

1 **STABILITY OF A NONAUTONOMOUS DELAYED PERIODIC**
2 **REACTION-DIFFUSION PREDATOR-PREY MODEL**

3 LILI JIA AND CHANGYOU WANG

ABSTRACT. This article focuses on the stability of periodic solution of a delayed nonautonomous reaction-diffusion predator-prey model. The fine combination of upper and lower solution methods and Lyapunov stability theory is used to transform the study of the stability problem of delayed reaction-diffusion equations into the stability problem of their corresponding delayed ordinary differential equations. Some sufficient conditions are given to ensure the globally asymptotically stability of the periodic solution for this model. Unlike existing results, the stable solution which are obtained in this article is a time-periodic solution rather than a constant periodic solution or a solution for a steady-state system. We extend a stability theorem on predator-prey model introduced by V. Ortega and C. Rebelo in 2023 to nonautonomous delayed reaction-diffusion model. Finally, in order to show the application of the theoretical results, the proposed conditions are numerically validated over a 2-periodic delayed nonautonomous reaction-diffusion predator-prey model.

4 1. INTRODUCTION

5 The study of the properties of reaction-diffusion equations (RDEs) can be used to describe
6 many phenomena in the population and epidemic system, which has attracted increasing attention
7 from scholars [1–7]. Especially, the research on predator-prey RDEs has recently achieved many
8 excellent results. For example, in 2013, Ko and Ahn [8] studied RDEs with two competing pred-
9 ators and one prey and obtained some sufficient conditions to ensure the persistence and global
10 attractiveness for solutions of the system. In 2015, Yang et al. [9] studied RDEs with Leslie-Gower
11 functional response and gained sufficient conditions to guarantee the coexistence state and attrac-
12 tor existence of the model by using fixed point index theory. In 2017, Wang [10] studied RDEs with
13 Neumann boundary conditions and Holling Type III functional response and obtained sufficient
14 conditions to guarantee the existence of periodic orbits by using coincidence theory and bifurcation
15 methods. In 2020, Wu and Zhao [11] studied RDEs with the Allee effect and threshold hunting and
16 analyzed the asymptotic stability of the equilibrium point of the model by constructing a Jacobian
17 matrix. In 2021, Bentout et al. [12] studied an age-structured predator-prey infection model, and
18 explored the impact of predator maturity on interspecies interactions and infectious disease spread.
19 The findings reveal that the minimum maturation time of predators can influence the behavior
20 of the system’s solutions. In the same year, Djilali and Cattani [13] analyzed a superdiffusive
21 predator-prey system with a hunting cooperation functional response. The presence of superdiffu-
22 sion represents the fear effect of the prey and the organized hunting strategy of the predator. The
23 study indicates that superdiffusion leads to complex dynamical behaviors of the system’s solutions
24 and can influence the stability of certain equilibria. In 2022, Yan and Zhang [14] studied RDEs
25 with a Beddington-DeAngelis functional response and obtained stability and instability criteria
26 for the positive constant equilibrium point of the model. In 2023, Chen and Wu [15] studied the
27 spatiotemporal behavior of RDEs with a Beddington-DeAngelis functional response function by
28 using the Leray-Schauder degree theory and Poincare inequality. It is worth mentioning that the
29 above models are autonomous RDEs. Due to the fact that the birth rate, the death rate and the
30 interaction between population are not invariable, nonautonomous RDEs can better simulate the
31 interactions among species in predator-prey models. However, the methods used in the previous
32 literature are difficult to study multi-species nonautonomous predator-prey RDEs. More recently,
33 Jia et al. [16] has considered a 3-species nonautonomous predator-prey RDEs and has obtained

Date: November 30, 2024.

Key words and phrases. Reaction-diffusion predators-prey model; nonautonomous model; periodic solution; stability and permanent property; method of upper and lower solutions.

2010 Mathematics Subject Classification. 35B10, 35B35, 34K20, 92B05.

34 some judgment criteria to ensure the globally asymptotically stability of strictly positive homoge-
 35 nous periodic solution for the system by using the upper and lower solutions method and Lyapunov
 36 stability theory.

37 In a large amount of the real world, the state of a system is influenced not only by its current
 38 state, but also often by their past state. Even in some phenomena, if you ignore the impact of
 39 the system's past state on its future state, the entire research is meaningless. Therefore, when
 40 describe the impact of the interaction between time delayed feedback and spatial transfer on the
 41 system state, scientists have proposed a new type of mathematical model-delayed reaction-diffusion
 42 equations (DRDEs). Using these equations, many real natural phenomena are described and well
 43 explained. In recent years, the research on DRDEs has attracted more and more attention of
 44 scholars. Early research on DRDEs was mostly included in academic works [17,18]. In recent years,
 45 some excellent achievements have been achieved in the study of periodic solutions to DRDEs. For
 46 example, in 2016, Chen and Yu [19] considered a DRDEs with nonlocal delay effect and Dirichlet
 47 boundary condition and obtained stability criteria for the positive equilibrium point of the model.
 48 In 2017, Shi et al. [20] studied a DRDEs with distributed delay and Dirichlet boundary condition
 49 and obtained stability conditions of the positive steady state for the model. In 2018, Yuan and
 50 Guo [21] studied a nonlocal DRDE and achieved the the existence and stability of solutions for
 51 the model with the help of monotone iteration methods. In 2019, Shen and Wei [22] studied a
 52 mussel-algae DRDEs with Neumann boundary conditions and obtained the stability conditions for
 53 the positive constant steady state. In 2021, Zuo and Shi [23] researched a general DRDEs with
 54 spatiotemporal nonlocal delay effect and Dirichlet boundary conditions and obtained some criteria
 55 to ensure the existence and stability of positive steady-state solutions for the system. In 2022,
 56 Xu et al. [24] analyzed a general DRDEs with predator maturation delay and obtained global
 57 asymptotic stability of the positive constant steady state. In 2023, Yuan and Guo [25] studied
 58 a class of DRDEs with spatial nonlocality and achieved some criteria to ensure the stability of
 59 positive steady-state solutions. In the same year, Djilali et al. [26] studied a class of spatially
 60 heterogeneous DRDEs. By employing the Kuratowski measure of noncompactness, the existence
 61 of a global compact attractor for the system is demonstrated. Furthermore, sufficient conditions
 62 for the uniform persistence of solutions and the asymptotic stability of equilibrium solutions are
 63 obtained. In 2024, Kumar [27] investigated a prey-predator DRDEs with Leslie-Gower functional
 64 response and Smith growth functions and established sufficient conditions of the global stability
 65 for the positive constant steady state. It is worth mentioning that the issues studied in the
 66 above literatures are the stability of constant equilibrium solutions or steady-state solutions of
 67 autonomous DRDEs. Moreover, the research methods in the previous papers, such as eigenvalues,
 68 which cannot be used to study nonautonomous DRDEs. To the best of our knowledge, the results
 69 about the stability of periodic solution to nonautonomous DRDEs rarely occurred.

70 Due to the fact that nonautonomous DRDEs can better simulate the interactions between species
 71 in predator-prey models and the study of its dynamic properties has very important practical
 72 significance. In this article, we focus on the following nonautonomous periodic DRDEs

$$\begin{cases} \partial u_1(x,t)/\partial t - d_1(t)\Delta u_1(x,t) = u_1(x,t)[r_1(t) - a_{11}(t)u_1(x,t - \tau_1) - a_{12}(t)u_2(x,t)], \\ \partial u_2(x,t)/\partial t - d_2(t)\Delta u_2(x,t) = u_2(x,t)[-r_2(t) - a_{22}(t)u_2(x,t - \tau_2) + a_{21}(t)u_1(x,t - \tau_1)], \end{cases} \quad (1.1)$$

73 with the Neumann boundary and initial conditions

$$\partial u_i(x,t)/\partial n = 0, (x,t) \in \partial\Omega \times \mathbb{R}_+, u_i(x,t) = \eta_{i0}(x,t) > 0, (x,t) \in \Omega \times [-\tau, 0], i = 1, 2. \quad (1.2)$$

74 Here Ω is a bounded smooth domain in \mathbb{R}^n with boundary $\partial\Omega$, Δ is a Laplace operator on Ω ,
 75 $\partial/\partial n$ denotes the outward normal derivation on $\partial\Omega$, $u_i(x,t)$ represents the density of i -th species
 76 at location $x = (x_1, x_2, \dots, x_n)$ and the time t . τ_1 and τ_2 are two constants representing delay and
 77 $\tau = \max\{\tau_1, \tau_2\}$. $d_1(t)$ and $d_2(t)$ denote the diffusion rates of the prey and the predator species
 78 at time t respectively. $a_{ii}(t)$ represent the interaction within i -th species, $a_{12}(t)$ is the capturing
 79 rate of the predator, and $a_{21}(t)$ is the effective conversion rate of predator. $r_1(t)$ and $r_2(t)$ are
 80 the reproduction rate of prey (in the absence of predator) and the natural death rate of predator.
 81 All the coefficients of the model (1.1)-(1.2) are continuous and positive ω -periodic functions. The
 82 model (1.1)-(1.2) is an extension of the classic Lotka-Volterra model, and its degenerate model

83 has been extensively studied, for example see [28–30]. Amine and Ortega [28] obtained a stability
84 criterion on non-constant periodic solutions for the following model

$$\begin{cases} \frac{du_1(t)}{dt} = u_1(t)(a(t) - b(t)u_1(t) - c(t)u_2(t)), \\ \frac{du_2(t)}{dt} = u_2(t)(d(t) - e(t)u_2(t) + f(t)u_1(t)), \end{cases} \quad (1.3)$$

85 in terms of the L^∞ norm of the coefficients of a planar linear system associated to the model (1.3).
86 Ortega [29] gave another stability criteria in terms of the L^1 norm and Ortega and Rebelo [30]
87 obtained a new stability criterion which establishes a bridge between the stability criteria in [28]
88 and [29] in terms of L^p norm.

89 The stability on time-periodic solution for nonautonomous DRDEs has not been studied before.
90 In this article, we intend to study the time-periodic solutions for the predator-prey Lotka-Volterra
91 models governed by nonautonomous DRDEs and generalize the stability result on (1.3) obtained
92 in [30]. Meanwhile, the methods obtained in this article can also be used to extend the permanent
93 result obtained in [31] to nonautonomous cooperative DRDEs.

94 The article organization are showed as follows. In Section 2, we will investigate the existence
95 of the time-periodic solution of the nonautonomous predator-prey DRDEs. In Section 3, we pay
96 more attention to the globally asymptotically stability of the time-periodic solution. In Section 4,
97 we will give a numerical example to show the application of the theoretical findings obtained in
98 this article.

99 **Remark 1.1.** *The innovations and achievements of this article are listed as follows: (1) By*
100 *introducing the time delays and the variable coefficient into the known population models, a new*
101 *Lotka-Volterra predator-prey model (nonautonomous predator-prey DRDEs) that can more truly*
102 *depict the interaction among populations is proposed. (2) By considering of the upper and lower*
103 *solution methods and Lyapunov stability theory as well as fixed point theory, some new theories and*
104 *methods have been creatively developed, the existence and stability of the positive time-dependent*
105 *periodic solution of the new predator-prey DRDEs are obtained only a set of simplify verified*
106 *conditions are needed. (3) The technique of constructing Lyapunov functions for delayed differential*
107 *equations step by step can be used to study related problems, which will provide an effective method*
108 *to study the stability of solutions to delayed partial differential equations. (4) Compared with the*
109 *existing results, the stable solution obtained in this article is a time-periodic solution rather than a*
110 *constant periodic solution or a solution for a steady-state system, which will be more in line with*
111 *the objective law of seasonal cyclical changes in species density.*

112 2. EXISTENCE OF SPATIAL HOMOGENEOUS PERIODIC SOLUTIONS

113 Set $\varphi(t)$ be a ω -periodic function in \mathbb{R}_+ . We denote

$$\varphi^m = \sup\{\varphi(t), t \in \mathbb{R}_+\}, \varphi^l = \inf\{\varphi(t), t \in \mathbb{R}_+\}.$$

114 Next, we study the functional differential equations corresponding to the model (1.1)

$$\begin{cases} \frac{du_1(t)}{dt} = u_1(t)[r_1(t) - a_{11}(t)u_1(t - \tau_1) - a_{12}(t)u_2(t)], \\ \frac{du_2(t)}{dt} = u_2(t)[-r_2(t) - a_{22}(t)u_2(t - \tau_2) + a_{21}(t)u_1(t - \tau_1)], \end{cases} \quad (2.1)$$

115 with the initial conditions

$$u_i(t) = \eta_{i0}(t) > 0, t \in [-\tau, 0], i = 1, 2. \quad (2.2)$$

116 **Theorem 2.1.** *For any positive initial conditions, the solution of the models (2.1)-(2.2) is positive.*

117 *Proof.* Due to the continuity of the functions on the right-hand side of model (2.1) and its satisfac-
118 tion of the local Lipschitz condition, models (2.1)-(2.2) possesses a unique local solution, denoted
119 as $u_1(t)$ and $u_2(t)$, on a small interval $[0, T)$ according to the existence and uniqueness theorem for
120 solutions of functional differential equations. Next, we prove that for any positive initial values,
121 this local solution remains positive and can be extended to the entire positive time axis.

122 According to the first equation of model (2.1), when $u_1(t) > 0$ and $u_2(t) > 0$, the sign of the
123 right-hand side of the equation depends on $r_1(t) - a_{11}(t)u_1(t - \tau_1) - a_{12}(t)u_2(t) \triangleq \Lambda_1(t)$. If the
124 initial value results in $\Lambda_1(t) > 0$, then the rate of change of $u_1(t)$ is positive, and since the initial
125 value $\eta_{10}(t)$ is positive, $u_1(t)$ will remain positive. If the initial values result in $\Lambda_1(t) < 0$, then

126 the rate of change of $u_1(t)$ is negative, and due to the positive initial value, the prey population
 127 $u_1(t)$ will decrease. **Based on the interaction mechanisms among populations in ecosystems and the**
 128 **continuity of population dynamics.** Subsequently, the predator population $u_2(t)$ will also decrease
 129 due to insufficient food. Since $r_1(t) > 0$, as the populations of $u_1(t)$ and $u_2(t)$ decrease, eventually
 130 $\Lambda_1(t)$ will become positive, causing the population of $u_1(t)$ to increase before reaching zero. In
 131 summary, regardless of whether the initial values make $\Lambda_1(t) > 0$ or $\Lambda_1(t) < 0$, $u_1(t)$ remains
 132 positive.

133 Similarly, according to the second equation of model (2.1), when $u_1(t) > 0$ and $u_2(t) > 0$, the
 134 sign of the right-hand side of the equation depends on $-r_2(t) - a_{22}(t)u_2(t - \tau_2) + a_{21}(t)u_1(t - \tau_1) \triangleq$
 135 $\Lambda_2(t)$. If the initial values result in $\Lambda_2(t) > 0$, then the rate of change of $u_2(t)$ is positive, and
 136 since the initial value $\eta_{20}(t)$ is positive, $u_2(t)$ will remain positive. If the initial values result in
 137 $\Lambda_2(t) < 0$, then the rate of change of $u_2(t)$ is negative, and due to the positive initial value, the
 138 predator population $u_2(t)$ will continuously decrease. **Based on the interaction mechanisms among**
 139 **populations in ecosystems and the continuity of population dynamics.** Simultaneously, the prey
 140 population $u_1(t)$ will increase due to the reduction in predators. As $u_2(t)$ decreases and $u_1(t)$
 141 increases, eventually $\Lambda_2(t)$ will become positive (since $a_{12}(t) > 0$), causing the population of $u_2(t)$
 142 to increase before decreasing to zero. In summary, regardless of whether the initial values make
 143 $\Lambda_2(t) > 0$ or $\Lambda_2(t) < 0$, $u_2(t)$ remains positive.

144 Furthermore, since the local solution is unique and positive, we can utilize the continuous
 145 dependence theorem for solutions of functional differential equations to extend the local solution to
 146 the entire positive time axis while maintaining its positivity. Therefore, given the initial conditions
 147 $\eta_{10}(t) > 0$ and $\eta_{20}(t) > 0$, the solutions of the predator-prey models (2.1)-(2.2) remain positive on
 148 the entire positive time axis. \square

149 For the model (2.1), set

$$150 \quad M_1 = \frac{r_1^m}{a_{11}^l} \exp \{r_1^m \tau_1\}, M_2 = \frac{a_{21}^m M_1 - r_2^l}{a_{22}^l} \exp \{(a_{21}^m M_1 - r_2^l) \tau_2\},$$

$$151 \quad m_1 = \frac{r_1^m - a_{12}^m M_2}{a_{11}^m} \exp \{(r_1^l - a_{12}^m M_2 - a_{11}^m M_1) \tau_1\},$$

$$152 \quad m_2 = \frac{a_{21}^l m_1 - r_2^m}{a_{22}^m} \exp \{(a_{21}^l m_1 - r_2^m - a_{22}^m M_2) \tau_2\}.$$

152 **Theorem 2.2.** *Assume the following conditions satisfy*

$$153 \quad (H_1) \quad a_{21}^l m_1 > r_2^m,$$

$$154 \quad (H_2) \quad r_1^l > a_{12}^m M_2.$$

154 *Then the model (2.1)-(2.2) is permanent.*

155 *Proof.* By the first equation of model (2.1), it follows that

$$156 \quad \frac{du_1(t)}{dt} = u_1(t)[r_1(t) - a_{11}(t)u_1(t - \tau_1) - a_{12}(t)u_2(t)] \leq u_1(t)[r_1^m - a_{11}^l u_1(t - \tau_1)]. \quad (2.3)$$

156 From the Lemma 2.2 in [31], one has

$$157 \quad \limsup_{t \rightarrow +\infty} u_1(t) \leq \frac{r_1^m}{a_{11}^l} \exp \{r_1^m \tau_1\} = M_1. \quad (2.4)$$

157 Moreover, from the second equation of model (2.1), it holds that

$$\begin{aligned} \frac{du_2(t)}{dt} &\leq u_2(t)[-r_2(t) - a_{22}(t)u_2(t - \tau_2) + a_{21}(t)u_1(t - \tau_1)] \\ &\leq u_2(t)[-r_2^l - a_{22}^l u_2(t - \tau_2) + a_{21}^m M_1]. \end{aligned}$$

158 By (H_1) , we have $a_{21}^m M_1 > r_2^l$. Thus, by Lemma 2.2 in [31],

$$159 \quad \limsup_{t \rightarrow +\infty} u_2(t) \leq \frac{a_{21}^m M_1 - r_2^l}{a_{22}^l} \exp \{(a_{21}^m M_1 - r_2^l) \tau_2\} = M_2. \quad (2.5)$$

159 On the other hand, by (2.1),

$$\frac{du_1(t)}{dt} \geq u_1(t)[r_1^l - a_{11}^m u_1(t - \tau_1) - a_{12}^m M_2] = u_1(t)[r_1^l - a_{12}^m M_2 - a_{11}^m u_1(t - \tau_1)].$$

160 By (H_2) and Lemma 2.3 in [31],

$$\liminf_{t \rightarrow +\infty} u_1(t) \geq \frac{r_1^l - a_{12}^m M_2}{a_{11}^m} \exp[(r_1^l - a_{12}^m M_2 - a_{11}^m M_1)\tau_1] = m_1. \quad (2.6)$$

161 Similarly, by the second equation in model (2.1), we have

$$\begin{aligned} \frac{du_2(t)}{dt} &= u_2(t)[-r_2(t) - a_{22}(t)u_2(t - \tau_2) + a_{21}(t)u_1(t - \tau_1)] \geq u_2(t)[-r_2^m - a_{22}^m u_2(t - \tau_2) + a_{21}^l m_1] \\ &= u_2(t)[a_{21}^l m_1 - r_2^m - a_{22}^m u_2(t - \tau_2)]. \end{aligned}$$

162 By (H_1) and Lemma 2.3 in [31],

$$\liminf_{t \rightarrow +\infty} u_2(t) \geq \frac{a_{21}^l m_1 - r_2^m}{a_{22}^m} \exp[(a_{21}^l m_1 - r_2^m - a_{22}^m M_2)\tau_2] = m_2. \quad (2.7)$$

163 By (2.4)-(2.7), we see that the model (2.1)-(2.2) is permanent, see [Definition 2.1, [32]] for the
164 definition of permanent property. \square

165 **Theorem 2.3.** *Assume that $(H_1) - (H_2)$ hold. Then there is a strictly positive spatial homogeneity
166 ω -periodic solution of (1.1)-(1.2).*

167 *Proof.* Let $V = C([- \tau, +\infty), \mathbb{R}_+^2)$ be a Banach space consisting of continuous, bounded, ω -periodic
168 and positive functions defined on $[- \tau, +\infty)$, equipped with the infinite norm. Based on the exis-
169 tence and uniqueness theorem of solutions of the functional differential equations, see [Theorem
170 2.3, page 42 of [33]], we define a Poincaré mapping $\psi : V \rightarrow V$ in the following form

$$\psi(U_0) = U(t, \omega, U_0),$$

171 where $U(t, \omega, U_0) = (u_1(t), u_2(t))$ is a positive solution of the functional differential equations (2.1)
172 subject to the initial conditions $U_0 = (\eta_{10}(t), \eta_{20}(t)), t \in [- \tau, 0]$.

173 It easy to see that ψ is continuous mapping by using the continuity of solution of the functional
174 differential equations (2.1) with regard to the above initial conditions, see [Theorem 4.1, page 46
175 of [33]]. Assume that K is any bounded set in V . For any $U_0 \in K$, and let $L = \sqrt{M_1^2 + M_2^2}$.
176 From the permanence of solutions to models (2.1)-(2.2), we have $\|\psi(U_0)\| = \|U(t, \omega, U_0)\| =$
177 $\|(u_1(t), u_2(t))\| = \sqrt{u_1(t)^2 + u_2(t)^2} \leq \sqrt{M_1^2 + M_2^2} = L$. Hence, $\psi(K)$ is uniformly bounded.
178 Furthermore, according to Theorem 2.2, the derivative of the mapping ψ is also bounded, which
179 can then be used to prove that the $\psi(K)$ is equicontinuous. The Arzela-Ascoli theorem implies
180 that ψ is completely continuous.

181 We define

$$S = \{(u_1(t), u_2(t)) \in V \mid m_i \leq u_i(t) \leq M_i, i = 1, 2\}, \quad (2.8)$$

182 then it is obvious that S is a closed bounded convex subset of the Banach space V . By Theorem
183 2.2 we have that ψ is a mapping from S to S . Thus, by Schauder fixed-point theorem, see [Lemma
184 2.4, page 40 of [33]], the mapping ψ has a fixed point $(u_1^*(t), u_2^*(t))$. That is, the equations (2.1)-
185 (2.2) have a positive ω -periodic solution $(u_1^*(t), u_2^*(t))$ which is the spatial homogeneity ω -periodic
186 solution for models (1.1)-(1.2), see [Definition 2.2, [34]]. \square

187 3. STABILITY OF SPATIAL HOMOGENEITY PERIODIC SOLUTION

188 In this section, we provide some sufficient conditions to obtain the globally asymptotically stable
189 of spatial homogeneity ω -periodic solution of (1.1) by using the method of upper and lower solutions
190 for the delayed parabolic partial differential equations and Lyapunov stability theory.

191 **Theorem 3.1.** *Assume that $(H_1) - (H_2)$ and the following assumptions hold.*

192 $(H_3) A_1 = a_{11}^l - a_{11}^m \tau_1 [r_1^m + a_{11}^m M_1 + a_{12}^m M_2] - M_1 (a_{11}^m)^2 \tau_1 - a_{21}^m (1 + a_{22}^m M_2 \tau_2) > 0,$

$(H_4) A_2 = a_{22}^l - a_{22}^m \tau_2 [r_2^m + a_{22}^m M_2 + a_{21}^m M_1] - M_2 (a_{22}^m)^2 \tau_2 - (1 + M_1 a_{11}^m \tau_1) a_{12}^m > 0.$

193 *Then there exists a spatial homogeneity strictly positive and globally asymptotically stable ω -periodic
194 solution $(u_1^*(t), u_2^*(t))$, that is, the solution $(u_1(x, t), u_2(x, t))$ of (1.1)-(1.2) with any positive initial
195 values fulfills*

$$\lim_{t \rightarrow \infty} (u_i(x, t) - u_i^*(t)) = 0, \text{ uniformly for } x \in \bar{\Omega}, i = 1, 2. \quad (3.1)$$

196 *Proof.* By Theorem 2.3, (1.1)-(1.2) has a spatial homogeneity strictly positive ω -periodic solution.
 197 We prove the stability of the solution. Because the solutions and coefficients of model (1.1) are
 198 positive, it is easy to see that the reaction functions of model (1.1) are mixed quasimonotone.
 199 Let $l_i = \min_{x \in \bar{\Omega}, t \in [-\tau, 0]} \eta_{i0}(x, t)$, $r_i = \max_{x \in \bar{\Omega}, t \in [-\tau, 0]} \eta_{i0}(x, t)$. Then $0 < l_i \leq \eta_{i0}(x, t) \leq r_i$. Let
 200 $(\tilde{u}_1(t), \tilde{u}_2(t))$ and $(\hat{u}_1(t), \hat{u}_2(t))$ be the solutions of (2.1) subject to initial values $(\eta_{10}(t), \eta_{20}(t)) =$
 201 (r_1, r_2) and $(\eta_{10}(t), \eta_{20}(t)) = (l_1, l_2)$ respectively, then there exist upper and lower solutions
 202 $(\tilde{u}_1(t), \tilde{u}_2(t))$ and $(\hat{u}_1(t), \hat{u}_2(t))$ of (1.1)-(1.2). By Theorem 2.1 in [35], (1.1)-(1.2) has a unique
 203 solution $(u_1(x, t), u_2(x, t))$, $(x, t) \in \bar{\Omega} \times [-\tau, +\infty)$, which satisfies

$$(\hat{u}_1(t), \hat{u}_2(t)) \leq (u_1(x, t), u_2(x, t)) \leq (\tilde{u}_1(t), \tilde{u}_2(t)). \quad (3.2)$$

204 We prove

$$\lim_{t \rightarrow \infty} (\tilde{u}_i(t) - u_i^*(t)) = \lim_{t \rightarrow \infty} (\hat{u}_i(t) - u_i^*(t)) = 0, (i = 1, 2). \quad (3.3)$$

205 We first prove the solution $(u_1(t), u_2(t))$ for the functional differential equations (2.1) with any
 206 positive initial $(u_1(t), u_2(t)) = (\eta_{10}(t), \eta_{20}(t))$ satisfies

$$\lim_{t \rightarrow \infty} (u_i(t) - u_i^*(t)) = 0, i = 1, 2. \quad (3.4)$$

207 By Theorem 2.2, there exist five positive real numbers M_i , m_i and T such that

$$m_i \leq u_i(t) \leq M_i \text{ when } t > T.$$

208 Let

$$V_{11}(t) = |\ln u_1(t) - \ln u_1^*(t)|.$$

We denote by $D^+ V_{11}(t)$ the right-side derivative of $V_{11}(t)$, then

$$\begin{aligned} D^+ V_{11}(t) &= \text{sgn}(u_1(t) - u_1^*(t))[-a_{11}(t)(u_1(t - \tau_1) - u_1^*(t - \tau_1)) - a_{12}(t)(u_2(t) - u_2^*(t))] \\ &= \text{sgn}(u_1(t) - u_1^*(t))[-a_{11}(t)(u_1(t) - u_1^*(t)) - a_{12}(t)(u_2(t) - u_2^*(t)) \\ &\quad + a_{11}(t) \int_{t-\tau_1}^t (\dot{u}_1(\theta) - \dot{u}_1^*(\theta)) d\theta] \\ &= \text{sgn}(u_1(t) - u_1^*(t))[-a_{11}(t)(u_1(t) - u_1^*(t)) - a_{12}(t)(u_2(t) - u_2^*(t)) \\ &\quad + a_{11}(t) \int_{t-\tau_1}^t \{u_1(\theta)[r_1(\theta) - a_{11}(\theta)u_1(\theta - \tau_1) - a_{12}(\theta)u_2(\theta)] \\ &\quad - u_1^*(\theta)[r_1(\theta) - a_{11}(\theta)u_1^*(\theta - \tau_1) - a_{12}(\theta)u_2^*(\theta)]\} d\theta] \\ &= \text{sgn}(u_1(t) - u_1^*(t))[-a_{11}(t)(u_1(t) - u_1^*(t)) - a_{12}(t)(u_2(t) - u_2^*(t)) \\ &\quad + a_{11}(t) \int_{t-\tau_1}^t \{(u_1(\theta) - u_1^*(\theta))[r_1(\theta) - a_{11}(\theta)u_1^*(\theta - \tau_1) - a_{12}(\theta)u_2^*(\theta)] \\ &\quad - u_1(\theta)[a_{11}(\theta)(u_1(\theta - \tau_1) - u_1^*(\theta - \tau_1)) + a_{12}(\theta)(u_2(\theta) - u_2^*(\theta))]\} d\theta] \\ &\leq -a_{11}(t) |u_1(t) - u_1^*(t)| + a_{12}(t) |u_2(t) - u_2^*(t)| \\ &\quad + a_{11}(t) \int_{t-\tau_1}^t ([r_1(\theta) + a_{11}(\theta)u_1^*(\theta - \tau_1) + a_{12}(\theta)u_2^*(\theta)] |u_1(\theta) - u_1^*(\theta)| \\ &\quad + u_1(\theta)[a_{11}(\theta) |u_1(\theta - \tau_1) - u_1^*(\theta - \tau_1)| + a_{12}(\theta) |u_2(\theta) - u_2^*(\theta)|]) d\theta. \end{aligned} \quad (3.5)$$

209 Let

$$V_{12}(t) = \int_{t-\tau_1}^t \int_s^t a_{11}(s + \tau_1) ([r_1(\theta) + a_{11}(\theta)u_1^*(\theta - \tau_1) + a_{12}(\theta)u_2^*(\theta)] |u_1(\theta) - u_1^*(\theta)| \\ + u_1(\theta)[a_{11}(\theta) |u_1(\theta - \tau_1) - u_1^*(\theta - \tau_1)| + a_{12}(\theta) |u_2(\theta) - u_2^*(\theta)|]) d\theta ds. \quad (3.6)$$

210 By (3.5) and (3.6),

$$\begin{aligned} D^+ \sum_{i=1}^2 V_{1i}(t) &\leq -a_{11}(t) |u_1(t) - u_1^*(t)| + a_{12}(t) |u_2(t) - u_2^*(t)| \\ &\quad + \int_{t-\tau_1}^t a_{11}(s + \tau_1) ds ([r_1(t) + a_{11}(t)u_1^*(t - \tau_1) + a_{12}(t)u_2^*(t)] |u_1(t) - u_1^*(t)| \\ &\quad + u_1(t)[a_{11}(t) |u_1(t - \tau_1) - u_1^*(t - \tau_1)| + a_{12}(t) |u_2(t) - u_2^*(t)|]) \\ &\leq -a_{11}(t) |u_1(t) - u_1^*(t)| + a_{12}(t) |u_2(t) - u_2^*(t)| \\ &\quad + \int_{t-\tau_1}^t a_{11}(s + \tau_1) ds [r_1(t) + a_{11}(t)M_1 + a_{12}(t)M_2] |u_1(t) - u_1^*(t)| \\ &\quad + M_1 \int_{t-\tau_1}^t a_{11}(s + \tau_1) ds [a_{11}(t) |u_1(t - \tau_1) - u_1^*(t - \tau_1)| + a_{12}(t) |u_2(t) - u_2^*(t)|] \\ &\leq (-a_{11}^l + a_{11}^m \tau_1 [r_1^m + a_{11}^m M_1 + a_{12}^m M_2]) |u_1(t) - u_1^*(t)| \\ &\quad + M_1 (a_{11}^m)^2 \tau_1 |u_1(t - \tau_1) - u_1^*(t - \tau_1)| + a_{12}^m (1 + a_{11}^m M_1 \tau_1) |u_2(t) - u_2^*(t)|. \end{aligned} \quad (3.7)$$

211 Let

$$V_{13}(t) = M_1 (a_{11}^m)^2 \tau_1 \int_{t-\tau_1}^t |(u_1(w) - u_1^*(w))| dw, \quad (3.8)$$

212 and

$$V_1(t) = V_{11}(t) + V_{12}(t) + V_{13}(t). \quad (3.9)$$

213 By (3.7) and (3.8), we have

$$D^+V_1(t) \leq (-a_{11}^l + a_{11}^m \tau_1 [r_1^m + a_{11}^m M_1 + a_{12}^m M_2] + M_1 (a_{11}^m)^2 \tau_1) |u_1(t) - u_1^*(t)| \\ + (1 + M_1 a_{11}^m \tau_1) a_{12}^m |u_2(t) - u_2^*(t)|. \quad (3.10)$$

214 Similarly, we define $V_{21}(t) = |\ln u_2(t) - \ln u_2^*(t)|$, and we have

$$D^+V_{21}(t) = \operatorname{sgn}(u_2(t) - u_2^*(t)) [-a_{22}(t)(u_2(t - \tau_2) - u_2^*(t - \tau_2)) + a_{21}(t)(u_1(t - \tau_1) - u_1^*(t - \tau_1))] \\ = \operatorname{sgn}(u_2(t) - u_2^*(t)) [-a_{22}(t)(u_2(t) - u_2^*(t)) + a_{21}(t)(u_1(t - \tau_1) - u_1^*(t - \tau_1)) \\ + a_{22}(t) \int_{t-\tau_2}^t (\dot{u}_2(\theta) - \dot{u}_2^*(\theta)) d\theta] \\ = \operatorname{sgn}(u_2(t) - u_2^*(t)) [-a_{22}(t)(u_2(t) - u_2^*(t)) + a_{21}(t)(u_1(t - \tau_1) - u_1^*(t - \tau_1)) \\ + a_{22}(t) \int_{t-\tau_2}^t \{u_2(\theta) [-r_2(\theta) - a_{22}(\theta)u_2(\theta - \tau_2) + a_{21}(\theta)u_1(\theta - \tau_1)] \\ - u_2^*(\theta) [-r_2(\theta) - a_{22}(\theta)u_2^*(\theta - \tau_2) + a_{21}(\theta)u_1^*(\theta - \tau_1)]\} d\theta] \\ = \operatorname{sgn}(u_2(t) - u_2^*(t)) [-a_{22}(t)(u_2(t) - u_2^*(t)) + a_{21}(t)(u_1(t - \tau_1) - u_1^*(t - \tau_1)) \\ + a_{22}(t) \int_{t-\tau_2}^t \{(u_2(\theta) - u_2^*(\theta)) [-r_2(\theta) - a_{22}(\theta)u_2^*(\theta - \tau_2) + a_{21}(\theta)u_1^*(\theta - \tau_1)] \\ - u_2(\theta) [a_{22}(\theta)(u_2(\theta - \tau_2) - u_2^*(\theta - \tau_2)) - a_{21}(\theta)(u_1(\theta - \tau_1) - u_1^*(\theta - \tau_1))]\} d\theta] \\ \leq -a_{22}(t) |u_2(t) - u_2^*(t)| + a_{21}(t) |u_1(t - \tau_1) - u_1^*(t - \tau_1)| \\ + a_{22}(t) \int_{t-\tau_2}^t ([r_2(\theta) + a_{22}(\theta)u_2^*(\theta - \tau_2) + a_{21}(\theta)u_1^*(\theta - \tau_1)] |u_2(\theta) - u_2^*(\theta)| \\ + u_2(\theta) [a_{22}(\theta) |u_2(\theta - \tau_2) - u_2^*(\theta - \tau_2)| + a_{21}(\theta) |u_1(\theta - \tau_1) - u_1^*(\theta - \tau_1)|]) d\theta. \quad (3.11)$$

215 Let

$$V_{22}(t) = \int_{t-\tau_2}^t \int_s^t a_{22}(s + \tau_2) ([r_2(\theta) + a_{22}(\theta)u_2^*(\theta - \tau_2) + a_{21}(\theta)u_1^*(\theta - \tau_1)] |u_2(\theta) - u_2^*(\theta)| \\ + u_2(\theta) [a_{22}(\theta) |u_2(\theta - \tau_2) - u_2^*(\theta - \tau_2)| + a_{21}(\theta) |u_1(\theta - \tau_1) - u_1^*(\theta - \tau_1)|]) d\theta ds. \quad (3.12)$$

216 By (3.11) and (3.12),

$$D^+ \sum_{i=1}^2 V_{2i}(t) \leq -a_{22}(t) |u_2(t) - u_2^*(t)| + a_{21}(t) |u_1(t - \tau_1) - u_1^*(t - \tau_1)| \\ + \int_{t-\tau_2}^t a_{22}(s + \tau_2) ds ([r_2(t) + a_{22}(t)u_2^*(t - \tau_2) + a_{21}(t)u_1^*(t - \tau_1)] |u_2(t) - u_2^*(t)| \\ + u_2(t) [a_{22}(t) |u_2(t - \tau_2) - u_2^*(t - \tau_2)| + a_{21}(t) |u_1(t - \tau_1) - u_1^*(t - \tau_1)|]) \\ \leq -a_{22}(t) |u_2(t) - u_2^*(t)| + a_{21}(t) |u_1(t - \tau_1) - u_1^*(t - \tau_1)| + \\ \int_{t-\tau_2}^t a_{22}(s + \tau_2) ds [r_2(t) + a_{22}(t)M_2 + a_{21}(t)M_1] |u_2(t) - u_2^*(t)| \\ \leq +M_2 \int_{t-\tau_2}^t a_{22}(s + \tau_2) ds [a_{22}(t) |u_2(t - \tau_2) - u_2^*(t - \tau_2)| \\ + a_{21}(t) |u_1(t - \tau_1) - u_1^*(t - \tau_1)|] \\ \leq (-a_{22}^l + a_{22}^m \tau_2 [r_2^m + a_{22}^m M_2 + a_{21}^m M_1]) |u_2(t) - u_2^*(t)| \\ + M_2 (a_{22}^m)^2 \tau_2 |u_2(t - \tau_2) - u_2^*(t - \tau_2)| + a_{21}^m (1 + a_{22}^m M_2 \tau_2) |u_1(t - \tau_1) - u_1^*(t - \tau_1)|. \quad (3.13)$$

217 Let

$$V_{23}(t) = M_2 (a_{22}^m)^2 \tau_2 \int_{t-\tau_2}^t |(u_2(w) - u_2^*(w))| dw + a_{21}^m (1 + a_{22}^m M_2 \tau_2) \int_{t-\tau_1}^t |(u_1(w) - u_1^*(w))| dw, \quad (3.14)$$

218 and

$$V_2(t) = V_{21}(t) + V_{22}(t) + V_{23}(t). \quad (3.15)$$

219 By (3.13) and (3.14),

$$D^+V_2(t) \leq (-a_{22}^l + a_{22}^m \tau_2 [r_2^m + a_{22}^m M_2 + a_{21}^m M_1] + M_2 (a_{22}^m)^2 \tau_2) |u_2(t) - u_2^*(t)| \\ + a_{21}^m (1 + a_{22}^m M_2 \tau_2) |u_1(t) - u_1^*(t)|. \quad (3.16)$$

220 We define a Lyapunov function as follows

$$V(t) = V_1(t) + V_2(t).$$

221 By (3.10) and (3.16), we get

$$D^+V(t) \leq -A_1 |u_1(t) - u_1^*(t)| - A_2 |u_2(t) - u_2^*(t)|. \quad (3.17)$$

222 Integrating from ω to t on both sides of (3.17), we have

$$V(t) + \alpha \int_{\omega}^t (|u_1(s) - u_1^*(s)| + |u_2(s) - u_2^*(s)|) ds \leq V(\omega) < +\infty, \quad (3.18)$$

223 where $\alpha = \min\{A_1, A_2\} > 0$. Therefore, $V(t)$ is bounded on $[\omega, +\infty)$, and

$$\int_{\omega}^t (|u_1(s) - u_1^*(s)| + |u_2(s) - u_2^*(s)|) ds \leq \frac{V(\omega)}{\alpha} < +\infty. \quad (3.19)$$

224 By (3.19), we have

$$(|u_1(t) - u_1^*(t)| + |u_2(t) - u_2^*(t)|) \in L^1(T, +\infty). \quad (3.20)$$

225 From the uniform permanence of model (2.1), we have that $|u_1(t) - u_1^*(t)| + |u_2(t) - u_2^*(t)|$
226 and its derivative are bounded. Thus, $|u_1(t) - u_1^*(t)| + |u_2(t) - u_2^*(t)|$ is uniformly continuous on
227 $[\omega, +\infty)$. By Lemma 8.2 in [36], we get

$$\lim_{t \rightarrow +\infty} |u_i(t) - u_i^*(t)| = 0, (i = 1, 2).$$

228 From (3.2) and the squeeze theorem, (3.1) holds true. That is, (1.1)-(1.2) have a spatial homo-
229 geneity strictly positive and globally asymptotically stable ω -periodic solution $(u_1^*(t), u_2^*(t))$, see
230 [Definition 2.3, [34]]. This completes the proof of Theorem 3.1. \square

231 **Theorem 3.2.** *Suppose that the ω -periodic model (1.1) satisfies assumptions $(H_1) - (H_4)$, then*
232 *the model (1.1) is permanent, i.e., the solution $(u_1(x, t), u_2(x, t))$ of models (1.1)-(1.2) with any*
233 *initial values fulfills*

$$m_i \leq u_i(x, t) \leq M_i, \text{ uniformly for } (x, t) \in \bar{\Omega} \times [T, +\infty), i = 1, 2. \quad (3.21)$$

234 *Proof.* By means of Theorem 2.3, there exist four positive real numbers m_i, M_i , ($i = 1, 2$) such
235 that

$$m_i \leq u_i^*(t) = u_i^*(t + \omega) \leq M_i, t \in [-\tau, +\infty). \quad (3.22)$$

236 Moreover, from Theorem 3.1, one has

$$\lim_{t \rightarrow +\infty} u_i(x, t) = u_i^*(t), \text{ uniformly for } x \in \bar{\Omega}, i = 1, 2. \quad (3.23)$$

237 Therefore, from (3.22) and (3.23), the model (1.1) is permanent. \square

238

4. NUMERICAL SIMULATIONS

239 In this section, we provide a numerical example to show the application of Theorem 3.1. For the
240 convenience of calculation and numerical simulation, we choose 2-period functions as the coefficients
241 for the nonautonomous ω -periodic DRDEs (1.1)-(1.2).

242 **Example 4.1.** Consider the following 2-species DRDEs. In view of the conditions $(H_1) - (H_4)$
243 of Theorem 3.1, with the help of some calculations we choose some special values of parameters
244 shown in models (4.1)-(4.2). It should be noted that, the selection of above parameters is not
245 unique.

$$\begin{cases} \frac{\partial u_1(x, t)}{\partial t} - \Delta u_1(x, t) = u_1(x, t)[(24 + \cos \pi t) - (6 + \sin \pi t)u_1(x, t - 0.001) \\ \quad - (0.75 + 0.25 \sin \pi t)u_2(x, t)], 2\pi > x > 0, t > 0, \\ \frac{\partial u_2(x, t)}{\partial t} - \Delta u_2(x, t) = u_2(x, t)[-(2 + \cos \pi t) - (5 + \sin \pi t)u_2(x, t - 0.002) \\ \quad + (1.2 + 0.2 \sin \pi t)u_1(x, t - 0.001)], 2\pi > x > 0, t > 0, \end{cases} \quad (4.1)$$

246 with the Neumann boundary and initial conditions

$$\begin{cases} \frac{\partial u_1(x, t)}{\partial n} = \frac{\partial u_2(x, t)}{\partial n} = 0, \quad t > 0, x = 0, 2\pi, \\ u_1(x, t) = (4 + 3t)[1 - \sin(x + 0.5\pi)], \\ u_2(x, t) = (0.6 + 5t)[1 + \cos(x + \pi)], \\ (x, t) \in (0, 2\pi) \times [-0.002, 0]. \end{cases} \quad (4.2)$$

247 By calculating, we have

$$M_1 = \frac{r_1^m}{a_{11}^l} \exp \{r_1^m \tau_1\} \approx 5.1266,$$

248

$$M_2 = \frac{a_{21}^m M_1 - r_2^l}{a_{22}^l} \exp \{(a_{21}^m M_1 - r_2^l) \tau_2\} \approx 1.5635,$$

249

$$m_1 = \frac{r_1^m - a_{12}^m M_2}{a_{11}^m} \exp \{ (r_1^l - a_{12}^m M_2 - a_{11}^m M_1) \tau_1 \} \approx 3.0185,$$

250

$$m_2 = \frac{a_{21}^l m_1 - r_2^m}{a_{22}^m} \exp \{ (a_{21}^l m_1 - r_2^m - a_{22}^m M_2) \tau_2 \} \approx 0.0030,$$

251

$$a_{21}^l m_1 - r_2^m \approx 0.0185 > 0, r_1^l - a_{12}^m M_2 \approx 21.4365 > 0,$$

252

$$A_1 = a_{11}^l - a_{11}^m \tau_1 [r_1^m + a_{11}^m M_1 + a_{12}^m M_2] - M_1 (a_{11}^m)^2 \tau_1 - a_{21}^m (1 + a_{22}^m M_2 \tau_2) \approx 2.8854 > 0,$$

253

$$A_2 = a_{22}^l - a_{22}^m \tau_2 [r_2^m + a_{22}^m M_2 + a_{21}^m M_1] - M_2 (a_{22}^m)^2 \tau_2 - (1 + M_1 a_{11}^m \tau_1) a_{12}^m \approx 2.6166 > 0.$$

254 Based on the above calculation results, it is easy to see that that systems (4.1)-(4.2) satisfy the
 255 conditions of Theorem 3.1. From Theorem 3.1 it is easy to know that the systems (4.1)-(4.2) has
 256 a strictly positive spatial homogeneity 2-periodic solution $(u_1(x, t), u_2(x, t))$ which satisfies

$$\lim_{t \rightarrow +\infty} |u_i(t) - u_i^*(t)| = 0, (i = 1, 2), \text{ uniformly for } x \in (0, 2\pi).$$

257 By employing the software package MATLAB 7.1 and the finite differences method, we can obtain
 258 some numerical solutions of the model (4.1) with the boundary conditions and initial conditions
 259 (4.2) which are shown in Figure 4.1 to Figure 4.2. From Figures 4.1-4.2, it is not difficult to find that
 260 the model (4.1)-(4.2) have a strictly positive globally asymptotically stable spatial homogeneity 2-
 261 periodic solution. In model (4.1)-(4.2), the densities of prey and predator will oscillate periodically
 262 with a period of 2 and distribute homogeneously in space when the time is long enough. In order
 263 to verify that the periodic solution of the model (4.1)-(4.2) is globally asymptotically stable, we
 264 selected different initial values and conducted extensive numerical simulations. The results showed
 265 that the 2-periodic solution of the model (4.1)-(4.2) is asymptotically stable for any positive initial
 266 value. Please refer to Figure 4.3 for details.

267 From the theoretical research in this article, we can clearly see the dynamic stability mechanism
 268 of the predator-prey system. When the birth rate of prey species and the post-predation nutrient
 269 absorption rates of predator species reach sufficient levels, predator and prey species can maintain a
 270 long-term stable survival state, effectively avoiding the risk of population extinction (see Theorem
 271 2.1 and 3.2). More interestingly, under the basic conditions mentioned above, if key factors such as
 272 population diffusion rate, interaction strength, and predator natural mortality rate can also meet
 273 specific criteria, then under minor time delays, the density of species in the predator-prey system
 274 will exhibit periodic changes. This further reveals the beauty of dynamic equilibrium within e-
 275 cosystems (see Theorem 3.1). This theoretical result has been thoroughly validated through precise
 276 numerical simulations, providing a solid theoretical foundation for our understanding, evaluation,
 277 and maintenance of ecosystem balance. It is worth noting that the sufficient conditions established
 278 in this article are both concise and easy to verify. Specifically, the conditions proposed here are
 279 formulated as a series of inequalities rather than strict equations, providing great convenience and
 280 flexibility for the application of these theoretical results in practical ecosystem management.

281 **Remark 4.1.** *The method obtained in this article can not only be used to study the dynamic*
 282 *properties of solutions for various delayed reaction-diffusion predator-prey models, but also to study*
 283 *corresponding cooperative and competition models. In addition, this method may also be used to*
 284 *study the dynamic properties of solutions for time-varying delayed reaction-diffusion population*
 285 *models and fractional-order delayed reaction-diffusion population models.*

286

ACKNOWLEDGEMENTS

287 The authors express their sincere gratitude to the anonymous reviewer for providing insightful
 288 comments, which played a crucial role in enhancing the quality of the paper.

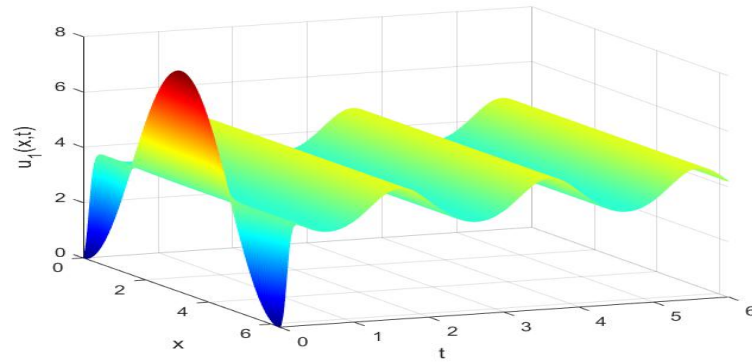


FIGURE 4.1. Evolution process of the density for the species $u_1(x,t)$ of model (4.1)-(4.2)

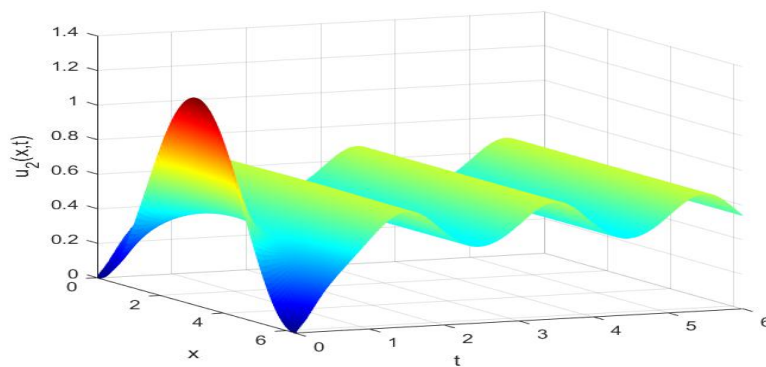


FIGURE 4.2. Evolution process of the density for the species $u_2(x,t)$ of model (4.1)-(4.2)

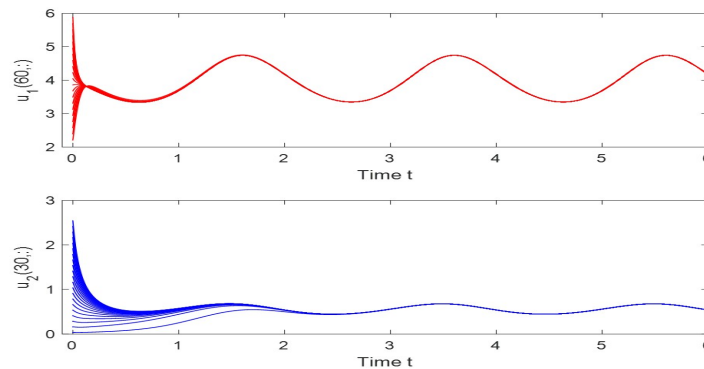


FIGURE 4.3. Evolution process of the densities for the species $u_1(x,t)$ and $u_2(x,t)$ of model (4.1)-(4.2) with different positive initial values

REFERENCES

- 290 [1] N. F. Britton, Reaction-diffusion equations and their applications to biology, Academic Press, New York, 1986.
 291 [2] J. Muller, C. Kuttler, Methods and models in mathematical biology: Deterministic and stochastic approaches,
 292 Springer, New York, 2015.
 293 [3] S. Djilali, Threshold asymptotic dynamics for a spatial age-dependent cell-to-cell transmission model with
 294 nonlocal disperse, Discrete and Continuous Dynamical Systems-Series B, 28 (7) (2023) 4108-4143.
 295 [4] S. Bentouta, S. Djilalia, Asymptotic profiles of a nonlocal dispersal SIR epidemic model with treat-age in a
 296 heterogeneous environment, Mathematics and Computers in Simulation, 203 (2023) 926-956.
 297 [5] S. Djilalia, Generalities on a delayed spatiotemporal host-pathogen infection model with distinct dispersal rates,
 298 Mathematical Modelling of Natural Phenomena, 19 (2024), ID: 11.

- 299 [6] S. Bentout, S. Djilali, Kuniya T, J. Wang, Mathematical analysis of a vaccination epidemic model with nonlocal
300 diffusion *Mathematical Methods in the Applied Sciences*, 46 (9) (2023) 10970-10994.
- 301 [7] S. Djilali, Dynamics of a spatiotemporal SIS epidemic model with distinct mobility range, *Applicable Analysis*,
302 2024, DOI: 10.1080/00036811.2024.2382267.
- 303 [8] W. Ko, I. Ahn, A diffusive one-prey and two-competing-predator system with a ratio-dependent functional
304 response: II stationary pattern formation, *Journal of Mathematical Analysis and Applications*, 397(1) (2013)
305 29-45.
- 306 [9] W. B. Yang, Y. L. Li, J. H. Wu, H. X. Li, Dynamics of a food chain model with ratio-dependent and modified
307 Leslie-Gower functional responses, *Discrete and Continuous Dynamical Systems-Series B*, 20 (7) (2015) 2269-
308 2290.
- 309 [10] J. F. Wang, Spatiotemporal patterns of a homogeneous diffusive predator-prey system with Holling type III
310 functional response, *Journal of Dynamics and Differential Equations*, 29 (2017) 1383-1409.
- 311 [11] D. Y. Wu, H. Y. Zhao, Spatiotemporal dynamics of a diffusive predator-prey system with allee effect and
312 threshold hunting, *Journal of Nonlinear Science*, 30 (2020) 1015-1054.
- 313 [12] S. Bentout, S. Djilali, A. Atangana, Bifurcation analysis of an age-structured prey-predator model with infection
314 developed in prey, *Mathematical Methods in the Applied Sciences*, 45 (3) (2021) 1189-1208.
- 315 [13] S. Djilali, C. Cattani, Patterns of a superdiffusive consumer-resource model with hunting cooperation functional
316 response, *Chaos, Solitons and Fractals*, 151 (2021), ID: 111258.
- 317 [14] X. P. Yan, C. H. Zhang, Spatiotemporal dynamics in a diffusive predator-prey system with Beddington-
318 DeAngelis functional response, *Qualitative Theory of Dynamical Systems*, 21 (4) (2022), ID: 166.
- 319 [15] M. X. Chen, R. C. Wu, Steady states and spatiotemporal evolution of a diffusive predator-prey model, *Chaos*
320 *Solitons & Fractals*, 170 (2023), ID: 113397.
- 321 [16] L. L. Jia, J. Huang, C. Y. Wang, Global stability of periodic solution for a 3-species nonautonomous ratio-
322 dependent diffusive predator-prey system, *Journal of Applied Analysis and Computation*, 14 (4) (2024) 1-19.
- 323 [17] J. Wu, *Theory and applications of partial functional differential equations*, Springer, New York, 1996.
- 324 [18] C. Y. Wang, Z. G. Yang, *Method of upper and lower solutions for reaction diffusion systems with delay*, Science
325 Press, Beijing, 2013. (in Chinese)
- 326 [19] S. S. Chen, J. S. Yu, Stability and bifurcations in a nonlocal delayed reaction-diffusion population model,
327 *Journal of Differential Equations*, 260 (2016) 218-240.
- 328 [20] Q. Y. Shi, J. P. Shi, Y. L. Song, Hopf bifurcation in a reaction-diffusion equation with distributed delay and
329 Dirichlet boundary condition, *Journal of Differential Equations*, 263 (2017) 6537-6575.
- 330 [21] Y. D. Yuan, Z. M. Guo, Monotone methods and stability results for nonlocal reaction-diffusion equations with
331 time delay, *Journal of Applied Analysis & Computation*, 8 (5) (2018) 1342-1368.
- 332 [22] Z. L. Shen, J. J. Wei, Spatiotemporal patterns in a delayed reaction-diffusion mussel-algae model, *International*
333 *Journal of Bifurcation and Chaos*, 29 (12) (2019), ID: 1950164.
- 334 [23] W. J. Zuo, J. P. Shi, Existence and stability of steady-state solutions of reaction-diffusion equations with
335 nonlocal delay effect, *Zeitschrift für angewandte Mathematik und Physik*, 72 (2) (2021), ID: 43.
- 336 [24] W. X. Xu, H. Y. Shu, Z. Tang, H. Wang, Complex dynamics in a general diffusive predator-
337 prey model with predator maturation delay, *Journal of Dynamics and Differential Equations*, 2022,
338 <https://doi.org/10.1007/s10884-022-10176-9>.
- 339 [25] Y. D. Yuan, Z. M. Guo, Global dynamics of a class of delayed differential systems with spatial non-locality,
340 *Journal of Differential Equations*, 349 (2023) 176-235.
- 341 [26] S. Djilali, S. Bentout, A. Zeb, Dynamics of a diffusive delayed viral infection model in a heterogeneous envi-
342 ronment, *Mathematical Methods in the Applied Sciences*, 46 (16) (2023) 16596-16624.
- 343 [27] V. Kumar, Pattern formation and delay-induced instability in a Leslie-Gower type prey-predator system with
344 Smith growth function, *Mathematics and Computers in Simulation*, 225 (2024) 78-97.
- 345 [28] Z. Amine, R. Ortega, A periodic prey-predator system, *Journal of Mathematical Analysis and Applications*,
346 185 (1994) 477-489.
- 347 [29] R. Ortega, Variations on Lyapunov's stability criterion and periodic prey-predator systems, *Electronic Research*
348 *Archive*, 29 (6) (2021) 3995-4008.
- 349 [30] V. Ortega, C. Rebelo, A note on stability criteria in the periodic Lotka-Volterra predator-prey model, *Applied*
350 *Mathematics Letters*, 145 (2023), ID: 10873.
- 351 [31] Y. Nakata, Y. Muroya, Permanence for nonautonomous Lotka-Volterra cooperative systems with delays, *Non-*
352 *linear Analysis: Real World Applications*, 11 (2010) 528-534.
- 353 [32] C. Wang, N. Li, Y. Zhou, X. Pu, R. Li, On a multi-delay Lotka-Volterra predator-prey model with feedback
354 controls and prey diffusion, *Acta Mathematica Scientia, Series B*, 39 (2019) 429-448.
- 355 [33] J. K. Hale, *Theory of Functional Differential Equations*, Springer-Verlag, New York, 1977.
- 356 [34] Y. Zhang, C. Wang, Stability analysis of n-species Lotka-Volterra almost periodic competition models with
357 grazing rates and diffusion, *International Journal of Biomathematics*, 7 (2) (2014), ID: 1450011.
- 358 [35] C. Wang, Existence and stability of periodic solutions for parabolic systems with time delays, *Journal of*
359 *Mathematical Analysis and Applications*, 339 (2008) 1354-1361.
- 360 [36] H. K. Khalil, *Nonlinear Systems*, 3rd ed, Prentice-Hall, Englewood Cliffs, 2002.

361 DEPARTMENT OF BASIC TEACHING, DIANCHI COLLEGE OF YUNNAN UNIVERSITY, KUNMING, YUNNAN, 650228, P.
362 R. CHINA, SCHOOL OF MATHEMATICAL SCIENCES, AND V.C. & V. R. KEY LAB OF SICHUAN PROVINCE, SICHUAN
363 NORMAL UNIVERSITY, CHENGDU, SICHUAN, 610066, P. R. CHINA
364 *E-mail address:* `lilijadianchi@163.com`

365 CORRESPONDING AUTHOR. COLLEGE OF APPLIED MATHEMATICS, CHENGDU UNIVERSITY OF INFORMATION TECH-
366 NOLOGY, CHENGDU, SICHUAN, 610225, P. R. CHINA
367 *E-mail address:* `wangchangyou417@163.com`