

AIMS Mathematics, 7(11): 19865–19890. DOI: 10.3934/math.20221088 Received: 02 July 2022 Revised: 16 August 2022 Accepted: 26 August 2022 Published: 08 September 2022

http://www.aimspress.com/journal/Math

Research article

Determining the global threshold of an epidemic model with general interference function and high-order perturbation

Yassine Sabbar¹, Asad Khan^{2,*}, Anwarud Din^{3,*}, Driss Kiouach¹ and S. P. Rajasekar⁴

- ¹ LPAIS Laboratory, Faculty of Sciences Dhar El Mahraz, Sidi Mohamed Ben Abdellah University, Fez, Morocco
- ² School of Computer Science and Cyber Engineering, Guangzhou University, Guangzhou 510006, China
- ³ Department of Mathematics, Sun Yat-Sen University, Guangzhou, China
- ⁴ Department of Mathematics, Government Arts College for Women, Nilakottai-624202, Tamilnadu, India
- * **Correspondence:** Email: asad@gzhu.edu.cn, anwarud@mail.sysu.edu.cn.

Abstract: This research provides an improved theoretical framework of the Kermack-McKendrick system. By considering the general interference function and the polynomial perturbation, we give the sharp threshold between two situations: the disappearance of the illness and the ergodicity of the higher-order perturbed system. Obviously, the ergodic characteristic indicates the continuation of the infection in the population over time. Our study upgrades and enhances the work of Zhou et al. (2021) and suggests a new path of research that will serve as a basis for future investigations. As an illustrative application, we discuss some special cases of the polynomial perturbation to examine the precision of our outcomes. We deduce that higher order fluctuations positively affect the illness extinction time and lead to its rapid disappearance.

Keywords: epidemic system; stochastic process; extinction; ergodicity; stationarity **Mathematics Subject Classification:** 34A26, 34A12, 92D30, 37C10, 60H30

1. Introduction

Mathematical formulations play a central role in exploring and predicting the future of the communicable diseases [1]. For example, in the case of the coronavirus disease and of course its new mutations, many researchers have contributed to modeling the mechanisms of its spread and providing scientific recommendations to control its expansion [2–5]. It should be noted that most of the epidemiological models presented in these articles are an improved and adapted version of the SIR

model [6]. This compartmentalized system was constructed by Kermack and McKendrick [7] and its philosophy is based on the idea of subdividing individuals according to their different characteristics. Explicitly, individuals are grouped into three main groups: susceptible class (C_1), infected class (C_2) and permanently recovered class (C_3). The transmit rates among these groups are determined by the following dynamical system:

$$\begin{cases} d\mathbf{C}_{1}(t) = \left(\Theta - \mathbf{u}\mathbf{C}_{1}(t) - \mathbf{b}\mathbf{C}_{1}(t)\mathbf{C}_{2}(t)\right)dt, \\ d\mathbf{C}_{2}(t) = \left(\mathbf{b}\mathbf{C}_{1}(t)\mathbf{C}_{2}(t) - (\mathbf{u} + \mathbf{a} + \mathbf{c})\mathbf{C}_{2}(t)\right)dt, \\ d\mathbf{C}_{3}(t) = \left(\mathbf{c}\mathbf{C}_{2}(t) - \mathbf{u}\mathbf{C}_{3}(t)\right)dt, \\ \mathbf{C}_{k}(0) > 0, \quad k = 1, 2, 3, \end{cases}$$
(1.1)

where $\Theta > 0$ designates the inflow rate into \mathbf{C}_1 , $\mathfrak{u} > 0$ indicates the normal death rate, $\mathfrak{b} > 0$ is the dissemination rate of the epidemic, $\mathfrak{a} > 0$ is the mortality ratio due to the infection, and $\mathfrak{c} > 0$ is the cure rate. We note that the above system is one of the most straightforward epidemiological models used to depict the first wave of COVID-19 [8]. Lately, Zhou et al. [9] proposed a general version of system (1.1) by including the pre-existing immunity presumption and the nonlinear incidence function $\mathfrak{b}\mathbf{C}_1(t)g(\mathbf{C}_2(t))$. By mandating certain conditions on g, they introduced the following system:

$$\begin{cases}
 d\mathbf{C}_{1}(t) = \left(\Theta - (u_{1} + z_{3})\mathbf{C}_{1}(t) - b\mathbf{C}_{1}(t)g(\mathbf{C}_{2}(t))\right)dt, \\
 d\mathbf{C}_{2}(t) = \left(b\mathbf{C}_{1}(t)g(\mathbf{C}_{2}(t)) - (u_{2} + a + c)\mathbf{C}_{2}(t)\right)dt, \\
 d\mathbf{C}_{3}(t) = \left(z_{1}(t) + c\mathbf{C}_{2}(t) - u_{3}\mathbf{C}_{3}(t)\right)dt, \\
 \mathbf{C}_{k}(0) > 0, \quad k = 1, 2, 3,
\end{cases}$$
(1.2)

where 3 is the pre-existing immunity rate, and u_1 , u_2 , u_3 are respectively the normal mortality rates of C_1 , C_2 , C_3 . The function *g* covers some functional responses, for instance, $g(C_2) = C_2$ [10], $g(C_2) = \frac{C_2}{m + C_2}$ (m > 0) [11] and $bg(C_2(t)) = b - \frac{b_c}{m + C_2(t)}$, where m > 0 is the media intrusion rate and $b_c > 0$ is the reduced active contact rate [12].

When dealing with epidemiological models, more characteristics can be considered such as crossindividual overlap [13]. It is worth to point out that the choice of functional response affects the prediction of illness behavior. Moreover, the previous setup of Zhou et al. [9] overlooks a large category of incidence rates that are often used in the literature. In this research, we exhibit an enhanced SIR illness system with an interference function that includes additional response examples (see Table 1). In line with this setting, the system (1.2) can be rewritten as follows:

$$\begin{aligned} \left(d\mathbf{C}_{1}(t) = \left(\Theta - (\mathfrak{u}_{1} + \mathfrak{z})\mathbf{C}_{1}(t) - \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t))\mathbf{C}_{2}(t) \right) dt, \\ d\mathbf{C}_{2}(t) = \left(\mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t))\mathbf{C}_{2}(t) - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c})\mathbf{C}_{2}(t) \right) dt, \\ d\mathbf{C}_{3}(t) = \left(\mathfrak{z}_{1}(t) + \mathfrak{c}\mathbf{C}_{2}(t) - \mathfrak{u}_{3}\mathbf{C}_{3}(t) \right) dt, \\ \mathbf{C}_{k}(0) > 0, \quad k = 1, 2, 3. \end{aligned}$$

$$(1.3)$$

We presume that the general interference response $\mathcal{H} \in C^2(\mathbb{R}_+ \times \mathbb{R}_+, \mathbb{R}_+)$ verifies these two hypotheses:

• Assumption (a): There exists a constant $\rho > 0$ such that for all $C_1, C_2 \ge 0$,

$$\frac{\partial \mathcal{H}(\mathbf{C}_1, \mathbf{C}_2)}{\partial \mathbf{C}_2} \le 0 \le \frac{\partial \mathcal{H}(\mathbf{C}_1, \mathbf{C}_2)}{\partial \mathbf{C}_1} \le \varrho.$$

AIMS Mathematics

• Assumption (b): $\lim_{\mathbf{C}_2 \to 0} \sup_{\mathbf{C}_1 > 0} \{ |\mathcal{H}(\mathbf{C}_1, \mathbf{C}_2) - \mathcal{H}(\mathbf{C}_1, 0)| \} = 0.$

The properties (a) and (b) are readily satisfied by the typical examples listed in Table 1.

	Table 1. List of some	prototypes of the	general interference	function \mathcal{F}
--	-----------------------	-------------------	----------------------	------------------------

Name	Expression	Source
Beddington-DeAngelis	$\mathcal{H}(\mathbf{C}_1, \mathbf{C}_2) = \frac{\mathbf{C}_1}{1 + \mathfrak{m}_1 \mathbf{C}_1 + \mathfrak{m}_2 \mathbf{C}_2} \ (\mathfrak{m}_1, \mathfrak{m}_2 > 0)$	[14]
Crowley-Martin	$\mathcal{H}(\mathbf{C}_1, \mathbf{C}_2) = \frac{\mathbf{C}_1}{(\mathfrak{m}_1 + \mathbf{C}_1)(\mathfrak{m}_2 + \mathbf{C}_2)} \ (\mathfrak{m}_1, \mathfrak{m}_2 > 0)$	[15]
Modified Crowley-Martin	$\mathcal{H}(\mathbf{C}_1, \mathbf{C}_2) = \frac{\mathbf{C}_1}{1 + \mathfrak{m}_1 \mathbf{C}_1 + \mathfrak{m}_2 \mathbf{C}_2 + \mathfrak{m}_3 \mathbf{C}_1 \mathbf{C}_2} (\mathfrak{m}_1, \mathfrak{m}_2, \mathfrak{m}_3 > 0)$	[13]

Substantially, the infection mechanism is random in nature at all scales [16–20]. From the attachment of the virus to the human cell to the encapsulation of repetitive genetic information, from the shedding of new virions to the transmission of a second individual, from individual behavior to global mobility, all the factors influence the diffusion of the infection and make it more uphill task to foretell its conduct [21–23]. The stochastic approach offers considerable advantages in providing insight into population dynamics under the said fluctuations [24–26]. To correctly describe this randomness, a series of perturbed compartmental models with different assumptions that simulate reality have been proposed [27–31]. Most of these models assume that random fluctuations can be modeled by integrating Brownian motions into the deterministic formulation [32]. By selecting this class of fluctuations in their linear shape, probabilistic systems are widely used in epidemiology to analyze disease prevalence [33–35]. In these works, the primary focus was the investigation of some biological long-run characteristics of the infection.

Since 2017, a new form of stochastic systems has emerged where the story begins with the work proposed by Liu and Jiang in [36]. By reason of the complexity of environmental changes, they claimed that the relative linear order of the disturbance can be raised to the second. Based on this assumption, scientific papers have proposed and analyzed various real systems with second-order fluctuations [37, 38]. Recently, the autrhors of [9] suggested an enhanced type of perturbation in its general representation. This frame generalizes the previously mentioned studies and offers a new line of research. Motivated by their arguments, we consider the following polynomial perturbed system:

$$\begin{cases} \underbrace{\operatorname{Deterministic part}}_{\operatorname{Deterministic part}} \left(\Theta - (\mathfrak{u}_{1} + \mathfrak{z}) \mathbf{C}_{1}(t) - \mathfrak{b} \mathcal{H}(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) \mathbf{C}_{2}(t) \right) dt + \sum_{h=0}^{N} \mathfrak{q}_{1h} \mathbf{C}_{1}^{h+1}(t) d\mathbb{W}_{1}(t), \\ d\mathbf{C}_{2}(t) = \left(\mathfrak{b} \mathcal{H}(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) \mathbf{C}_{2}(t) - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) \mathbf{C}_{2}(t) \right) dt + \sum_{h=0}^{N} \mathfrak{q}_{2h} \mathbf{C}_{2}^{h+1}(t) d\mathbb{W}_{2}(t), \\ d\mathbf{C}_{3}(t) = \left(\mathfrak{z}_{1}(t) + \mathfrak{c} \mathbf{C}_{2}(t) - \mathfrak{u}_{3} \mathbf{C}_{3}(t) \right) dt + \sum_{h=0}^{N} \mathfrak{q}_{3h} \mathbf{C}_{3}^{h+1}(t) d\mathbb{W}_{3}(t), \end{cases}$$
(1.4)

where $\mathbb{W}_1(t)$, $\mathbb{W}_2(t)$, $\mathbb{W}_3(t)$ are independent Brownian motions defined on a filtered probability space $\Omega^{\mathcal{E},\mathbb{P}} \equiv (\Omega, \mathcal{E}, \{\mathcal{E}_t\}_{t\geq 0}, \mathbb{P})$ such that $\{\mathcal{E}_t\}_{t\geq 0}$ follows the usual assumptions, and $\mathfrak{q}_{kh} > 0$ (k = 1, 2, 3, h = 0, 1, 2, ..., N) are the high-order intensities of white noises.

In [9], the authors indicated that the model (1.4) in the case of incidence $bC_1(t)g(C_2(t))$ and polynomial perturbation is well-posed mathematically and biologically, then they treated its longrun behavior. The problem is that they obtained two separate critical conditions for extinction and stationarity which is not ideal when addressing epidemiological models. Moreover, they mention that there was a large gap between the defined criteria; and the corresponding threshold value is still an open question (for more details see the discussion part of [9]). Compared to the results presented in [9], in this article, we address the said problem from a global angle by considering an epidemic model with a general incidence function. Taking the latter into account makes the analysis very complex, which has prompted us to innovate alternative techniques. By adding the polynomial perturbation, we present the acute threshold value between stationarity and extinction of the infection, which offers an excellent insight into the possible scenarios of epidemic status in a given population.

Technically, we introduce an analytical method based on some long-term characteristics of an auxiliary boundary equation [39]. By using the ergodic theorem and the stochastic comparison lemma, we establish the well-defined threshold between stationarity and infection extinction. Our method differs from the one used in [9] by using the mutually exclusive possibilities lemma and other analytical tools. Specifically, we focus on the long-term characteristics of the Markov process $\mathbf{D}(t)$ that verifies

$$\begin{cases} d\mathbf{D}(t) = \left\{ \Theta - (\mathfrak{u}_1 + \mathfrak{z})\mathbf{D}(t) \right\} dt + \sum_{h=0}^{N} \mathfrak{q}_{1h} \mathbf{D}^{h+1}(t) d\mathbb{W}_1(t), \\ \mathbf{D}(0) = \mathbf{C}_1(0). \end{cases}$$
(1.5)

In accordance with Lemma 5 of [9], **D** admits the following single stable distribution:

$$\pi^{\mathbf{D}}(y) = C^{\mathbf{D}} \left(\sum_{h=0}^{N} \mathfrak{q}_{1h} y^{h+1}(t) \right)^{-2} \exp\left\{ 2 \int_{\frac{\Theta}{\mathfrak{u}+\mathfrak{z}}}^{y} \left(\Theta - (\mathfrak{u}_{1}+\mathfrak{z})\tau \right) \left(\sum_{h=0}^{N} \mathfrak{q}_{1h} \tau^{h+1}(t) \right)^{-2} \mathrm{d}\tau \right\},\$$

where $C^{\mathbf{D}}$ is a specific constant that verifies $\int_{\mathbb{R}_{+}} \pi^{\mathbf{D}}(y) dy = 1$. From the probabilistic comparison result [40], we can compare the processes \mathbf{D} and \mathbf{C}_{1} as follows: $\mathbf{D}(t) \ge \mathbf{C}_{1}(t)$ almost surely (a.s.). Furthermore, the time average of $\mathbf{D}(t)$ converges almost surely to $\int_{\mathbb{R}_{+}} y\pi^{\mathbf{D}}(dy)$ as $t \to \infty$. In accordance with the above results, we clearly state that the present work aims to prove that the following quantity

$$\mathcal{T}_{\circ}^{\Sigma} = \mathfrak{b} \int_{\mathbb{R}_{+}} \mathcal{H}(y,0) \pi^{\mathbf{D}}(\mathrm{d}y) - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) - 0.5\mathfrak{q}_{20}^{2}$$

is the sharp threshold between the disappearance of the illness and the ergodic characteristic of the system (1.4), and its sign provides a stellar overview of the potential scenarios of the epidemic situation. In other words, this article proposes a nice generalization of the article [9] and presents a new treatment applicable to other complex models. Importantly, we show numerically that the extreme amount of disturbance reduces the disease extinction time.

The rest of the paper is organized as follows: In Section 2, we show that the disappearance scenario occurs when $T_{\circ}^{\Sigma} < 0$. In Section 3, we prove that the scenario of the ergodicity of system (1.4) occurs when $T_{\circ}^{\Sigma} > 0$. In Section 4, we verify numerically the correctness of our outcomes. Finally, we conclude this paper in Section 5 with a discussion.

2. Scenario 1: disease disappearance

This section aims to exhibit the criterion for the demise of the infection.

Theorem 2.1. The disappearance of the disease occurs if $\mathcal{T}^{\Sigma}_{\circ} < 0$.

Proof. By employing Itô's lemma for drift-diffusion processes [41], we obtain

$$\mathrm{d}\ln\mathbf{C}_{2}(t) = \left(\mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t)) - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) - 0.5\left(\sum_{h=0}^{N}\mathfrak{q}_{2h}\mathbf{C}_{2}^{h}(t)\right)^{2}\right)\mathrm{d}t + \sum_{h=0}^{N}\mathfrak{q}_{2h}\mathbf{C}_{2}^{h}(t)\mathrm{d}\mathbb{W}_{2}(t).$$

In line with the probabilistic comparison lemma, we conclude that

$$\operatorname{d}\ln \mathbf{C}_{2}(t) \leq \left(\mathfrak{b}\mathcal{H}(\mathbf{D}(t),0) - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) - 0.5\left(\sum_{h=0}^{N}\mathfrak{q}_{2h}\mathbf{C}_{2}^{h}(t)\right)^{2}\right)\operatorname{d}t + \sum_{h=0}^{N}\mathfrak{q}_{2h}\mathbf{C}_{2}^{h}(t)\operatorname{d}\mathbb{W}_{2}(t).$$
(2.1)

After that, we make two operations on both sides of (2.1): integration from 0 to *t* and division by *t*, then the result is

$$t^{-1} \ln \mathbf{C}_{2}(t) - t^{-1} \ln \mathbf{C}_{2}(0) \leq t^{-1} \mathfrak{b} \int_{0}^{t} \mathcal{H}(\mathbf{D}(\tau), 0) d\tau - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) + t^{-1} \underbrace{\left(\int_{0}^{t} \sum_{h=0}^{N} \mathfrak{q}_{2h} \mathbf{C}_{2}^{h}(\tau) d\mathbb{W}_{2}(\tau) - 0.5 \int_{0}^{t} \left(\sum_{h=0}^{N} \mathfrak{q}_{2h} \mathbf{C}_{2}^{h}(\tau) \right)^{2} d\tau \right)}_{=\mathcal{G}(t)}.$$
(2.2)

The next step is based on the use of the exponential inequality for martingales [41], which leads to

$$\mathbb{P}\left\{\sup_{t\in[0,T_1]}\left(\int_0^t\sum_{h=0}^N\mathfrak{q}_{2h}\mathbf{C}_2^h(\tau)\mathrm{d}\mathbb{W}_2(\tau)-0.5\alpha_1\int_0^t\left(\sum_{h=0}^N\mathfrak{q}_{2h}\mathbf{C}_2^h(\tau)\right)^2\mathrm{d}\tau\right)>\frac{2\ln T_1}{\alpha_1}\right\}\leq T_1^{-2},$$

for all $0 < \alpha_1 < 1$ and $T_1 > 0$. From the Borel-Cantelli result [41], we assure the existence of $T_{1,\omega} = T_1(\omega)$, $\forall \omega$ in Ω , such that

$$\int_{0}^{t} \sum_{h=0}^{N} \mathfrak{q}_{2h} \mathbf{C}_{2}^{h}(\tau) \mathrm{d} \mathbb{W}_{2}(\tau) \leq \frac{2 \ln T_{1}}{\alpha_{1}} + 0.5 \alpha_{1} \int_{0}^{t} \left(\sum_{h=0}^{N} \mathfrak{q}_{2h} \mathbf{C}_{2}^{h}(\tau) \right)^{2} \mathrm{d} \tau$$

holds for all $T_1 \ge T_{1,\omega}$ and $T_1 - 1 < t \le T_1$ a.s. Therefore,

$$\begin{split} t^{-1}\mathcal{G}(t) &\leq \frac{2\ln T_1}{\alpha_1 t} + t^{-1}0.5\alpha_1 \int_0^t \left(\sum_{h=0}^N \mathfrak{q}_{2h} \mathbf{C}_2^h(\tau)\right)^2 \mathrm{d}\tau - t^{-1}0.5 \int_0^t \left(\sum_{h=0}^N \mathfrak{q}_{2h} \mathbf{C}_2^h(\tau)\right)^2 \mathrm{d}\tau \\ &\leq \frac{2\ln T_1}{\alpha_1(T_1 - 1)} - t^{-1}0.5(1 - \alpha_1) \int_0^t \left(\sum_{h=0}^N \mathfrak{q}_{2h} \mathbf{C}_2^h(\tau)\right)^2 \mathrm{d}\tau \\ &\leq \frac{2\ln T_1}{\alpha(T_1 - 1)} - 0.5(1 - \alpha_1)\mathfrak{q}_{20}^2. \end{split}$$

AIMS Mathematics

By taking the limitsup on two sides of (2.2), we infer that

$$\begin{split} \limsup_{t \to \infty} t^{-1} \ln \mathbf{C}_2(t) &\leq \operatorname{blim}_{t \to \infty} t^{-1} \int_0^t \mathcal{H}(\mathbf{D}(\tau), 0) \mathrm{d}\tau - (\mathfrak{u}_2 + \mathfrak{a} + \mathfrak{c}) + \operatorname{lim}_{t \to \infty} t^{-1} \mathcal{G}(t) \\ &\leq \mathfrak{b} \int_{\mathbb{R}_+} \mathcal{H}(y, 0) \pi^{\mathbf{D}}(\mathrm{d}y) - (\mathfrak{u}_2 + \mathfrak{a} + \mathfrak{c}) + \operatorname{lim}_{T_1 \to \infty} \frac{2 \ln T_1}{\alpha(T_1 - 1)} - 0.5(1 - \alpha_1) \mathfrak{q}_{20}^2 \\ &= \mathfrak{b} \int_{\mathbb{R}_+} \mathcal{H}(y, 0) \pi^{\mathbf{D}}(\mathrm{d}y) - (\mathfrak{u}_2 + \mathfrak{a} + \mathfrak{c}) - 0.5(1 - \alpha_1) \mathfrak{q}_{20}^2 \quad \text{a.s.} \end{split}$$

We let α_1 tends to 0^+ , then the obtained result is

$$\limsup_{t\to\infty} t^{-1} \ln \mathbf{C}_2(t) \le \mathcal{T}_{\circ}^{\Sigma} < 0 \quad \text{a.s.}$$

That implies the stochastic extinction $\lim_{t\to\infty} C_2(t) = 0$ a.s. In other words, the class of individuals carrying the infection will disappear.

3. Scenario 2: the ergodicity of the model (1.4)

This section introduces a new approach to establish the criterion of the ergodicity of our probabilistic system (1.4).

Theorem 3.1. The single ergodic stable distribution of the probabilistic model (1.4) exists if $\mathcal{T}_{\circ}^{\Sigma} > 0$.

Proof. In order to reduce notations and provide clear mathematical writing, we set

$$\left(\sum_{h=0}^{N}\mathfrak{q}_{2h}\mathbf{C}_{2}^{h}(t)\right)^{2} = \sum_{h=0}^{2N} \underbrace{\left(\sum_{n+m=h}\mathfrak{q}_{2n}\mathfrak{q}_{2m}\right)}_{\mathfrak{p}_{h}}\mathbf{C}_{2}^{h}(t) = \sum_{h=0}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h}(t).$$

The Itô differential operator \mathcal{L} associated with the stochastic equation of $C_2(t)$ is given by

$$\mathcal{L}(-\ln \mathbf{C}_{2}(t)) = -b\mathcal{H}(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) + (u_{2} + a + c) + 0.5 \left(\sum_{h=0}^{N} \mathfrak{q}_{2h} \mathbf{C}_{2}^{h}(t)\right)^{2}$$

= $-b\mathcal{H}(\mathbf{D}(t), 0) + b\mathcal{H}(\mathbf{D}(t), 0) - b\mathcal{H}(\mathbf{C}_{1}(t), 0) + b\mathcal{H}(\mathbf{C}_{1}(t), 0) - b\mathcal{H}(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t))$
+ $(u_{2} + a + c) + 0.5\mathfrak{q}_{20}^{2} + \mathfrak{q}_{20}\mathfrak{q}_{21}\mathbf{C}_{2}(t) + 0.5\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h}(t).$

From Assumption (a), we have

$$\mathcal{L}(-\ln \mathbf{C}_{2}(t)) \leq -\mathfrak{b}\mathcal{H}(\mathbf{D}(t),0) + (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) + 0.5\mathfrak{q}_{20}^{2} + \mathfrak{b}\varrho(\mathbf{D}(t) - \mathbf{C}_{1}(t)) + \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t),0) - \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t)) + \mathfrak{q}_{20}\mathfrak{q}_{21}\mathbf{C}_{2}(t) + 0.5\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h}(t).$$
(3.1)

AIMS Mathematics

Now, the application of \mathcal{L} on $(\ln \mathbf{D}(t) - \ln \mathbf{C}_1(t))$ gives

$$\mathcal{L}\left(\ln \mathbf{D}(t) - \ln \mathbf{C}_{1}(t)\right) \leq \mathfrak{A}\left(\mathbf{D}^{-1}(t) - \mathbf{C}_{1}^{-1}(t)\right) + \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t))\mathbf{C}_{2}(t)\mathbf{C}_{1}^{-1}(t)$$
$$- 0.5\left(\sum_{h=0}^{N}\mathfrak{q}_{1h}\mathbf{D}^{h}(t)\right)^{2} + 0.5\left(\sum_{h=0}^{N}\mathfrak{q}_{1h}\mathbf{C}_{1}^{h}(t)\right)^{2}.$$

Since $\mathbf{D}(t) \ge \mathbf{C}_1(t)$ a.s., we obtain

$$\mathcal{L}\left(\ln \mathbf{D}(t) - \ln \mathbf{C}_{1}(t)\right) \leq -\mathfrak{q}_{10}\mathfrak{q}_{11}(\mathbf{D}(t) - \mathbf{C}_{1}(t)) + \mathfrak{b}\varrho\mathbf{C}_{2}(t).$$
(3.2)

We define the function $\Phi(t)$ as follows:

$$\Phi(t) = -\ln \mathbf{C}_2(t) + \frac{b\varrho}{\mathfrak{q}_{10}\mathfrak{q}_{11}} \Big(\ln \mathbf{D}(t) - \ln \mathbf{C}_1(t) \Big).$$

Based on (3.1) and (3.2), we get

$$\mathcal{L}\Phi(t) \leq -\mathfrak{b}\mathcal{H}(\mathbf{D}(t), 0) + (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) + 0.5\mathfrak{q}_{20}^{2} + \left(\mathfrak{q}_{20}\mathfrak{q}_{21} + \frac{\mathfrak{b}^{2}\varrho^{2}}{\mathfrak{q}_{10}\mathfrak{q}_{11}}\right)\mathbf{C}_{2}(t) + 0.5\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h}(t) + \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t), 0) - \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)).$$
(3.3)

We add and subtract at the same time the quantity $\mathfrak{b} \int_{\mathbb{R}_+} \mathcal{H}(y,0) \pi^{\mathbf{D}}(dy)$ in (3.3) as follows:

$$\begin{aligned} \mathcal{L}\Phi(t) &\leq -\mathfrak{b} \int_{\mathbb{R}_{+}} \mathcal{H}(y,0)\pi^{\mathbf{D}}(\mathrm{d}y) + (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) + 0.5\mathfrak{q}_{20}^{2} + \mathfrak{b}\left(\int_{\mathbb{R}_{+}} \mathcal{H}(y,0)\pi^{\mathbf{D}}(\mathrm{d}y) - \mathcal{H}(\mathbf{D}(t),0)\right) \\ &+ \left(\mathfrak{q}_{20}\mathfrak{q}_{21} + \frac{\mathfrak{b}^{2}\varrho^{2}}{\mathfrak{q}_{10}\mathfrak{q}_{11}}\right)\mathbf{C}_{2}(t) + 0.5\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h}(t) + \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t),0) - \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t)). \end{aligned}$$

To eliminate the term associated with $C_2(t)$, we set

$$\Phi_{\Theta}(t) = -\ln \mathbf{C}_{2}(t) + \frac{\mathfrak{b}\varrho}{\mathfrak{q}_{10}\mathfrak{q}_{11}} \Big(\ln \mathbf{D}(t) - \ln \mathbf{C}_{1}(t) \Big) + \Theta \mathbf{C}_{2}(t),$$

where the positive constant Θ verifies

$$\Theta(\mathfrak{u}_{2}+\mathfrak{a}+\mathfrak{c})\geq\left(\mathfrak{q}_{20}\mathfrak{q}_{21}+\frac{\mathfrak{b}^{2}\varrho^{2}}{\mathfrak{q}_{10}\mathfrak{q}_{11}}\right).$$

The application of \mathcal{L} on $\Phi_{\Theta}(t)$ gives

$$\mathcal{L}\Phi_{\Theta}(t) \leq \underbrace{-\mathfrak{b}\int_{\mathbb{R}_{+}}\mathcal{H}(y,0)\pi^{\mathbf{D}}(\mathrm{d}y) + (\mathfrak{u}_{2}+\mathfrak{a}+\mathfrak{c}) + 0.5\mathfrak{q}_{20}^{2}}_{h=2} + \mathfrak{b}\left(\int_{\mathbb{R}_{+}}\mathcal{H}(y,0)\pi^{\mathbf{D}}(\mathrm{d}y) - \mathcal{H}(\mathbf{D}(t),0)\right) + \Theta\mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t))\mathbf{C}_{2}(t) + 0.5\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h}(t) + \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t),0) - \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t)).$$

In the same vein, we apply \mathcal{L} on the function $\zeta^{-1}(1 + \mathbf{C}_1(t))^{\zeta} + \zeta^{-1}\mathbf{C}_2^{\zeta}(t), \zeta \in (0, 1)$, then

$$\begin{aligned} \mathcal{L}(\zeta^{-1}(1+\mathbf{C}_{1}(t))^{\zeta}+\zeta^{-1}\mathbf{C}_{2}^{\zeta}(t)) &= (1+\mathbf{C}_{1}(t))^{\zeta-1} \Big(\Theta - (\mathfrak{u}_{1}+\mathfrak{z})\mathbf{C}_{1}(t) - \mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t))\mathbf{C}_{2}(t)\Big) \\ &+ 0.5(\zeta-1)(1+\mathbf{C}_{1}(t))^{\zeta-2} \left(\sum_{h=0}^{N} \mathfrak{q}_{1h}\mathbf{C}_{1}^{h+1}(t)\right)^{2} \\ &+ \mathbf{C}_{2}^{\zeta-1}(t) \Big(\mathfrak{b}\mathcal{H}(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t))\mathbf{C}_{2}(t) - (\mathfrak{u}_{2}+\mathfrak{a}+\mathfrak{c})\mathbf{C}_{2}(t)\Big) \\ &+ 0.5(\zeta-1)\mathbf{C}_{2}^{\zeta-2}(t) \left(\sum_{h=0}^{N} \mathfrak{q}_{2h}\mathbf{C}_{2}^{h+1}(t)\right)^{2}. \end{aligned}$$

Accordingly, we derive that

$$\begin{split} \mathcal{L} \Big(\zeta^{-1} \big(1 + \mathbf{C}_{1}(t) \big)^{\zeta} + \zeta^{-1} \mathbf{C}_{2}^{\zeta}(t) \Big) &\leq \mathfrak{A} - 0.5 (1 - \zeta) \mathfrak{q}_{11}^{2} \mathbf{C}_{1}^{\zeta+2}(t) + \mathfrak{b} \varrho \mathbf{C}_{1}(t) \mathbf{C}_{2}^{\zeta}(t) \\ &- \big((\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) + 0.5 (1 - \zeta) \mathfrak{q}_{20}^{2} \big) \mathbf{C}_{2}^{\zeta}(t) \\ &+ (1 - \zeta) \mathfrak{q}_{20} \mathfrak{q}_{21} \mathbf{C}_{2}^{\zeta+1}(t) - 0.5 (1 - \zeta) \mathfrak{q}_{21}^{2} \mathbf{C}_{2}^{\zeta+2}(t) \\ &- 0.5 (1 - \zeta) \sum_{h=2}^{2N} \mathfrak{p}_{h} \mathbf{C}_{2}^{h+\zeta}(t) \\ &\leq \Theta - 0.5 (1 - \zeta) \mathfrak{q}_{11}^{2} \mathbf{C}_{1}^{\zeta+2}(t) + \mathfrak{b} \varrho (\zeta + 1)^{-1} \mathbf{C}_{1}^{\zeta+1}(t) \\ &+ \mathfrak{b} \varrho \zeta (\zeta + 1)^{-1} \mathbf{C}_{2}^{\zeta+1}(t) - 0.5 (1 - \zeta) \mathfrak{q}_{21}^{2} \mathbf{C}_{2}^{\zeta+2}(t) \\ &- 0.5 (1 - \zeta) \sum_{h=2}^{2N} \mathfrak{p}_{h} \mathbf{C}_{2}^{h+\zeta}(t). \end{split}$$

Now, we define a new function $\Phi_{\Theta,\zeta}$ as follows:

$$\Phi_{\Theta,\zeta}(\mathbf{C}_1(t),\mathbf{C}_2(t)) = S\Phi_{\Theta}(t) + \zeta^{-1}(1+\mathbf{C}_1(t))^{\zeta} + \zeta^{-1}\mathbf{C}_2^{\zeta}(t),$$

where S > 0 satisfies that $-S\mathcal{T}_{\circ}^{\Sigma} + \mathcal{Z} + 2 \le 0$ and \mathcal{Z} is given by

$$\mathcal{Z} = \max \left\{ \sup_{(\mathbf{C}_{1},\mathbf{C}_{2})\in\mathbb{R}^{2}_{+,\star}} \left\{ \Theta + \mathfrak{b}\varrho(\zeta+1)^{-1}\mathbf{C}_{1}^{\zeta+1}(t) - 0.25(1-\zeta)\mathfrak{q}_{11}^{2}\mathbf{C}_{1}^{\zeta+2}(t) + \mathfrak{b}\varrho\zeta(\zeta+1)^{-1}\mathbf{C}_{2}^{\zeta+1}(t) - 0.25(1-\zeta)\mathfrak{q}_{21}^{2}\mathbf{C}_{2}^{\zeta+2}(t) + 0.5\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h}(t) - 0.5(1-\zeta)\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h+\zeta}(t) \right\}, 1 \right\}.$$

AIMS Mathematics

Clearly, the function $\Phi_{\Theta,\zeta}$ reaches its minimum value at a point $(\mathbf{C}_1^{\ell}, \mathbf{C}_2^{\ell})$. For this reason, we will consider a new non-negative function defined as follows:

$$\Phi_{\Theta,\zeta}^{\Sigma}(\mathbf{C}_1(t),\mathbf{C}_2(t)) = S\Phi_{\Theta}(t) + \zeta^{-1}(1+\mathbf{C}_1(t))^{\zeta} + \zeta^{-1}\mathbf{C}_2^{\zeta}(t) - \Phi_{\Theta,\zeta}(\mathbf{C}_1^{\ell},\mathbf{C}_2^{\ell}).$$

From the above calculation, we obtain

$$\begin{split} \mathcal{L}\Phi_{\Theta,\zeta}^{\Sigma}(t) &\leq -S \mathcal{T}_{\circ}^{\Sigma} + S \,\Theta b \varrho \mathbf{C}_{1}(t) \mathbf{C}_{2}(t) + b S \left(\mathcal{H}(\mathbf{C}_{1}(t),0) - \mathcal{H}(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t)) \right) + 0.5 \sum_{h=2}^{2N} \mathfrak{p}_{h} \mathbf{C}_{2}^{h}(t) \\ &+ \Theta - 0.5(1-\zeta) \mathfrak{q}_{11}^{2} \mathbf{C}_{1}^{\zeta+2}(t) + \mathfrak{b} \varrho(\zeta+1)^{-1} \mathbf{C}_{1}^{\zeta+1}(t) + \mathfrak{b} \varrho\zeta(\zeta+1)^{-1} \mathbf{C}_{2}^{\zeta+1}(t) \\ &- 0.5(1-\zeta) \mathfrak{q}_{21}^{2} \mathbf{C}_{2}^{\zeta+2}(t) - 0.5(1-\zeta) \sum_{h=2}^{2N} \mathfrak{p}_{h} \mathbf{C}_{2}^{h+\zeta}(t) + \mathfrak{b} \left(\int_{\mathbb{R}_{+}} \mathcal{H}(y,0) \pi^{\mathbf{D}}(\mathrm{d} y) - \mathcal{H}(\mathbf{D}(t),0) \right) \\ &= \Psi(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t)) + \mathfrak{b} \left(\int_{\mathbb{R}_{+}} \mathcal{H}(y,0) \pi^{\mathbf{D}}(\mathrm{d} y) - \mathcal{H}(\mathbf{D}(t),0) \right). \end{split}$$

Now, we define five sets:

$$\begin{aligned} \mathcal{J}_{a,a_{\star}} &= \left\{ (\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) \in \mathbb{R}^{2,\star}_{+} | \ a \leq \mathbf{C}_{1}(t) \leq a^{-1}, \ a_{\star} \leq \mathbf{C}_{2}(t) \leq a_{\star}^{-1} \right\}, \\ \mathcal{J}_{a,1} &= \left\{ (\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) \in \mathbb{R}^{2,\star}_{+} | \ 0 < \mathbf{C}_{1}(t) < a \right\}, \\ \mathcal{J}_{a_{\star},2} &= \left\{ (\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) \in \mathbb{R}^{2,\star}_{+} | \ 0 < \mathbf{C}_{2}(t) < a_{\star} \right\}, \\ \mathcal{J}_{a,3} &= \left\{ (\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) \in \mathbb{R}^{2,\star}_{+} | \ \mathbf{C}_{1}(t) > a^{-1} \right\}, \\ \mathcal{J}_{a_{\star},4} &= \left\{ (\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) \in \mathbb{R}^{2,\star}_{+} | \ \mathbf{C}_{2}(t) > a_{\star}^{-1} \right\}. \end{aligned}$$

Here, $\mathbb{R}^{2,\star}_+ = \{(x, y) : x > 0, y > 0\}, a_{\star} = \min\{a_{\circ}, a\}$, where $a_{\circ} > 0$ verifies (3.9), and a > 0 is chosen carefully such that

$$S \Theta \mathfrak{b}\varrho a + \mathfrak{b} S \varrho a + \frac{\zeta S \Theta \mathfrak{b}\varrho a}{\zeta + 2} \left(\frac{2S \Theta \mathfrak{b}\varrho a}{0.25(\zeta + 2)(1 - \zeta)\mathfrak{q}_{21}^2} \right)^{2\zeta^{-1}} - 1 \le 0, \tag{3.4}$$

$$S \Theta b\varrho a + \frac{\zeta S \Theta b\varrho a}{\zeta + 2} \left(\frac{2S \Theta b\varrho a}{0.25(\zeta + 2)(1 - \zeta)\mathfrak{q}_{11}^2} \right)^{2\zeta} - 1 < 0, \tag{3.5}$$

$$-S\mathcal{T}_{\circ}^{\Sigma} + \mathcal{U} - 0.25(1-\zeta)\mathfrak{q}_{11}^2 a^{-\zeta-2} + 1 \le 0,$$
(3.6)

$$-S\mathcal{T}_{\circ}^{\Sigma} + \mathcal{U} - 0.25(1-\zeta)\mathfrak{q}_{21}^{2}a^{-\zeta-2} + 1 \le 0,$$
(3.7)

where

$$\begin{aligned} \mathcal{U} &= \sup_{(\mathbf{C}_1, \mathbf{C}_2) \in \mathbb{R}^{2,\star}_+} \Big\{ 0.5S \,\Theta \mathfrak{b}_{\mathcal{Q}} \mathbf{C}_1^2(t) + 0.5S \,\Theta \mathfrak{b}_{\mathcal{Q}} \mathbf{C}_2^2(t) + S \,\mathfrak{b}_{\mathcal{Q}} \mathbf{C}_1(t) + \Theta + \mathfrak{b}_{\mathcal{Q}}(\zeta + 1)^{-1} \mathbf{C}_1^{\zeta + 1}(t) \\ &+ \mathfrak{b}_{\mathcal{Q}} \zeta(\zeta + 1)^{-1} \mathbf{C}_2^{\zeta + 1}(t) - 0.25(1 - \zeta) \mathfrak{q}_{11}^2 \mathbf{C}_1^{\zeta + 2}(t) - 0.25(1 - \zeta) \mathfrak{q}_{21}^2 \mathbf{C}_2^{\zeta + 2}(t) \\ &+ 0.5 \sum_{h=2}^{2N} \mathfrak{p}_h \mathbf{C}_2^h(t) - 0.5(1 - \zeta) \sum_{h=2}^{2N} \mathfrak{p}_h \mathbf{C}_2^{h+\zeta}(t) \Big\}. \end{aligned}$$

AIMS Mathematics

Plainly, $\mathcal{J}_{a,a_{\star}}^{c} = \mathbb{R}^{2,\star}_{+} \setminus \mathcal{J}_{a,a_{\star}} = \mathcal{J}_{a,1} \cup \mathcal{J}_{a_{\star},2} \cup \mathcal{J}_{a,3} \cup \mathcal{J}_{a_{\star},4}$. In the following, we will verify that

$$\Psi(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) + 1 \le 0, \tag{3.8}$$

for any $(\mathbf{C}_1(t), \mathbf{C}_2(t)) \in \mathcal{J}_{a,a_{\star}}^c$ which is equivalent to showing it on $\mathcal{J}_{a,1}, \mathcal{J}_{a_{\star},2}, \mathcal{J}_{a,3}$ and $\mathcal{J}_{a_{\star},4}$, respectively. For this reason, we have the following situations:

(1) Assume that $(\mathbf{C}_1(t), \mathbf{C}_2(t)) \in \mathcal{J}_{a,1}$. From (3.4), we obtain

$$\begin{split} \Psi(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t)) &\leq -S\mathcal{T}_{\circ}^{\Sigma} + S\Theta b\varrho a + bS\varrho a + S\Theta b\varrho a \mathbf{C}_{2}^{2}(t) - 0.25(1-\zeta)\mathfrak{q}_{11}^{2}\mathbf{C}_{1}^{\zeta+2}(t) \\ &+ 0.5\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h}(t) + \Theta + b\varrho(\zeta+1)^{-1}\mathbf{C}_{1}(t)^{\zeta+1} + b\varrho\zeta(\zeta+1)^{-1}\mathbf{C}_{2}^{\zeta+1}(t) \\ &- 0.25(1-\zeta)\mathfrak{q}_{21}^{2}\mathbf{C}_{2}^{\zeta+2}(t) - 0.25(1-\zeta)\mathfrak{q}_{21}^{2}\mathbf{C}_{2}^{\zeta+2}(t) - 0.5(1-\zeta)\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h+\zeta}(t) \\ &\leq -S\mathcal{T}_{\circ}^{\Sigma} + \mathcal{Z} + S\Theta b\varrho a + bS\varrho a + \frac{\zeta S\Theta b\varrho a}{\zeta+2} \left(\frac{2S\Theta b\varrho a}{0.25(\zeta+2)(1-\zeta)\mathfrak{q}_{21}^{2}}\right)^{2\zeta^{-1}} \\ &\leq -1. \end{split}$$

(2) Here, we use the uniform continuity at $C_2 = 0$ of the function $\mathcal{H}(C_1(t), C_2(t))$. By Assumption (b), $\exists a_0 > 0$ such that as $0 < C_2 \le a_0$,

$$S \Theta \mathfrak{b} \varrho a + \frac{\zeta S \Theta \mathfrak{b} \varrho a}{\zeta + 2} \left(\frac{2S \Theta \mathfrak{b} \varrho a}{0.25(\zeta + 2)(1 - \zeta)\mathfrak{q}_{11}^2} \right)^{2\zeta^{-1}} + \mathfrak{b} S \left(\mathcal{H}(\mathbf{C}_1(t), 0) - \mathcal{H}(\mathbf{C}_1(t), \mathbf{C}_2(t)) \right) < 1.$$
(3.9)

Consequently, if $C_2 < a_{\star} = \min\{a_{\circ}, a\}$, we get from (3.5) that

$$\begin{split} \Psi(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) \\ &\leq -S\mathcal{T}_{\circ}^{\Sigma} + S \,\Theta b \varrho a + S \,\Theta b \varrho a \mathbf{C}_{1}^{2}(t) - 0.25(1 - \zeta) \mathfrak{q}_{11}^{2} \mathbf{C}_{1}^{\zeta+2}(t) + 0.5 \sum_{h=2}^{2N} \mathfrak{p}_{h} \mathbf{C}_{2}^{h}(t) + \Theta \\ &+ b S \left(\mathcal{H}(\mathbf{C}_{1}(t), 0) - \mathcal{H}(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) \right) + b \varrho (\zeta + 1)^{-1} \mathbf{C}_{1}(t)^{\zeta+1} + b \varrho \zeta (\zeta + 1)^{-1} \mathbf{C}_{2}^{\zeta+1}(t) \\ &- 0.25(1 - \zeta) \mathfrak{q}_{21}^{2} \mathbf{C}_{2}^{\zeta+2}(t) - 0.5(1 - \zeta) \sum_{h=2}^{2N} \mathfrak{p}_{h} \mathbf{C}_{2}^{h+\zeta}(t) \\ &\leq -S\mathcal{T}_{\circ}^{\Sigma} + \mathcal{Z} + S \,\Theta b \varrho a + \frac{\zeta S \,\Theta b \varrho a}{\zeta + 2} \left(\frac{2S \,\Theta b \varrho a}{0.25(\zeta + 2)(1 - \zeta) \mathfrak{q}_{11}^{2}} \right)^{2\zeta^{-1}} \\ &+ b S \left(\mathcal{H}(\mathbf{C}_{1}(t), 0) - \mathcal{H}(\mathbf{C}_{1}(t), \mathbf{C}_{2}(t)) \right) \\ &\leq -1. \end{split}$$

(3) Assume that $(\mathbf{C}_1(t), \mathbf{C}_2(t)) \in \mathcal{J}_{a,3}$. From (3.6), we have

$$\begin{split} \Psi(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t)) &\leq -S\mathcal{T}_{\circ}^{\Sigma} - 0.25(1-\zeta)\mathfrak{q}_{11}^{2}\mathbf{C}_{1}^{\zeta+2}(t) + 0.5S\,\Theta\mathfrak{b}\varrho\mathbf{C}_{1}^{2}(t) + 0.5S\,\Theta\mathfrak{b}\varrho\mathbf{C}_{2}^{2}(t) + S\,\mathfrak{b}\varrho\mathbf{C}_{1}(t) \\ &+ \Theta + \mathfrak{b}\varrho(\zeta+1)^{-1}\mathbf{C}_{1}(t)^{\zeta+1} + \mathfrak{b}\varrho\zeta(\zeta+1)^{-1}\mathbf{C}_{2}^{\zeta+1}(t) - 0.25(1-\zeta)\mathfrak{q}_{11}^{2}\mathbf{C}_{1}^{\zeta+2}(t) \end{split}$$

AIMS Mathematics

$$-0.25(1-\zeta)\mathfrak{q}_{21}^{2}\mathbf{C}_{2}^{\zeta+2}(t) + 0.5\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h}(t) - 0.5(1-\zeta)\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h+\zeta}(t) \\ -S\mathcal{T}_{\circ}^{\Sigma} + \mathcal{U} - 0.25(1-\zeta)\mathfrak{q}_{11}^{2}a^{-\zeta-2} \\ -1.$$

(4) Assume that $(\mathbf{C}_1(t), \mathbf{C}_2(t)) \in \mathcal{J}_{a_{\star},4}$. From (3.7), we get

 \leq

$$\begin{split} \Psi(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t)) &\leq -S\mathcal{T}_{\circ}^{\Sigma} - 0.25(1-\zeta)\mathfrak{q}_{21}^{2}\mathbf{C}_{2}^{\zeta+2}(t) + 0.5S\,\Theta\mathfrak{b}_{\mathcal{Q}}\mathbf{C}_{1}^{2}(t) + 0.5S\,\Theta\mathfrak{b}_{\mathcal{Q}}\mathbf{C}_{2}^{2}(t) + S\,\mathfrak{b}_{\mathcal{Q}}\mathbf{C}_{1}(t) \\ &+ \Theta + \mathfrak{b}_{\mathcal{Q}}(\zeta+1)^{-1}\mathbf{C}_{1}(t)^{\zeta+1} + \mathfrak{b}_{\mathcal{Q}}\zeta(\zeta+1)^{-1}\mathbf{C}_{2}^{\zeta+1}(t) - 0.25(1-\zeta)\mathfrak{q}_{11}^{2}\mathbf{C}_{1}^{\zeta+2}(t) \\ &- 0.25(1-\zeta)\mathfrak{q}_{21}^{2}\mathbf{C}_{2}^{\zeta+2}(t) + 0.5\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h}(t) - 0.5(1-\zeta)\sum_{h=2}^{2N}\mathfrak{p}_{h}\mathbf{C}_{2}^{h+\zeta}(t) \\ &\leq -S\mathcal{T}_{\circ}^{\Sigma} + \mathcal{U} - 0.25(1-\zeta)\mathfrak{q}_{21}^{2}a^{-\zeta-2} \\ &\leq -1. \end{split}$$

In summary, the assertion (3.8) is obtained. On the other hand, we can easily show that $\exists O > 0$ such that $\Psi(\mathbf{C}_1, \mathbf{C}_2) \leq O$, for all $(\mathbf{C}_1, \mathbf{C}_2) \in \mathbb{R}^{2, \star}_+$. Accordingly, we get

$$\begin{split} &\int_{0}^{t} \mathbb{E}(\Psi(\mathbf{C}_{1}(\tau),\mathbf{C}_{2}(\tau))) d\tau + S \varrho \mathbb{E}\left(\int_{0}^{t} \int_{0}^{\infty} \mathcal{H}(y,0) \pi^{\mathbf{D}}(dy) d\tau - \int_{0}^{t} \mathcal{H}(\mathbf{D}(\tau),0) d\tau\right) \\ &\geq \int_{0}^{t} \mathbb{E}\left(\mathcal{L}\Phi_{\Theta,\zeta}^{\Sigma}(t)(\mathbf{C}_{1}(\tau),\mathbf{C}_{2}(\tau))\right) d\tau \\ &= \mathbb{E}\left(\Phi_{\Theta,\zeta}^{\Sigma}(t)(\mathbf{C}_{1}(t),\mathbf{C}_{2}(t))\right) - \mathbb{E}\left(\Phi_{\Theta,\zeta}^{\Sigma}(t)(\mathbf{C}_{1}(0),\mathbf{C}_{2}(0))\right) \\ &\geq - \mathbb{E}\left(\Phi_{\Theta,\zeta}^{\Sigma}(t)(\mathbf{C}_{1}(0),\mathbf{C}_{2}(0))\right). \end{split}$$

By using the ergodic property of $\mathbf{D}(t)$, we conclude that

$$0 \leq \liminf_{t \to \infty} t^{-1} \int_0^t \left(\mathbb{E}\Psi(\mathbf{C}_1(\tau), \mathbf{C}_2(\tau)) \mathbb{1}_{\{(\mathbf{C}_1(\tau), \mathbf{C}_2(\tau)) \in \mathcal{J}_{a,a_\star}^c\}} + \mathbb{E}\Psi(\mathbf{C}_1(\tau), \mathbf{C}_2(\tau)) \mathbb{1}_{\{(\mathbf{C}_1(\tau), \mathbf{C}_2(\tau)) \in \mathcal{J}_{a,a_\star}\}} \right) d\tau$$

$$\leq \liminf_{t \to \infty} t^{-1} \int_0^t \left(-\mathbb{P}((\mathbf{C}_1(\tau), \mathbf{C}_2(\tau)) \in \mathcal{J}_{a,a_\star}^c) + O\mathbb{P}((\mathbf{C}_1(\tau), \mathbf{C}_2(\tau)) \in \mathcal{J}_{\epsilon,\epsilon_\star})) d\tau$$

$$= -1 + (1 + O)\liminf_{t \to \infty} t^{-1} \int_0^t \mathbb{P}((\mathbf{C}_1(\tau), \mathbf{C}_2(\tau)) \in \mathcal{J}_{a,a_\star}) d\tau.$$

Consequently,

$$\liminf_{t\to\infty} t^{-1} \int_0^t \mathbb{P}((\mathbf{C}_1(\tau), \mathbf{C}_2(\tau)) \in \mathcal{Z}_{a,a_\star}) \mathrm{d}\tau \ge (1+O)^{-1} > 0.$$

Hence,

$$\liminf_{t\to\infty} t^{-1} \int_0^t \mathbb{P}((\mathbf{C}_1(0), \mathbf{C}_2(0), \mathbf{C}_3(0)); \tau, \mathcal{J}_{a,a_\star}) ds > 0, \qquad \forall (\mathbf{C}_1(0), \mathbf{C}_2(0), \mathbf{C}_3(0)) \in \mathbb{R}^{3,\star}_+,$$

where $\mathbb{R}^{3,\star}_+ = \{(x, y, z) : x > 0, y > 0, z > 0\}$. From Lemma 3.2 of [42] and also the mutually exclusive possibilities lemma [43], we confirm the existence of a single invariant distribution π^{Σ} .

AIMS Mathematics

Remark 3.1. From Theorem 3.1 of this paper and also Theorem 2.6 of [44], we can deduce interesting indications on the stochastic permanence of the Markovian processes C_1 , C_2 and C_3 . Explicitly, we obtain that

$$\lim_{t \to \infty} t^{-1} \int_0^t \mathbf{C}_1(\tau) d\tau = \int_{\mathbb{R}^{3,\star}_+} c_1 \pi^{\Sigma}(dc_1, dc_2, dc_3) < \infty,$$
$$\lim_{t \to \infty} t^{-1} \int_0^t \mathbf{C}_2(\tau) d\tau = \int_{\mathbb{R}^{3,\star}_+} c_2 \pi^{\Sigma}(dc_1, dc_2, dc_3) < \infty,$$
$$\lim_{t \to \infty} t^{-1} \int_0^t \mathbf{C}_3(\tau) d\tau = \int_{\mathbb{R}^{3,\star}_+} c_3 \pi^{\Sigma}(dc_1, dc_2, dc_3) < \infty.$$

By way of illustration, this indicates the persistence of the infection over time.

4. Numerical verification

In this section, we exhibit some simulations to shed some light on the exactitude of our global threshold. For that purpose, we present three situations of system (1.4), and in each case, we explore the complex long-run behavior of the illness. We will consider general saturated interference function introduced in Table 1. The model parameters are theoretically selected according to well audit the outcomes of this paper. By using the high-order discrete Milstein method, the associated discretization equations of our system are directly obtained as follows:

$$\begin{cases} \mathbf{C}_{1,k+1} = \mathbf{C}_{1,k} + \left[\Theta - (\mathbf{u}_{1} + \mathbf{3})\mathbf{C}_{1,k} - \mathfrak{b}\mathcal{H}(\mathbf{C}_{1,k}, \mathbf{C}_{2,k})\mathbf{C}_{2,k}\right]\Delta t + \sum_{h=0}^{N}\mathfrak{q}_{1h}\mathbf{C}_{1,k}^{h+1}\sqrt{\Delta t}\xi_{k} \\ + \frac{1}{2}\left(\sum_{h=0}^{N}\mathfrak{q}_{1h}\mathbf{C}_{1,k}^{h+1}\right)\left(\sum_{h=0}^{N}\mathfrak{q}_{1h}(h+1)\mathbf{C}_{1,k}^{h}\right)(\xi^{2} - 1)\Delta t, \\ \mathbf{C}_{2,k+1} = \mathbf{C}_{2,k} + \left[\mathfrak{b}\mathcal{H}(\mathbf{C}_{1,k}, \mathbf{C}_{2,k})\mathbf{C}_{2,k} - (\mathbf{u}_{2} + \mathfrak{a} + \mathfrak{c})\mathbf{C}_{2,k}\right]\Delta t + \sum_{h=0}^{N}\mathfrak{q}_{2h}\mathbf{C}_{2,k}^{h+1}\sqrt{\Delta t}\xi_{k} \\ + \frac{1}{2}\left(\sum_{h=0}^{N}\mathfrak{q}_{2h}\mathbf{C}_{2,k}^{h+1}\right)\left(\sum_{h=0}^{N}\mathfrak{q}_{2h}(h+1)\mathbf{C}_{2,k}^{h}\right)(\zeta^{2} - 1)\Delta t, \\ \mathbf{C}_{3,k+1} = \mathbf{C}_{3,k} + \left[\mathbf{3}\mathbf{C}_{1,k} + \mathfrak{c}\mathbf{C}_{2,k} - \mathfrak{u}_{3}\mathbf{C}_{3,k}\right]\Delta t + \sum_{h=0}^{N}\mathfrak{q}_{3h}\mathbf{C}_{3,k}^{h+1}\sqrt{\Delta t}\alpha_{k} \\ + \frac{1}{2}\left(\sum_{h=0}^{N}\mathfrak{q}_{3h}\mathbf{C}_{3,k}^{h+1}\right)\left(\sum_{h=0}^{N}\mathfrak{q}_{3h}(h+1)\mathbf{C}_{3,k}^{h}\right)(\alpha^{2} - 1)\Delta t, \end{cases}$$

where the time increment $\Delta t > 0$. ξ_k , ζ_k and α_k are three independent random variables which follow the Gaussian distribution $\mathcal{N}(0, 1)$. ($\mathbf{C}_{1,k}, \mathbf{C}_{2,k}, \mathbf{C}_{3,k}$) is the corresponding value of the *k*-th iteration.

4.1. Linear disturbance case (N = 0)

In this example, we will deal with the following stochastic system:

$$\begin{cases} d\mathbf{C}_{1}(t) = \left(\Theta - (\mathfrak{u}_{1} + \mathfrak{z})\mathbf{C}_{1}(t) - \frac{\mathbf{b}\mathbf{C}_{1}(t)\mathbf{C}_{2}(t)}{1 + \mathfrak{m}_{1}\mathbf{C}_{1}(t) + \mathfrak{m}_{2}\mathbf{C}_{2}(t)}\right) dt + \mathfrak{q}_{10}\mathbf{C}_{1}(t)d\mathbb{W}_{1}(t), \\ d\mathbf{C}_{2}(t) = \left(\frac{\mathbf{b}\mathbf{C}_{1}(t)\mathbf{C}_{2}(t)}{1 + \mathfrak{m}_{1}\mathbf{C}_{1}(t) + \mathfrak{m}_{2}\mathbf{C}_{2}(t)} - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c})\mathbf{C}_{2}(t)\right) dt + \mathfrak{q}_{20}\mathbf{C}_{2}(t)d\mathbb{W}_{2}(t), \\ d\mathbf{C}_{3}(t) = \left(\mathfrak{z}\mathbf{C}_{1}(t) + \mathfrak{c}\mathbf{C}_{2}(t) - \mathfrak{u}_{3}\mathbf{C}_{3}(t)\right) dt + \mathfrak{q}_{30}\mathbf{C}_{3}(t)d\mathbb{W}_{3}(t), \\ \mathbf{C}_{1}(0) = 0.5, \ \mathbf{C}_{2}(0) = 0.3, \ \mathbf{C}_{3}(0) = 0.2, \end{cases}$$
(4.1)

AIMS Mathematics

associated with the following auxiliary process:

$$d\mathbf{D}(t) = \left(\Theta - (\mathfrak{u}_1 + \mathfrak{z})\mathbf{D}(t)\right)dt + \mathfrak{q}_{10}\mathbf{D}(t)d\mathbb{W}_1(t), \quad \mathbf{D}(0) = 0.5$$

We choose $\Theta = 0.23$, $u_1 = 0.2$, $u_3 = 0.19$, $\mathfrak{z} = 0.02$, $u_2 = 0.2$, $\mathfrak{m}_1 = 0.1$, $\mathfrak{m}_2 = 0.1$, $\mathfrak{a} = 0.2$, $\mathfrak{c} = 0.02$, $\mathfrak{q}_{10} = 0.11$, $\mathfrak{q}_{20} = 0.112$ and $\mathfrak{q}_{30} = 0.1$. By setting $\mathfrak{b} = 0.4$ and considering a large time *T*, we get

$$\mathcal{T}_{\circ}^{\Sigma} = \lim_{T \to \infty} T^{-1} \int_{0}^{T} \frac{\mathfrak{b} \mathbf{D}(\tau)}{1 + m_{1} \mathbf{D}(\tau)} \mathrm{d}\tau - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) - 0.5\mathfrak{q}_{20}^{2} \cong -0.0478 < 0.$$

From Theorem 2.1, we can infer the disappearance of the illness. Numerically, it can be seen in Figure 1 that the disease disappears after 230 days in the population with a strong permanence of classes 1 and 3. Now, we select b = 0.55 to switch from the case of extinction to the case of persistence. Then,

 $\mathcal{T}_{\circ}^{\Sigma} = \lim_{T \to \infty} T^{-1} \int_{0}^{T} \frac{\mathbf{b} \mathbf{D}(\tau)}{1 + m_{1} \mathbf{D}(\tau)} d\tau - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) - 0.5\mathfrak{q}_{20}^{2} \cong 0.0879 > 0.$



Figure 1. Computer illustration of the trajectories of the probabilistic model (4.1) with linear white noise.

In accordance with Theorem 3.1, we infer that the properties of stationarity and ergodicity hold. From Figure 2, we offer a good illustration of these two statistical characteristics. Clearly, in this situation, the continuation of all classes is strongly happening which is depicted in Figure 3.



Figure 2. The 3*D* graphs and associated upper views of the joint probability density at time t = 600 of the classes C_1 , C_2 and C_3 .



Figure 3. Computer simulation of the solution of the stochastic model (4.1) with linear white noises.

4.2. Quadratic disturbance case (N = 1)

In this example, we deal with the following stochastic system:

$$\begin{cases} d\mathbf{C}_{1}(t) = \left(\Theta - (u_{1} + 3)\mathbf{C}_{1}(t) - \frac{b\mathbf{C}_{1}(t)\mathbf{C}_{2}(t)}{1 + w_{1}\mathbf{C}_{1}(t) + w_{2}\mathbf{C}_{2}(t)}\right) dt + \mathfrak{q}_{10}\mathbf{C}_{1}(t)d\mathbb{W}_{1}(t) + \mathfrak{q}_{11}\mathbf{C}_{1}^{2}(t)d\mathbb{W}_{1}(t), \\ d\mathbf{C}_{2}(t) = \left(\frac{b\mathbf{C}_{1}(t)\mathbf{C}_{2}(t)}{1 + w_{1}\mathbf{C}_{1}(t) + w_{2}\mathbf{C}_{2}(t)} - (u_{2} + \mathfrak{a} + \mathfrak{c})\mathbf{C}_{2}(t)\right) dt + \mathfrak{q}_{20}\mathbf{C}_{2}(t)d\mathbb{W}_{2}(t) + \mathfrak{q}_{21}\mathbf{C}_{2}^{2}(t)d\mathbb{W}_{2}(t), \\ d\mathbf{C}_{3}(t) = \left(3\mathbf{C}_{1}(t) + \mathfrak{c}\mathbf{C}_{2}(t) - u_{3}\mathbf{C}_{3}(t)\right) dt + \mathfrak{q}_{30}\mathbf{C}_{3}(t)d\mathbb{W}_{3}(t) + \mathfrak{q}_{31}\mathbf{C}_{3}^{2}(t)d\mathbb{W}_{3}(t), \\ \mathbf{C}_{1}(0) = 0.5, \ \mathbf{C}_{2}(0) = 0.3, \ \mathbf{C}_{3}(0) = 0.2, \end{cases}$$

$$(4.2)$$

associated with the following auxiliary process:

$$d\mathbf{D}(t) = \left(\Theta - (\mathfrak{u}_1 + \mathfrak{z})\mathbf{D}(t)\right)dt + \mathfrak{q}_{10}\mathbf{D}(t)d\mathbb{W}_1(t) + \mathfrak{q}_{11}\mathbf{D}^2(t)d\mathbb{W}_1(t), \quad \mathbf{D}(0) = 0.5.$$

For the comparison objective, we keep the same parameter values as the first case and we select $q_{11} = 0.022$, $q_{21} = 0.013$ and $q_{31} = 0.011$. As the above case, we select b = 0.4. Then, we get

$$\mathcal{T}_{\circ}^{\Sigma} = \lim_{T \to \infty} T^{-1} \int_{0}^{T} \frac{\mathbf{b} \mathbf{D}(\tau)}{1 + m_1 \mathbf{D}(\tau)} \mathrm{d}\tau - (\mathfrak{u}_2 + \mathfrak{a} + \mathfrak{c}) - 0.5\mathfrak{q}_{20}^2 \cong -0.0520 < 0.$$

AIMS Mathematics

Apparently, the condition of Theorem 2.1 holds. Numerically, the extinction phenomenon of the illness is illustrated in Figure 4. Classes 1 and 3 still persist. Now, we choose b = 0.55 to move from the case of extinction to the case of persistence. Then,



$$\mathcal{T}_{\circ}^{\Sigma} = \lim_{T \to \infty} T^{-1} \int_{0}^{T} \frac{\mathfrak{b} \mathbf{D}(\tau)}{1 + m_{1} \mathbf{D}(\tau)} \mathrm{d}\tau - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) - 0.5\mathfrak{q}_{20}^{2} \cong 0.0896 > 0.$$

Figure 4. Computer simulation of the trajectories of the probabilistic model (4.2) with quadratic white noises.

Based on Theorem 3.1, The last value implies that the system (4.2) admits a stable distribution. From Figure 5, we get an insight into the stationarity of the model (4.2). Furthermore, we offer Figure 6 to clarify the continuation of all classes C_1 , C_2 and C_3 .



Figure 5. The 3D graphs and associated upper views of the joint probability density at time t = 600 of the classes C_1 , C_2 and C_3 .



Figure 6. Computer illustration of the solution of the probabilistic model (4.2) with quadratic white noises.

4.3. Cubic disturbance case (N = 2)

In this part, we numerically prove that $\mathcal{T}_{\circ}^{\Sigma}$ is the sill of the system (1.4) in the special case of cubic perturbation. So, we firstly introduce this probabilistic model:

$$\begin{cases} d\mathbf{C}_{1}(t) = \left(\Theta - (u_{1} + 3)\mathbf{C}_{1}(t) - \frac{b\mathbf{C}_{1}(t)\mathbf{C}_{2}(t)}{1 + m_{1}\mathbf{C}_{1}(t) + m_{2}\mathbf{C}_{2}(t)}\right) dt + q_{10}\mathbf{C}_{1}(t)d\mathbb{W}_{1}(t) \\ + q_{11}\mathbf{C}_{1}^{2}(t)d\mathbb{W}_{1}(t) + q_{12}\mathbf{C}_{1}^{3}(t)d\mathbb{W}_{1}(t), \\ d\mathbf{C}_{2}(t) = \left(\frac{b\mathbf{C}_{1}(t)\mathbf{C}_{2}(t)}{1 + m_{1}\mathbf{C}_{1}(t) + m_{2}\mathbf{C}_{2}(t)} - (u_{2} + a + c)\mathbf{C}_{2}(t)\right) dt + q_{20}\mathbf{C}_{2}(t)d\mathbb{W}_{2}(t) \\ + q_{21}\mathbf{C}_{2}^{2}(t)d\mathbb{W}_{2}(t) + q_{22}\mathbf{C}_{2}^{3}(t)d\mathbb{W}_{2}(t), \\ d\mathbf{C}_{3}(t) = \left(3\mathbf{C}_{1}(t) + c\mathbf{C}_{2}(t) - u_{3}\mathbf{C}_{3}(t)\right) dt + q_{30}\mathbf{C}_{3}(t)d\mathbb{W}_{3}(t) + q_{31}\mathbf{C}_{3}^{2}(t)d\mathbb{W}_{3}(t) + q_{32}\mathbf{C}_{3}^{3}(t)d\mathbb{W}_{3}(t), \\ \mathbf{C}_{1}(0) = 0.5, \ \mathbf{C}_{2}(0) = 0.3, \ \mathbf{C}_{3}(0) = 0.2, \end{cases}$$

$$(4.3)$$

associated with the following auxiliary process:

$$\mathbf{d}\mathbf{D}(t) = \left(\Theta - (\mathfrak{u}_1 + \mathfrak{z})\mathbf{D}(t)\right)\mathbf{d}t + \mathfrak{q}_{10}\mathbf{D}(t)\mathbf{d}\mathbb{W}_1(t) + \mathfrak{q}_{11}\mathbf{D}^2(t)\mathbf{d}\mathbb{W}_1(t) + \mathfrak{q}_{12}\mathbf{D}^3(t)\mathbf{d}\mathbb{W}_1(t), \quad \mathbf{D}(0) = 0.5.$$

AIMS Mathematics

Here, we select $q_{12} = 0.014$, $q_{22} = 0.0135$, $q_{32} = 0.012$ and we keep the other coefficient values as the above two cases. Again, if we select b = 0.4, then the result is

$$\mathcal{T}_{\circ}^{\Sigma} = \lim_{T \to \infty} T^{-1} \int_{0}^{T} \frac{\mathbf{b} \mathbf{D}(\tau)}{1 + m_{1} \mathbf{D}(\tau)} \mathrm{d}\tau - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) - 0.5\mathfrak{q}_{20}^{2} \cong -0.0616 < 0.$$

Theoretically, we have the disappearance of the illness according to Theorem 2.1. It remains to verify it numerically. From Figure 7, the disease will clear up in about 70 days with long-term persistence of categories 1 and 3. Now, we opt b = 0.55. Then,

$$\mathcal{T}_{\circ}^{\Sigma} = \lim_{T \to \infty} T^{-1} \int_{0}^{T} \frac{\mathfrak{b} \mathbf{D}(\tau)}{1 + m_{1} \mathbf{D}(\tau)} \mathrm{d}\tau - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) - 0.5\mathfrak{q}_{20}^{2} \cong 0.0905 > 0.$$



Figure 7. Trajectories of the probabilistic model (4.3) with cubic white noises.

From Theorem 3.1, we establish that there is a single stable distribution for (4.3) which is depicted in Figure 8. The persistence of all classes is depicted in Figure 9.



Figure 8. The 3*D* graphs and associated upper views of the joint probability density at time t = 600 of the classes C_1 , C_2 and C_3 .



Figure 9. Computer simulation of the solution of the probabilistic model (4.3) with cubic white noises.

5. Conclusions

Mathematical formulation plays a major role in understanding epidemics and also in supporting public health decision-making. The regularly used models provide deterministic predictions, that is to say, a strict behavior of the system studied, thus ignoring individual and environmental variations. Actually, we group these unpredictable variations under the name of stochasticity (or randomness), and the present study is devoted to the analysis of an epidemic strewing under heavy stochasticity. The non-linearity and the complexity of the fluctuations pushed us to consider a general form of the probabilistic part. Focusing on these motivations, we have offered an improved generalization of the recent paper [9]. In the following, we present our substantial enhancements of the mentioned research.

- ★ In [9], the authors considered a nonlinear prevalence function of the form: $bC_1(t)g(C_2(t))$. This type of function has certain limitations and is not suitable for covering certain well-known functional responses. By assuming mutual interference between classes C_1 and C_2 , we have proposed a general function \mathcal{H} which includes all the existing functional incidences.
- ★ In [9], the authors presented two distinct criteria to classify the asymptotic attitude of system (1.4) with the incidence function $bC_1(t)g(C_2(t))$. Explicitly, they obtained the following results:

(1) The sufficient condition of the disease disappearance is

$$\mathcal{R}_{0}^{C} = \mathfrak{b}g'(0) \int_{\mathbb{R}_{+}} y\pi^{\mathbf{D}}(\mathrm{d}y) - \left(\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c} + 0.5\mathfrak{q}_{20}^{2}\right) < 0.$$

(2) The sufficient condition of the ergodicity is

$$\mathcal{R}_{0}^{S} = \Theta \mathfrak{b}g'(0) - \left(\mathfrak{u}_{1} + \mathfrak{z} + 0.5\mathfrak{q}_{10}^{2} + \sum_{h=1}^{2N} \sqrt[h+1]{2^{2h-1}\sum_{i+j=h}\mathfrak{q}_{1i}\mathfrak{q}_{1j}\Theta^{h}}\right) \left(\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c} + 0.5\mathfrak{q}_{20}^{2}\right) > 0.$$

However, in this article, we have unified the criterion of the above-mentioned characteristics by providing the following acute threshold value:

$$\mathcal{T}_{\circ}^{\Sigma} = \mathfrak{b} \int_{\mathbb{R}_{+}} \mathcal{H}(y,0) \pi^{\mathbf{D}}(\mathrm{d}y) - (\mathfrak{u}_{2} + \mathfrak{a} + \mathfrak{c}) - 0.5\mathfrak{q}_{20}^{2}.$$

Specifically,

(1) The condition of the disease disappearance is $\mathcal{T}^{\Sigma}_{\circ} < 0$.

(2) The condition of the ergodicity is $\mathcal{T}_{\circ}^{\Sigma} > 0$.

As a special case, we mention that the sharp threshold of the perturbed model studied in [9] is exactly \mathcal{R}_0^C .

Numerically, we have chosen the first three values of h, that are, linear, quadratic and cubic perturbations. In all cases, we have confirmed that $\mathcal{T}_{\circ}^{\Sigma}$ is the global sill among disappearance of infection and ergodicity. From Figure 10, we infer that when we raise the perturbation order, the illness disappears swiftly. This indicates that the intense environmental variations have a negative effect on the illness duration. This remark requires further theoretical clarification and explanation. We will deal with this interesting question in our future work.

Some fascinating topics deserve more attention. For example, we can consider our model with fractal-fractional differentiation. This framework is an attractive branch of applied mathematics that deals with derivatives and integrals of non-integer order. Due to its amazing features, it is preferred for describing and simulating real-world problems in various fields such as biological mechanisms, material science, hydrological modeling, economic phenomena. We will address this idea in our future work.



Figure 10. Stochastic paths of infected class C_2 in the case of N = 3 ($q_{13} = 0.0116$, $q_{23} = 0.012$, $q_{33} = 0.0108$); in the case of N = 4 ($q_{14} = 0.016$, $q_{24} = 0.024$, $q_{34} = 0.038$) and in the case of N = 5 ($q_{15} = 0.06$, $q_{25} = 0.08$, $q_{35} = 0.017$). The other coefficients are selected above.

Acknowledgements

This research was sponsored by the Guangzhou Government Project under Grant No. 62216235, also supported by SERB of India (EEQ/2021/001003).

Conflict of interest

The authors declare no conflicts of interest.

References

1. Z. C. Wang, K. Tang, Combating COVID-19: health equity matters, *Nat. Med.*, **26** (2020), 458. https://doi.org/10.1038/s41591-020-0823-6

- S. Djilali, L. Benahmadi, A. Tridane, K. Niri, Modeling the impact of unreported cases of the COVID-19 in the North African countries, *Biology*, 9 (2020), 1–18. https://doi.org/10.3390/biology9110373
- 3. S. Bentout, A. Tridane, S. Djilali, T. M. Touaoula, Age-structured modeling of COVID-19 epidemic in the USA, UAE and Algeria, *Alex. Eng. J.*, **60** (2021), 401–411.
- M. Abdy, S. Side, S. Annas, W. Nur, W. Sanusi, An SIR epidemic model for COVID-19 spread with fuzzy parameter: the case of Indonesia, *Adv. Differ. Equ.*, 2021 (2021), 1–17. https://doi.org/10.1186/s13662-021-03263-6
- 5. P. Brodin, Immune determinants of COVID-19 disease presentation and severity, *Nat. Med.*, **27** (2021), 28–33. https://doi.org/10.1038/s41591-020-01202-8
- W. J. Li, J. C. Ji, L. H. Huang, Z. Y. Guo, Global dynamics of a controlled discontinuous diffusive SIR epidemic system, *Appl. Math. Lett.*, **121** (2021), 107420. https://doi.org/10.1016/j.aml.2021.107420
- W. O. Kermack, A. G. McKendrick, A contribution to the mathematical theory of epidemics, *Proc. R. Soc. A*, **115** (1927), 700–721. https://doi.org/10.1098/rspa.1927.0118
- 8. N. A. Kudryashov, M. A. Chmykhov, M. Vigdorowitsch, Analytical features of the SIR model and their applications to COVID-19, *Appl. Math. Model.*, **90** (2021), 466–473. https://doi.org/10.1016/j.apm.2020.08.057
- B. Q. Zhou, B. T. Han, D. Q. Jiang, Ergodic property, extinction and density function of a stochastic SIR epidemic model with nonlinear incidence and general stochastic perturbations, *Chaos Solitons Fract.*, **152** (2021), 111338. https://doi.org/10.1016/j.chaos.2021.111338
- 10. J. J. Wang, J. Z. Zhang, Z. Jin, Analysis of an SIR model with bilinear incidence rate, *Nonlinear Anal.*, **11** (2010), 2390–2402. https://doi.org/10.1016/j.nonrwa.2009.07.012
- 11. Y. G. Lin, D. Q. Jiang, M. L. Jin, Stationary distribution of a stochastic SIR model with saturated incidence and its asymptotic stability, *Acta Math. Sci.*, **35** (2015), 619–629. https://doi.org/10.1016/S0252-9602(15)30008-4
- C. J. Sun, W. Yang, J. Arino, K. Khan, Effect of media-induced social distancing on disease transmission in a two patch setting, *Math. Biosci.*, 230 (2011), 87–95. https://doi.org/10.1016/j.mbs.2011.01.005
- D. Kiouach, Y. Sabbar, Stability and threshold of a stochastic SIRS epidemic model with vertical transmission and transfer from infectious to susceptible individuals, *Discrete Dyn. Nat. Soc.*, 2018 (2018), 1–13. https://doi.org/10.1155/2018/7570296
- 14. N. H. Du, N. N. Nhu, Permanence and extinction of certain stochastic SIR models perturbed by a complex type of noises, *Appl. Math. Lett.*, **64** (2017), 223–230. https://doi.org/10.1016/j.aml.2016.09.012
- A. Kumar, Nilam, Dynamic behavior of an SIR epidemic model along with time delay; Crowley-Martin type incidence rate and Holling type II treatment rate, *Int. J. Nonlinear Sci. Numer. Simul.*, 20 (2019), 757–771. https://doi.org/10.1515/ijnsns-2018-0208
- 16. M. J. Faddy, Nonlinear stochastic compartmental models, *Math. Med. Biol.*, **2** (1985), 287–297. https://doi.org/10.1093/imammb/2.4.287

- Y. Sabbar, A. Din, D. Kiouach, Predicting potential scenarios for wastewater treatment under unstable physical and chemical laboratory conditions: a mathematical study, *Results Phys.*, 39 (2022), 105717. https://doi.org/10.1016/j.rinp.2022.105717
- Y. Sabbar, A. Zeb, D. Kiouach, N. Gul, T. Sitthiwirattham, D. Baleanu, et al., Dynamical bifurcation of a sewage treatment model with general higher-order perturbation, *Results Phys.*, **39** (2022), 105799. https://doi.org/10.1016/j.rinp.2022.105799
- 19. Y. Sabbar, A. Khan, A. Din, Probabilistic analysis of a marine ecological system with intense variability, *Mathematics*, **10** (2022), 1–19. https://doi.org/10.3390/math10132262
- 20. Y. Sabbar, D. Kiouach, New method to obtain the acute sill of an ecological model with complex polynomial perturbation, *Math. Methods Appl. Sci.*, 2022. https://doi.org/10.1002/mma.8654
- S. Ditlevsen, A. Samson, Introduction to stochastic models in biology, In: *Stochastic biomathematical models*, Berlin, Heidelber: Springer, 2013, 3–35. https://doi.org/10.1007/978-3-642-32157-3_1
- S. P. Rajasekar, M. Pitchaimani, Q. X. Zhu, Dynamic threshold probe of stochastic SIR model with saturated incidence rate and saturated treatment function, *Phys. A*, **535** (2019), 122300. https://doi.org/10.1016/j.physa.2019.122300
- 23. D. Kiouach, Y. Sabbar, The threshold of a stochastic SIQR epidemic model with Lévy jumps, In: *Trends in biomathematics: mathematical modeling for health, harvesting, and population dynamics*, Cham: Springer, 2019, 87–105. https://doi.org/10.1007/978-3-030-23433-1_7
- 24. S. Winkelmann, C. Schütte, *Stochastic dynamics in computational biology*, Cham: Springer, 2020. https://doi.org/10.1007/978-3-030-62387-6
- 25. S. P. Rajasekar, M. Pitchaimani, Q. X. Zhu, K. B. Shi, Exploring the stochastic host-pathogen tuberculosis model with adaptive immune response, *Math. Probl. Eng.*, **2021** (2021), 1–23. https://doi.org/10.1155/2021/8879538
- 26. S. P. Rajasekar, M. Pitchaimani, Q. X. Zhu, Probing a stochastic epidemic hepatitis C virus model with a chronically infected treated population, *Acta Math. Sci.*, 42 (2022), 2087–2112. https://doi.org/10.1007/s10473-022-0521-1
- 27. N. S. Goel, N. Richter-Dyn, Stochastic models in biology, Academic Press, 1974.
- 28. D. Kiouach, Y. Sabbar, S. E. A. El-idrissi, New results on the asymptotic behavior of an SIS epidemiological model with quarantine strategy, stochastic transmission, and Lévy disturbance, *Math. Methods Appl. Sci.*, 44 (2021), 13468–13492. https://doi.org/10.1002/mma.7638
- 29. D. Kiouach, Y. Sabbar, Developing new techniques for obtaining the threshold of a stochastic SIR epidemic model with 3-dimensional Levy process, *J. Appl. Nonlinear Dyn.*, **11** (2022), 401–414. https://doi.org/10.5890/JAND.2022.06.010
- 30. Y. Sabbar, D. Kiouach, S. P. Rajasekar, S. E. A. El-idrissi, The influence of quadratic Lévy noise on the dynamic of an SIC contagious illness model: new framework, critical comparison and an application to COVID-19 (SARS-CoV-2) case, *Chaos Solitons Fract.*, **159** (2022), 112110. https://doi.org/10.1016/j.chaos.2022.112110
- D. Kiouach, Y. Sabbar, Threshold analysis of the stochastic Hepatitis B epidemic model with successful vaccination and Lévy jumps, In: 2019 4th World Conference on Complex Systems (WCCS), 2019, 1–6. https://doi.org/10.1109/ICoCS.2019.8930709

- 32. S. P. Rajasekar, M. Pitchaimani, Q. X. Zhu, Higher order stochastically perturbed SIRS epidemic model with relapse and media impact, *Math. Methods Appl. Sci.*, **45** (2022), 843–863. https://doi.org/10.1002/mma.7817
- 33. D. Kiouach, Y. Sabbar, The long-time behavior of a stochastic SIR epidemic model with distributed delay and multidimensional Lévy jumps, *Int. J. Biomath.*, **15** (2021), 2250004. https://doi.org/10.1142/S1793524522500048
- 34. D. Kiouach, Y. Sabbar, Dynamic characterization of a stochastic SIR infectious Biomath., model with dual perturbation, Int. J. 14 (2021),2150016. disease https://doi.org/10.1142/S1793524521500169
- D. Kiouach, Y. Sabbar, Ergodic stationary distribution of a stochastic Hepatitis B epidemic model with interval-valued parameters and compensated Poisson process, *Comput. Math. Methods Med.*, 2020 (2020), 1–12. https://doi.org/10.1155/2020/9676501
- Q. Liu, D. Jiang, Stationary distribution and extinction of a stochastic SIR modelwith nonlinear perturbation, *Appl. Math. Lett.*, 73 (2017), 8–15. https://doi.org/10.1016/j.aml.2017.04.021
- 37. Q. Liu, D. Q. Jiang, Dynamical behavior of a higher order stochastically perturbed HIV/AIDS model with differential infectivity and amelioration, *Chaos Solitons Fract.*, **141** (2020), 110333. https://doi.org/10.1016/j.chaos.2020.110333
- 38. B. T. Han, D. Q. Jiang, T. Hayat, A. Alsaedi, B. Ahmed, Stationary distribution and extinction of a stochastic staged progression AIDS model with staged treatment and second-order perturbation, *Chaos Solitons Fract.*, **140** (2020), 110238. https://doi.org/10.1016/j.chaos.2020.110238
- 39. Y. Sabbar, D. Kiouach, S. P. Rajasekar, Acute threshold dynamics of an epidemic system with quarantine strategy driven by correlated white noises and Lévy jumps associated with infinite measure, *Int. J. Dyn. Control*, 2022. https://doi.org/10.1007/s40435-022-00981-x
- 40. N. Ikeda, S. Watanabe, A comparison theorem for solutions of stochastic differential equations and its applications, *Osaka J. Math.*, **14** (1977), 619–633.
- 41. X. R. Mao, Stochastic differential equations and applications, Elsevier, 2007.
- 42. J. Y. Tong, Z. Z. Zhang, J. H. Bao, The stationary distribution of the facultative population model with a degenerate noise, *Stat. Probabil. Lett.*, **83** (2013), 655–664. https://doi.org/10.1016/j.spl.2012.11.003
- 43. D. L. Zhao, S. L. Yuan, Sharp conditions for the existence of a stationary distribution in one classical stochastic chemostat, *Appl. Math. Comput.*, **339** (2018), 199–205. https://doi.org/10.1016/j.amc.2018.07.020
- 44. N. T. Dieu, V. H. Sam, N. H. Du, Threshold of a stochastic SIQS epidemic model with isolation, *Discrete Cont. Dyn. Syst. B*, **27** (2022), 5009–5028. https://doi.org/10.3934/dcdsb.2021262



© 2022 the Author(s), licensee AIMS Press. This is an open access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0)