)^{\frac{p}{2}} F_2(p).$$

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Thus, for $0 < t_1 < t_2 < \infty$, $t_2 - t_1 \le 1$, $\frac{1}{p} + \frac{1}{q} = 1$ and by Eqs (A.6) and (A.7), we achieve

$$\begin{split} \mathrm{E}[y_{1}(t_{2}) - y_{1}(t_{1})]^{p} &= \mathrm{E}[\int_{t_{1}}^{t_{2}} f_{1}(s)\mathrm{d}s + \int_{t_{1}}^{t_{2}} f_{2}(s)\mathrm{d}B_{1}(s)]^{p} \\ &\leq 2^{p-1}\mathrm{E}[\int_{t_{1}}^{t_{2}} f_{1}(s)\mathrm{d}s]^{p} + 2^{p-1}\mathrm{E}[\int_{t_{1}}^{t_{2}} f_{2}(s)\mathrm{d}B_{1}(s)]^{p} \\ &\leq 2^{p-1}(t_{2} - t_{1})^{\frac{p}{q}}\mathrm{E}[\int_{t_{1}}^{t_{2}} |f_{1}(s)|^{p}\mathrm{d}s] + 2^{p-1}(\frac{p(p-1)}{2})^{\frac{p}{2}}(t_{2} - t_{1})^{\frac{p}{2}}F_{2}(p) \\ &\leq 2^{p-1}(t_{2} - t_{1})^{\frac{p}{q}}F_{1}(p)(t_{2} - t_{1}) + 2^{p-1}(\frac{p(p-1)}{2})^{\frac{p}{2}}(t_{2} - t_{1})^{\frac{p}{2}}F_{2}(p) \\ &= 2^{p-1}(t_{2} - t_{1})^{p}F_{1}(p) + 2^{p-1}(\frac{p(p-1)}{2})^{\frac{p}{2}}(t_{2} - t_{1})^{\frac{p}{2}}F_{2}(p) \\ &= 2^{p-1}(t_{2} - t_{1})^{\frac{p}{2}}\{(t_{2} - t_{1})^{\frac{p}{2}}F_{1}(p) + (\frac{p(p-1)}{2})^{\frac{p}{2}}F_{2}(p)\} \\ &\leq 2^{p-1}(t_{2} - t_{1})^{\frac{p}{2}}\{1 + (\frac{p(p-1)}{2})^{\frac{p}{2}}\}F(p), \end{split}$$

where $F(p) = \max\{F_1(p), F_2(p)\}$. By Lemma 5 in [11] and the definition in [30], we obtain that almost every sample path of $y_1(t)$ is locally but uniformly *Hölder*-continuous with exponent v for every $v \in (0, \frac{p-2}{2p})$. Similarly, it can be proved that almost every sample path of $y_2(t)$ is also uniformly continuous on $t \ge 0$. The proof is completed.



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