



**HAL**  
open science

# Inside dynamics for stage-structured integrodifference equations

Nathan Marculis, Jimmy Garnier, Roger Lui, Mark Lewis

► **To cite this version:**

Nathan Marculis, Jimmy Garnier, Roger Lui, Mark Lewis. Inside dynamics for stage-structured integrodifference equations. *Journal of Mathematical Biology*, 2019, 10.1007/s00285-019-01378-9 . hal-02367984v1

**HAL Id: hal-02367984**

**<https://hal.science/hal-02367984v1>**

Submitted on 22 Dec 2020 (v1), last revised 21 Jan 2021 (v2)

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

# 1 Inside Dynamics for Stage-Structured Integrodifference Equations

2 Nathan G. Marculis · Jimmy Garnier · Roger Lui ·  
3 Mark A. Lewis

4  
5 Received: date / Accepted: date

6 **Abstract** A stage-structured model of integrodifference equations is used to study the asymptotic neutral  
7 genetic structure of populations undergoing range expansion. That is, we study the inside dynamics of  
8 solutions to stage-structured integrodifference equations. To analyze the genetic consequences for long term  
9 population spread, we decompose the solution into neutral genetic components called neutral fractions.  
10 The inside dynamics are then given by the spatiotemporal evolution of these neutral fractions. We show  
11 that, under some mild assumptions on the dispersal kernels and population projection matrix, the spread  
12 is dominated by individuals at the leading edge of the expansion. This result is consistent with the founder  
13 effect. In the case where there are multiple neutral fractions at the leading edge we are able to explicitly cal-  
14 culate the asymptotic proportion of these fractions found in the long-term population spread. This formula  
15 is simple and depends only on the right and left eigenvectors of the population projection matrix evaluated  
16 at zero and the initial proportion of each neutral fraction at the leading edge of the range expansion. In  
17 the absence of a strong Allee effect, multiple neutral fractions can drive the long-term population spread,  
18 a situation not possible with the scalar model.

19 **Keywords** integrodifference equations · stage-structure · inside dynamics · neutral genetic structure ·  
20 founder effect

21 **Mathematics Subject Classification (2000)** 39A10 · 45G10 · 92D25 · 92D40

## 22 1 Introduction

23 There are a wide array of observational (Cullingham et al. 2011), empirical (Liebhold et al. 1992; Lubina  
24 and Levin 1988), and theoretical studies (Li et al. 2009; Lui 1989a; Weinberger 1982) for the spatial spread  
25 of populations by range expansion. Over the last decades, theoretical studies about range expansion mainly

---

This research was supported by a grant to MAL from the Natural Science and Engineering Research Council of Canada (Grant No. NET GP 434810-12) to the TRIA Network, with contributions from Alberta Agriculture and Forestry, Foothills Research Institute, Manitoba Conservation and Water Stewardship, Natural Resources Canada-Canadian Forest Service, Northwest Territories Environment and Natural Resources, Ontario Ministry of Natural Resources and Forestry, Saskatchewan Ministry of Environment, West Fraser and Weyerhaeuser. MAL is also grateful for support through NSERC and the Canada Research Chair Program. NGM acknowledges support from NSERC TRIA-Net Collaborative Research Grant. JG acknowledges NONLOCAL project (ANR-14-CE25-0013) and the European Research Council (ERC) under the European Unions Horizon 2020 research and innovation programme (grant agreement No 639638, MesoProbio). The authors also gratefully acknowledge the helpful suggestions from the anonymous reviewers.

---

Nathan G. Marculis  
Department of Mathematical and Statistical Sciences, University of Alberta, Edmonton, AB T6G 2G1, Canada  
E-mail: marculis@ualberta.ca

Jimmy Garnier  
Université Savoie MontBlanc, CNRS, LAMA, F-73000 Chambéry, France

Roger Lui  
Department of Mathematical Sciences, Worcester Polytechnic Institute, Worcester, MA 01609, USA

Mark A. Lewis  
Department of Mathematical and Statistical Sciences, University of Alberta, Edmonton, AB T6G 2G1, Canada  
Department of Biological Sciences, University of Alberta, Edmonton, AB T6G 2G1, Canada

26 focused on the asymptotic speed of propagation of the expanding population or the profile of invasion  
 27 (Hastings et al. 2005). Spatial models in population genetics have also been developed for studying the  
 28 spread of an advantageous gene in a population (Lui 1982a,b, 1983; Weinberger 1978, 1982). Recently,  
 29 much effort have been invested to understand the genetic consequences of range expansion (Hallatschek  
 30 and Nelson 2008; Roques et al. 2012). Indeed, range expansions are known to have significant effects on  
 31 genetic diversity (Davis and Shaw 2001; Hewitt 2000). For instance, if range expansion occurs through  
 32 successive founder effects, genetic diversity is likely to decrease. However, empirical and theoretical studies  
 33 have shown that many mechanisms may reduce or reverse the loss of diversity in an expanding population  
 34 (Pluess 2011). In particular, the presence of an Allee effect (Roques et al. 2012) which reduces the per-capita  
 35 growth rate at low density, the occurrence of long distance dispersal events (Bonnefon et al. 2014; Ibrahim  
 36 et al. 1996), or the existence of a juvenile stage (Austerlitz and Garnier-Géré 2003) may promote neutral  
 37 genetic diversity in traveling waves of colonization. In this work, we are interested in the neutral genetic  
 38 dynamics of a stage-structured population undergoing range expansion.

39 It is well known that the structure of the population is important for understanding the asymptotic  
 40 dynamics. For example, individuals often must undergo a maturation period before they can produce off-  
 41 spring. For discrete population models, the dynamics of the life history traits have typically been structured  
 42 according to age, Leslie matrix (Leslie 1945), or developmental stage, Lefkovich matrix (Lefkovich 1965),  
 43 but matrix models can be easily generalized to include other physiological characteristics. It is also com-  
 44 mon for sessile species to typically have a motile stage in their development, such as seed dispersal in plant  
 45 populations (Howe and Smallwood 1982) and larval dispersal in marine environments (Levin 2006).

46 Our study considers a stage-structured integrodifference equation describing range expansion for a  
 47 population of the form:

$$\mathbf{u}_{t+1}(x) = \int_{-\infty}^{\infty} [\mathbf{K}(x-y) \circ \mathbf{B}(\mathbf{u}_t(y))] \mathbf{u}_t(y) dy, \quad (1)$$

48 where  $\mathbf{u}_t(x)$  corresponds to the population density at time  $t$  and location  $x$ . The population is structured  
 49 into  $m$  stages, whose densities are given by  $\mathbf{u}_t(x) = [u_{1,t}(x), \dots, u_{m,t}(x)]$ . Each stage distribution changes  
 50 in time and space through the successive effects of dispersal, described by the dispersal matrix  $\mathbf{K} = [k_{jl}]$ ,  
 51 and the demography, embodied in the population projection matrix  $\mathbf{B}(\mathbf{u}) = [b_{jl}(\mathbf{u})]$  which takes into  
 52 account density-dependence. The succession of the reproduction stage and dispersal stage is described by  
 53 the Hadamard product  $\circ$  (element-wise multiplication of matrix). This model allows the different stages to  
 54 spread, reproduce, and interact in a variety of ways that cannot be captured by scalar models (Neubert and  
 55 Caswell 2000). More precisely, if we consider stage  $j$ , where  $j = 1, \dots, m$ , then its density,  $u_{j,t}(x)$ , satisfies  
 56 the following equation

$$u_{j,t+1} = \int_{-\infty}^{\infty} \sum_{l=1}^m k_{jl}(x-y) b_{jl}(u_{1,t}(y), \dots, u_{m,t}(y)) u_{l,t}(y) dy \quad (2)$$

57 where  $k_{jl}(x-y) dy$  is the probability that an individual transitioning from stage  $l$  to stage  $j$  disperses from  
 58 the interval  $(y, y + dy]$  to location  $x$ , and the function  $b_{jl}$  is the per-capita production of stage  $j$  individuals  
 59 from stage  $l$  individuals. Such a model has been used to describe epidemic spread (Lui 1989b), biological  
 60 invasions (Bateman et al. 2017; Veit and Lewis 1996), and critical domain size (Lutscher and Lewis 2004).

61 The model (1) is biologically valid if the stages are chosen in a way such that the life history and  
 62 dispersal parameters vary within stages as little as possible. In some cases this is easy; for example, a  
 63 division between juvenile and adult individuals is normally determined by the ability to reproduce. In other  
 64 cases, the division may not be so clear, and partitions may be difficult to decide. Fortunately, there are  
 65 algorithms that can be used to minimize errors associated with partitioning a population into distinct stages  
 66 (Moloney 1986; Vandermeer 1978). If the division of population structure is modeled using a continuous  
 67 variable such as size or mass, and there is no natural break point to structure the population into distinct  
 68 stages then an integral projection model may be more appropriate (Easterling et al. 2000).

69 The goal of this work is to understand the neutral genetic patterns of structured populations. Neutral  
 70 genetic markers are genes that have no direct effect on individual fitness. Even though this type of gene  
 71 tells us nothing about the adaptive or evolutionary potential of a population, neutral genetic markers can  
 72 be used to understand processes such as gene flow, genetic drift, migration, or dispersal (Holderegger et al.  
 73 2006). It has also been shown by simulations that high levels of neutral genetic diversity can be correlated  
 74 with increased allelic richness at loci under selection (Bataillon et al. 1996). Our analysis will be focused  
 75 on the inside dynamics of stage-structured integrodifference equations.

76 This paper is organized as follows. Section 2 is dedicated to providing necessary background material  
 77 for understanding the main results. Within this section, we break it into two subsections: Section 2.1

78 provides background to the analysis of inside dynamics and the stage-structured integrodifference equation  
79 used in our analysis and Section 2.2 lays out four of the major assumptions made about the demographic  
80 and dispersal processes. In Section 3, we provide asymptotic results regarding population structure. This  
81 section is broken into three parts. Section 3.1 covers the inside dynamics of neutral fractions not present  
82 at the leading edge, Section 3.2 discusses the inside dynamics of neutral fractions that are located at the  
83 leading edge, and Section 3.3 contains proofs for our main theorems. To complement the analytical results,  
84 numerical simulations are given in Section 4. Finally, in Section 5, we discuss the modeling technique,  
85 results, numerical simulations, and implications of our work.

## 86 **2 Materials and methods**

### 87 2.1 Inside dynamics

88 To study the neutral genetic distribution of a population, we consider the inside dynamics of the population.  
89 The term inside dynamics refers to the inside structure of the population rather than the total density.  
90 The key assumption in the analysis of inside dynamics is that all individuals grow and disperse in the  
91 same manner, differing only with respect to neutral genetic markers. In other words, all individuals in the  
92 population have the same fitness. This allows us to split up the population into distinct subgroups called  
93 neutral fractions with which we track the spatiotemporal evolution of these subgroups.

94 Inside dynamics have been studied for reaction-diffusion equations (Garnier and Lewis 2016; Garnier  
95 et al. 2012; Roques et al. 2012), delay reaction-diffusion equations (Bonnefon et al. 2013), integro-differential  
96 equations (Bonnefon et al. 2014), and integrodifference equations (Lewis et al. 2018; Marculis et al. 2017).  
97 In these works, the subject for analysis was a scalar population model. Indeed, to date, there is only one  
98 study of the inside dynamics of systems of equations. This study concentrated on the analysis on a diffusive  
99 Lotka-Volterra competition system (Roques et al. 2015). Our mathematical contribution to this area of  
100 research is to extend the analysis of inside dynamics to stage-structured integrodifference equations.

101 Recall the stage-structured population model in (1). Separating the initial population up into distinct  
102 neutral fractions, we obtain the initial condition

$$\mathbf{u}_0(x) = \sum_{i=1}^n \mathbf{v}_0^i(x), \quad (3)$$

103 where  $\mathbf{v}_0^i(x) \geq 0$  is the initial population density for neutral fraction  $i$  and  $n$  is the finite number of neutral  
104 fractions. An illustration of this decomposition can be seen in Figures 1(a) and 2(a). By assuming that  
105 individuals in each neutral fraction grow and disperse similarly, we obtain the following system of equations:

$$\mathbf{v}_{t+1}^i(x) = \int_{-\infty}^{\infty} [\mathbf{K}(x-y) \circ \mathbf{B}(\mathbf{u}_t(y))] \mathbf{v}_t^i(y) dy, \quad i = 1, \dots, n, \quad (4)$$

106 where  $\mathbf{u}_t(y) = \sum_{i=1}^n \mathbf{v}_t^i(y)$ . Throughout the remaining sections, we use the superscript  $i$  to denote the  
107 neutral fraction and, when not written in vector form, subscript  $j$  to denotes the stage. Note that the  
108 number of neutral fractions,  $n$ , and the number of stages in the population,  $m$ , need not be the same  
109 ( $n \neq m$ ). Also, observe the model given in Equation (4) is natural extension of the scalar model to a system  
110 of recursions (Marculis et al. 2017). Thus, it can be expected that many of the results proven for the scalar  
111 equation can be extended to systems of cooperative equations. This is the approach we take in what follows.

### 112 2.2 Demographic and dispersal assumptions

113 For each of our main theorems, we make five assumptions regarding Equations (3)-(4). The first three as-  
114 sumptions are related to the population projection matrix, the fourth assumption is related to the dispersal  
115 kernel, and the fifth and final assumption is related to the decay of the initial conditions. In this section,  
116 we outline the first four assumptions related to the demography and dispersal of the population.

117 *Population projection matrix*

118 We begin with looking at the population projection matrix  $\mathbf{B}(\mathbf{u})$ . Here, we outline three assumptions about  
 119 the population projection matrix. The population projection matrix describes reproduction, survival, and  
 120 interactions between stages. As a projection matrix, its entries should be nonnegative:

121 *A1 : The matrix  $\mathbf{B}(\mathbf{u})$  is nonnegative for any  $\mathbf{u} \in (0, \infty)^m$ .*

122 Moreover, we can see from (1) that  $\mathbf{0}$  is a steady state of the problem. Define

$$\mathbf{B}_0 := \mathbf{B}(\mathbf{u})|_{\mathbf{u}=\mathbf{0}}. \quad (5)$$

123 Notice that  $\mathbf{B}_0$  is the population projection matrix evaluated at  $\mathbf{u} = \mathbf{0}$ . We will assume that this steady  
 124 state is unstable. More precisely, we assume:

125 *A2 :  $\mathbf{B}_0$  is a primitive matrix, that is there exists  $k > 0$  such that  $\mathbf{B}_0^k$  is positive, and its dominant  
 126 eigenvalue,  $\lambda_1$ , is greater than 1,  $\lambda_1 > 1$ .*

127 Finally, we assume that there are no Allee effects. That is:

128 *A3 :  $\mathbf{B}(\mathbf{u})$  is bounded by its linearization at the steady state  $\mathbf{0}$ ,  $\mathbf{B}(\mathbf{u})\mathbf{v} \leq \mathbf{B}_0\mathbf{v}$  for all  $\mathbf{v} \in (0, \infty)^m$ .*

129 *Dispersal kernel*

130 In our model, we assume that individuals in the population may disperse at long distance but those events  
 131 are rare in the following sense:

132 **Definition 1** *A dispersal kernel,  $k(x)$ , is called thin-tailed if there exists a  $\xi > 0$ , such that*

$$\int_{-\infty}^{\infty} k(x)e^{\xi|x|} dx < \infty. \quad (6)$$

133 A dispersal kernel that is not thin-tailed is called a fat-tailed dispersal kernel, and in this case, the long  
 134 distance dispersal events become frequent, which leads to different behaviors for some solutions, such as  
 135 accelerating waves. Many of the classical mathematical results for (1), such as traveling wave solutions  
 136 and the asymptotic speed of propagation, rely on the assumption that the dispersal kernel is thin-tailed. A  
 137 common dispersal kernel that we consider throughout our work is the Gaussian probability density function:

$$k(x; \mu, \sigma) = \frac{1}{\sqrt{2\pi\sigma^2}} e^{-\frac{(x-\mu)^2}{2\sigma^2}}, \quad (7)$$

138 where  $\mu$  is the mean shift in location and  $\sigma^2$  is the variance in dispersal distance. In the following sections,  
 139 we use the following shorthand notation to denote that the dispersal kernel is Gaussian by  $k$  is  $N(\mu, \sigma^2)$ .  
 140 In what follows, we will make one of two assumptions about the dispersal kernels.

141 *A4 : Each kernel,  $k_{ji}(x - y)$ , is thin-tailed.*

142 *A4' : Each kernel,  $k_{ji}(x - y)$ , is  $N(\mu, \sigma^2)$ .*

143 From above, we see that our fourth assumption provides a condition on the dispersal kernels. In both cases,  
 144 we assume, at a minimum, that every dispersal kernel is thin-tailed in order to calculate the asymptotic speed  
 145 of propagation. The above assumption implies that we are not considering a population with accelerating  
 146 waves (Kot et al. 1996).

147 *Asymptotic speed of propagation*

148 Under the previous assumptions A1-A4 we can deduce from the work of Lui (1989a) that solutions of (1)  
 149 will spread to the right with an asymptotic spreading speed  $c$  greater than or equal to a critical spreading  
 150 speed  $c^* > 0$  for appropriately chosen initial conditions. Moreover, the critical spreading speed  $c^*$  can be  
 151 computed explicitly by the following formula

$$c^* := \min_{0 < s < s^+} \frac{1}{s} \ln \rho(s), \quad (8)$$

152 where  $\rho(s) := \rho(\mathbf{H}(s)) > 1$  is the dominant eigenvalue of  $\mathbf{H}(s)$  defined by

$$\mathbf{H}(s) := \mathbf{M}(s) \circ \mathbf{B}_0. \quad (9)$$

153 The moment generating function matrix  $\mathbf{M}(s)$  is calculated by applying the reflected bilateral Laplace  
 154 transform to the dispersal kernel matrix  $\mathbf{K}$  and is defined by

$$\mathbf{M}(s) := \int_{-\infty}^{\infty} \mathbf{K}(x)e^{sx} dx. \quad (10)$$

155 Since the entries of the dispersal kernel matrix,  $k_{jl}$ , are thin-tailed by Assumption A5, this matrix is well  
 156 defined over  $(0, s^+)$  where  $s^+ \in (0, \infty]$ . Throughout our analysis, we let  $s_0(c)$  be the smallest positive root  
 157 of the equation

$$cs = \ln(\rho(s)) \text{ for } c \geq c^*. \quad (11)$$

158 We know that  $s_0(c)$  exists because  $\rho(s)$  is log convex; see Lemma 6.4 by Lui (1989a). In particular, when  
 159 each kernel is Gaussian,  $k_{jl}$  is  $N(\mu, \sigma^2)$ , then we have an explicit formula for the asymptotic speed of  
 160 propagation given by

$$c^* = \sqrt{2\sigma^2 \ln(\lambda_1)} + \mu, \quad (12)$$

161 where  $\lambda_1$  is the dominant eigenvalue of  $\mathbf{B}_0$ . The technical details for the asymptotic speed of propagation  
 162 are provided in Appendix A.

### 163 3 Main results

164 Henceforth, we assume that the structured population,  $\mathbf{u}_t(x)$ , satisfies (1) with an initial condition  $\mathbf{u}_0(x)$ .  
 165 With such initial condition, the population is spreading to the right with an asymptotic speed of propaga-  
 166 tion,  $c$ , greater than or equal to  $c^*$ , given by formula (8). We first consider neutral fractions that are not  
 167 present at the leading edge of the solution and then afterwards consider neutral fractions that are at the  
 168 leading edge of the expanding population.

169 Our fifth and final assumption places a requirement on the initial conditions for the neutral fractions.  
 170 This requirement is closely connected to the decay rate of the solution for the population and determines  
 171 whether or not an individual is at the leading edge of the population spread. The technical details of whether  
 172 or not a neutral fraction is located at the leading edge is defined in the statement of our main theorems.  
 173 We do not explicitly write these out here, but rather save them for the statement of our theorems because  
 174 this assumption takes different forms based on our assumptions. We are now ready to present our first two  
 175 theorems, that provides sufficient conditions for when the density of neutral fractions converges to zero in  
 176 the moving half-frame.

#### 177 3.1 Inside dynamics not at the leading edge

178 **Theorem 3.1** *Let us assume that A1-A4 hold true. Let  $\mathbf{v}_t^i(x)$  be a neutral fraction satisfying (4) with*  
 179 *initial condition  $\mathbf{v}_0^i(x)$  satisfying (3) that is not present at the leading edge of the expanding population, in*  
 180 *the sense that*

181  $A5 : x^2 \mathbf{v}_0^i(x) e^{s_0(c)x} \in L^1(\mathbb{R}) \cap L^\infty(\mathbb{R}) \text{ for a given } c \geq c^* .$

182 *Then, for any  $A \in \mathbb{R}$ , the density of neutral fraction  $i$ ,  $\mathbf{v}_t^i(x)$ , converges to  $\mathbf{0}$  uniformly as  $t \rightarrow \infty$  in the*  
 183 *moving half-frame  $[A + ct, \infty)$ .*

184 In summary, Theorem 3.1 provides sufficient conditions for the neutral fractions in the population to  
 185 approach zero asymptotically. As stated in the theorem, Assumption A5 corresponds to an initial condition  
 186 for the neutral fractions that are not present at the leading edge of the population. This result implies that  
 187 the only neutral fractions that will contribute to the spread of the population are those that are initially  
 188 at the leading edge. In this scenario, we observe an extreme founder effect for the population spread. For  
 189 this proof, see Section 3.3.

190 By making a stronger assumption on the dispersal kernels, we are able to relax Assumption A5 on the  
 191 initial conditions in Theorem 3.1. In particular, for the next theorem we assume that all dispersal kernels  
 192 are Gaussian with the same mean and variance as given by Assumption A4'. This allows us to relax the  
 193 assumptions on the initial conditions to a much simpler integrability assumption.

194 **Theorem 3.2** *Let us assume that A1-A3 and A4' hold true. Let  $\mathbf{v}_t^i(x)$  be a neutral fraction satisfying (4)*  
 195 *with initial condition  $\mathbf{v}_0^i(x)$  satisfying (3) that is not present at the leading edge of the expanding population,*  
 196 *in the sense that*

197  $A5' : \int_{-\infty}^{\infty} e^{\frac{c-\mu}{\sigma^2}y} \mathbf{v}_0^i(y) dy < \infty$  for a given  $c \geq c^*$ .

198 Then, for any  $A \in \mathbb{R}$ , the density of neutral fraction  $i$ ,  $\mathbf{v}_t^i(x)$ , converges to  $\mathbf{0}$  uniformly as  $t \rightarrow \infty$  in the  
199 moving half-frame  $[A + ct, \infty)$ .

200 In summary, Theorem 3.2 provides the same result as Theorem 3.1 but with different assumptions on  
201 the dispersal kernels and initial conditions. That is, Theorem 3.2 provides sufficient conditions for when  
202 the neutral fractions do not contribute to the population spread. Under Assumption  $A5'$ , we see that the  
203 leading edge is determined by the decaying exponential  $e^{-\frac{c-\mu}{\sigma^2}x}$ . This condition is much different than those  
204 given by Assumption  $A5$  in Theorem 3.1. As in the previous theorem, we also observe here that the only  
205 neutral fractions that will contribute to the spread of the population are those that are initially at the  
206 leading edge. For this proof, see Section 3.3.

207 The proof of Theorem 3.1 is more complicated than that of Theorem 3.2, even though the method of  
208 proof and conclusions are the same. The difference is due to the assumptions made about the dispersal  
209 kernels. In Theorem 3.1 we assume the dispersal kernels are thin-tailed and must use the definition of the  
210 inverse reflected bilateral Laplace transform. In Theorem 3.2 we assume all dispersal kernels are Gaus-  
211 sian with the same mean and variance. This assumption simplifies the proof because convolving Gaussian  
212 distributions results in another Gaussian.

213 If the initial conditions are all compactly supported, then all neutral fractions will satisfy Assumption  
214  $A5$  and  $A5'$  respectively in Theorems 3.1 and 3.2. If the initial conditions decay according to the traveling  
215 wave solution, then all neutral fractions except those at the leading edge will satisfy Assumption  $A5$  and  
216  $A5'$  in Theorems 3.1 and 3.2 respectively. This means that the only neutral fractions that we will see in  
217 the moving half-frame are those that were initially at the leading edge. However, Theorems 3.1 and 3.2 do  
218 not tell us anything about these neutral fractions.

### 219 3.2 Inside dynamics at the leading edge

220 In the next theorem, we look at initial data that decay slower than Assumption  $A5'$  in Theorem 3.2. Here  
221 we are able to calculate the asymptotic proportion of each neutral fraction provided we move at the slowest  
222 speed  $c^*$ .

223 **Theorem 3.3** *Let us assume that  $A1$ - $A3$  and  $A4'$  hold true. Let  $\mathbf{v}_t^i(x)$  be a neutral fraction satisfying (4)  
224 with initial condition  $\mathbf{v}_0^i(x)$  satisfying (3) that is present at the leading edge of the expanding population,  
225 in the sense that for  $c = c^*$*

226  $A5'' : \mathbf{v}_0^i(x) = (\mathbf{p}_0^i \circ \mathbf{r}) e^{-\frac{c-\mu}{\sigma^2}x}$ , where  $\mathbf{p}_0^i$  is the initial proportion for neutral fraction  $i$  in each stage,  $\mathbf{r}$  is  
227 the right eigenvector of  $\mathbf{B}_0$  corresponding to  $\lambda_1$ .

228 Then, for any  $A \in \mathbb{R}$ , the density of neutral fraction  $i$ ,  $\mathbf{v}_t^i(x)$ , asymptotically approaches a proportion,  $p^i$ ,  
229 of the traveling wave for the linear equation as  $t \rightarrow \infty$  in the moving half-frame  $[A + ct, \infty)$ . That is,

$$\lim_{t \rightarrow \infty} \mathbf{v}_t^i(x_0 + ct) = e^{-\frac{c-\mu}{\sigma^2}x_0} \mathbf{r} \mathbf{p}^i \quad (13)$$

230 for  $x_0 \geq A$ . Moreover, the proportion can be calculated to be the scalar

$$p^i = \ell \left( \mathbf{p}_0^i \circ \mathbf{r} \right) \quad (14)$$

231 where  $\ell$  is the left eigenvector of  $\mathbf{B}_0$  corresponding to  $\lambda_1$  with  $\ell$  normalized by  $\langle \ell^T, \mathbf{r} \rangle$ .

232 Theorem 3.3 provides a formula for the asymptotic proportion of neutral fractions based on the initial  
233 distribution at the leading edge of the population. The formula is simple because it depends only on the  
234 right and left eigenvectors of  $\mathbf{B}_0$  and the initial proportion of neutral fractions. This theorem characterizes  
235 the fate of neutral fractions at the leading edge. One drawback to this theorem is that it is only valid  
236 for initial conditions that decay at a specific rate,  $e^{-\frac{c-\mu}{\sigma^2}x}$ , with a solution that moves at a specific speed,  
237  $c = c^*$ . The reason why we cannot prove this theorem for  $c > c^*$  and a slower decay rate for the initial  
238 condition is because we do not have an explicit formula for the spreading speed  $c > c^*$ . For this proof, see  
239 Section 3.3.

240 It is also important to note that  $A5''$  in Theorem 3.3 is not completely biologically realistic since the  
241 population grows without bound as  $x \rightarrow -\infty$ . However, this type of initial condition is needed based  
242 on the construction of our sub-solution and super-solutions. It may be possible to relax this assumption  
243 by studying the nonlinear operator and considering a more biologically realistic class of initial conditions.  
244 Next, we present the proofs of Theorems 3.1-3.3 in Section 3.3. For a comprehensive review of the necessary  
245 mathematical material needed in the proofs of the theorems, we direct the reader to Appendix B.

247 *Proof of Theorem 3.1*248 *Proof* For simplicity, we drop the superscript  $i$  in (4) and focus on a single neutral fraction. Our equation  
249 of interest is

$$\mathbf{v}_{t+1}(x) = \int_{-\infty}^{\infty} [\mathbf{K}(x-y) \circ \mathbf{B}(\mathbf{u}_t(y))] \mathbf{v}_t(y) dy. \quad (15)$$

250 Let

$$\mathbf{w}_0(x) = \frac{\mathbf{C}e^{-s_0(c)x}}{1+x^2} \quad (16)$$

251 where  $\mathbf{C} = \kappa\phi$  and  $\phi$  is the eigenvector of  $\mathbf{H}(s_0(c))$  with dominant eigenvalue  $\rho(s_0(c))$ . From Lemma B.1,  
252 we know that  $\mathbf{w}_0(x)$  is an upper bound for  $\mathbf{v}_0(x)$ . By Assumption A3, we know that  $\mathbf{B}(\mathbf{u}_t(y))\mathbf{v} \leq \mathbf{B}_0\mathbf{v}$  for  
253 all  $\mathbf{v} \geq 0$ . Hence, we can construct a super-solution  $\mathbf{w}_t(x)$  that satisfies the following equation

$$\mathbf{w}_{t+1}(x) = \int_{-\infty}^{\infty} [\mathbf{K}(x-y) \circ \mathbf{B}_0] \mathbf{w}_t(y) dy \quad (17)$$

254 with initial condition given by (16). By iterating we can write the solution to the above system as the  $t$ -fold  
255 convolution

$$\mathbf{w}_t(x) = [\mathbf{K}(x-y) \circ \mathbf{B}_0]^{*t} \mathbf{w}_0(y). \quad (18)$$

256 Applying the bilateral Laplace transform

$$\mathbf{W}_t(s) = [\mathbf{M}(s) \circ \mathbf{B}_0]^t \mathbf{W}_0(s) \quad (19)$$

$$= [\mathbf{H}(s)]^t \mathbf{W}_0(s). \quad (20)$$

257 Recall that  $s_0(c)$  is the smallest positive root of  $sc = \ln(\rho(s))$  for  $c \geq c^*$ . Then, the inverse transform as  
258 defined in Appendix B, see (129), yields

$$\mathbf{w}_t(x) = \frac{1}{2\pi i} \lim_{R \rightarrow \infty} \int_{s_0(c)-iR}^{s_0(c)+iR} [\mathbf{H}(s)]^t \mathbf{W}_0(s) e^{-sx} ds \quad (21)$$

$$= \frac{1}{2\pi} \int_{-\infty}^{\infty} [\mathbf{H}(s_0(c) + i\omega)]^t \mathbf{W}_0(s_0(c) + i\omega) e^{-(s_0(c)+i\omega)x} d\omega \quad (22)$$

259 for  $c \geq c^*$ . In the moving frame we have

$$\mathbf{w}_t(x_0 + ct) = \frac{1}{2\pi} \int_{-\infty}^{\infty} [\mathbf{H}(s_0(c) + i\omega)]^t \mathbf{W}_0(s_0(c) + i\omega) e^{-(s_0(c)+i\omega)x_0} e^{-(s_0(c)+i\omega)ct} d\omega. \quad (23)$$

260 Using the results from Lemma B.1, see Appendix B for details, we are able to write the initial condition in  
261 terms of a Fourier transform that is known. This is seen as follows,

$$\mathbf{W}_0(s_0(c) + i\omega) = \int_{-\infty}^{\infty} \mathbf{w}_0(x) e^{(s_0(c)+i\omega)x} dx \quad (24)$$

$$= \int_{-\infty}^{\infty} \mathbf{w}_0(x) e^{s_0(c)x} e^{i\omega x} dx \quad (25)$$

$$= \mathcal{F}[\mathbf{w}_0(x) e^{s_0(c)x}](-\omega) \quad (26)$$

$$= \mathbf{C}\pi e^{-|\omega|} \quad (27)$$

262 for all  $\omega \in \mathbb{R}$ . Recall that  $\mathbf{C} = \kappa\phi$ . This gives

$$\mathbf{w}_t(x_0 + ct) = \frac{1}{2\pi} \int_{-\infty}^{\infty} [\mathbf{H}(s_0(c) + i\omega)]^t \mathbf{C}\pi e^{-|\omega|} e^{-(s_0(c)+i\omega)x_0} e^{-(s_0(c)+i\omega)ct} d\omega \quad (28)$$

$$= \frac{1}{2} \int_{-\infty}^{\infty} [\mathbf{H}(s_0(c) + i\omega)]^t \kappa e^{-s_0(c)ct} \phi e^{-|\omega|} e^{-(s_0(c)+i\omega)x_0} e^{-i\omega ct} d\omega. \quad (29)$$



263 Since  $s_0(c)c = \ln(\rho(s_0(c)))$ , we have

$$\mathbf{w}_t(x_0 + ct) = \frac{\kappa e^{-s_0(c)x_0}}{2} \int_{-\infty}^{\infty} [\mathbf{H}(s_0(c) + i\omega)]^t e^{-\ln(\rho(s_0(c)))t} \phi e^{-|\omega|} e^{-i\omega x_0} e^{-i\omega ct} d\omega \quad (30)$$

$$= \frac{\kappa e^{-s_0(c)x_0}}{2} \int_{-\infty}^{\infty} [\mathbf{H}(s_0(c) + i\omega)]^t (\rho(s_0(c)))^{-t} \phi e^{-|\omega|} e^{-i\omega x_0} e^{-i\omega ct} d\omega. \quad (31)$$

264 Since  $\rho(s_0(c))$  is the dominant eigenvalue of  $\mathbf{H}(s_0(c))$  with eigenvector  $\phi$ ,

$$\mathbf{w}_t(x_0 + ct) = \frac{\kappa e^{-s_0(c)x_0}}{2} \int_{-\infty}^{\infty} [\mathbf{H}(s_0(c) + i\omega)]^t [\mathbf{H}(s_0(c))]^{-t} \phi e^{-|\omega|} e^{-i\omega x_0} e^{-i\omega ct} d\omega. \quad (32)$$

265 Applying the matrix norm and using the sub-additive property, we find that

$$\|\mathbf{w}_t(x_0 + ct)\| \leq \frac{\kappa e^{-s_0(c)x_0}}{2} \int_{-\infty}^{\infty} \|[\mathbf{H}(s_0(c) + i\omega)]^t\| \|[\mathbf{H}(s_0(c))]^{-t}\| \|\phi\| e^{-|\omega|} |e^{-i\omega x_0}| |e^{-i\omega ct}| d\omega \quad (33)$$

$$= \frac{\kappa e^{-s_0(c)x_0}}{2} \int_{-\infty}^{\infty} \|[\mathbf{H}(s_0(c) + i\omega)]^t\| \|[\mathbf{H}(s_0(c))]^{-t}\| \|\phi\| e^{-|\omega|} d\omega. \quad (34)$$

266 We can also see that

$$|\mathbf{H}(s_0(c) + i\omega)| = |\mathbf{M}(s_0(c) + i\omega) \circ \mathbf{B}_0| \quad (35)$$

$$= \left| \int_{-\infty}^{\infty} [\mathbf{K}(x) \circ \mathbf{B}_0] e^{(s_0(c) + i\omega)x} dx \right| \quad (36)$$

$$= \left| \int_{-\infty}^{\infty} [\mathbf{K}(x) \circ \mathbf{B}_0] e^{s_0(c)x} (\cos(\omega x) + i \sin(\omega x)) dx \right| \quad (37)$$

$$= I, \quad (38)$$

267 where  $I$  is defined to be

$$I := \sqrt{\left( \int_{-\infty}^{\infty} [\mathbf{K}(x) \circ \mathbf{B}_0] e^{s_0(c)x} \cos(\omega x) dx \right)^2 + \left( \int_{-\infty}^{\infty} [\mathbf{K}(x) \circ \mathbf{B}_0] e^{s_0(c)x} \sin(\omega x) dx \right)^2}. \quad (39)$$

268 By the Cauchy-Schwarz inequality, using a similar technique as in Theorem 3 of (Marculis et al. 2017),

$$I < \int_{-\infty}^{\infty} [\mathbf{K}(x) \circ \mathbf{B}_0] e^{s_0(c)x} dx \quad (40)$$

$$= \mathbf{M}(s_0(c)) \circ \mathbf{B}_0 \quad (41)$$

$$= \mathbf{H}(s_0(c)) \quad (42)$$

269 for  $\omega \neq 0$ . From the above calculation we can conclude that  $|\mathbf{H}(s_0(c) + i\omega)| < \mathbf{H}(s_0(c))$  for  $\omega \neq 0$ .

270 Consequently,  $\rho(|\mathbf{H}(s_0(c) + i\omega)|) < \rho(s_0(c))$  for  $\omega \neq 0$ . By Gelfand's formula,

$$\lim_{t \rightarrow \infty} \|[\mathbf{H}(s_0(c) + i\omega)]^t\|^{\frac{1}{t}} = \rho(|\mathbf{H}(s_0(c) + i\omega)|) \quad \text{and} \quad (43)$$

$$\lim_{t \rightarrow \infty} \|[\mathbf{H}(s_0(c))]^{-t}\|^{\frac{1}{t}} = \frac{1}{\rho(s_0(c))}. \quad (44)$$

271 Thus, for  $\omega \neq 0$ , we can choose  $\varepsilon > 0$  such that  $(\rho(|\mathbf{H}(s_0(c) + i\omega)|) + \varepsilon) \left( \frac{1}{\rho(s_0(c))} + \varepsilon \right) < 1$ . Therefore,

$$\|[\mathbf{H}(s_0(c) + i\omega)]^t\| \|[\mathbf{H}(s_0(c))]^{-t}\| < 1 \quad (45)$$

272 for large  $t$  and

$$\lim_{t \rightarrow \infty} \|[\mathbf{H}(s_0(c) + i\omega)]^t\| \|[\mathbf{H}(s_0(c))]^{-t}\| = 0. \quad (46)$$

273 From (34) and the dominated convergence theorem,

$$\lim_{t \rightarrow \infty} \|\mathbf{w}_t(x_0 + ct)\| \leq \frac{\kappa e^{-s_0(c)x_0}}{2} \int_{-\infty}^{\infty} \lim_{t \rightarrow \infty} \|[\mathbf{H}(s_0(c) + i\omega)]^t\| \|[\mathbf{H}(s_0(c))]^{-t}\| \|\phi\| e^{-|\omega|} d\omega \quad (47)$$

$$= \mathbf{0}. \quad (48)$$

274 Therefore, for any  $A \in \mathbb{R}$  and  $c \geq c^*$ ,

$$\lim_{t \rightarrow \infty} \max_{[A, \infty)} \mathbf{w}_t(x + ct) = \mathbf{0}. \quad (49)$$

275 Since  $\mathbf{w}$  was constructed as a super-solution, we can conclude that

$$\lim_{t \rightarrow \infty} \max_{[A, \infty)} \mathbf{v}_t(x + ct) = \mathbf{0}. \quad (50)$$

The proof of Theorem 3.1 is complete.  $\square$

276 *Proof of Theorem 3.2*

277 *Proof* For simplicity, we focus on a single neutral fraction and drop the superscript  $i$ . By Assumption A3,  
278  $\mathbf{B}(\mathbf{u}_t(y))\mathbf{v} \leq \mathbf{B}_0\mathbf{v}$  for all  $\mathbf{v} \geq 0$ , we can use a comparison principle to show that a new sequence  $\mathbf{w}_t(x)$   
279 defined by

$$\mathbf{w}_{t+1}(x) = \int_{-\infty}^{\infty} [\mathbf{K}(x-y) \circ \mathbf{B}_0] \mathbf{w}_t(y) dy \quad (51)$$

280 is always greater than the solution to any neutral fraction  $\mathbf{v}_t(x)$  with the same initial condition,  $\mathbf{w}_0(x) =$   
281  $\mathbf{v}_0(x)$ . By iterating we can write the solution to Equation (51) as the  $t$ -fold convolution

$$\mathbf{w}_t(x) = [\mathbf{K}(x-y) \circ \mathbf{B}_0]^{*t} \mathbf{w}_0(y). \quad (52)$$

282 Taking the bilateral Laplace transform

$$\mathcal{M}[\mathbf{w}_t(x)](s) = [\mathcal{M}[\mathbf{K}(x)](s) \circ \mathbf{B}_0]^t \mathcal{M}[\mathbf{w}_0(x)](s). \quad (53)$$

283 Since all of the dispersal kernels are Gaussian, we know that  $\mathcal{M}[\mathbf{K}(x)](s) = e^{\frac{\sigma^2 s^2}{2} + \mu s} \mathbf{1}$  where  $\mathbf{1}$  is a matrix  
284 of all ones. Then,

$$[\mathcal{M}[\mathbf{K}(x)](s) \circ \mathbf{B}_0]^t \mathcal{M}[\mathbf{w}_0(x)](s) = \left[ e^{\frac{\sigma^2 s^2}{2} + \mu s} \mathbf{1} \circ \mathbf{B}_0 \right]^t \mathcal{M}[\mathbf{w}_0(x)](s) \quad (54)$$

$$= \left[ e^{\frac{\sigma^2 s^2}{2} + \mu s} \mathbf{B}_0 \right]^t \mathcal{M}[\mathbf{w}_0(x)](s) \quad (55)$$

$$= e^{\frac{\sigma^2 t s^2}{2} + \mu t s} [\mathbf{B}_0]^t \mathcal{M}[\mathbf{w}_0(x)](s) \quad (56)$$

$$= [\mathbf{B}_0]^t \mathcal{M} \left[ \frac{1}{\sqrt{2\pi\sigma^2 t}} e^{-\frac{(x-\mu t)^2}{2\sigma^2 t}} \right] (s) \mathcal{M}[\mathbf{w}_0(x)](s) \quad (57)$$

$$= [\mathbf{B}_0]^t \mathcal{M} [(K_t * \mathbf{w}_0)(x)](s) \quad (58)$$

285 where  $K_t$  is  $N(\mu t, \sigma^2 t)$ . From (53)

$$\mathcal{M}[\mathbf{w}_t(x)](s) = [\mathbf{B}_0]^t \mathcal{M} [(K_t * \mathbf{w}_0)(x)](s). \quad (59)$$

286 Applying the inverse bilateral Laplace transform,

$$\mathbf{w}_t(x) = [\mathbf{B}_0]^t (K_t * \mathbf{w}_0)(x) \quad (60)$$

$$= [\mathbf{B}_0]^t \int_{-\infty}^{\infty} \frac{1}{\sqrt{2\pi\sigma^2 t}} e^{-\frac{(x-y-\mu t)^2}{2\sigma^2 t}} \mathbf{w}_0(y) dy \quad (61)$$

287 In the moving half-frame  $[A + ct, \infty)$  with  $c \geq c^*$  we have

$$\mathbf{w}_t(x_0 + ct) = [\mathbf{B}_0]^t \int_{-\infty}^{\infty} \frac{1}{\sqrt{2\pi\sigma^2 t}} e^{-\frac{(x_0+ct-y-\mu t)^2}{2\sigma^2 t}} \mathbf{w}_0(y) dy. \quad (62)$$

288 From (12), we know that  $c^* = \sqrt{2\sigma^2 \ln(\lambda_1)} + \mu$ , expanding the exponent, yields

$$\frac{(x_0 + ct - y - \mu t)^2}{2\sigma^2 t} = \frac{(x_0 - y)^2}{2\sigma^2 t} + \frac{2(c - \mu)t(x_0 - y) + (c - \mu)^2 t^2}{2\sigma^2 t} \quad (63)$$

$$\geq \frac{(x_0 - y)^2}{2\sigma^2 t} + \frac{c - \mu}{\sigma^2} (x_0 - y) + \ln(\lambda_1)t. \quad (64)$$

289 Thus,

$$\mathbf{w}_t(x_0 + ct) \leq \frac{[\mathbf{B}_0]^t}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(x_0-y)^2}{2\sigma^2 t}} e^{-\frac{c-\mu}{\sigma^2}(x_0-y)} e^{-\ln(\lambda_1)t} \mathbf{w}_0(y) dy \quad (65)$$

$$= \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t \frac{1}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(x_0-y)^2}{2\sigma^2 t}} e^{-\frac{c-\mu}{\sigma^2}(x_0-y)} \mathbf{w}_0(y) dy. \quad (66)$$

290 Since  $x_0 \geq A$  and  $e^{-\frac{(x_0-y)^2}{2\sigma^2 t}} \leq 1$ , we have

$$\mathbf{w}_t(x_0 + ct) \leq \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t \frac{e^{-\frac{A(c-\mu)}{\sigma^2}}}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{\frac{c-\mu}{\sigma^2} y} \mathbf{w}_0(y) dy. \quad (67)$$

291 From Lemma B.2, see Appendix B for details, we know that

$$\lim_{t \rightarrow \infty} \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t = \mathbf{r}\boldsymbol{\ell}, \quad (68)$$

where  $\mathbf{r}$  and  $\boldsymbol{\ell}$  are the right and left eigenvectors of  $\mathbf{B}_0$  corresponding to  $\lambda_1$  respectively with  $\boldsymbol{\ell}$  normalized by  $\langle \boldsymbol{\ell}^T, \mathbf{r} \rangle$  to account for the scaling in  $\mathbf{r}$ . Note that  $\mathbf{r}\boldsymbol{\ell}$  is a  $m \times m$  matrix since  $\mathbf{r}$  is  $m \times 1$  and  $\boldsymbol{\ell}$  is  $1 \times m$ . Thus since  $\int_{-\infty}^{\infty} e^{\frac{c-\mu}{\sigma^2} y} \mathbf{w}_0(y) dy < \infty$  by Assumption A5' and (68) we have  $\mathbf{w}_t(x_0 + ct) \rightarrow \mathbf{0}$  uniformly as  $t \rightarrow \infty$  in  $[A, \infty)$ . Recall that  $\mathbf{w}_t(x)$  was constructed as a super-solution,  $\mathbf{0} \leq \mathbf{v}_t(x) \leq \mathbf{w}_t(x)$ . This implies the uniform convergence of  $\mathbf{v}_t(x) \rightarrow \mathbf{0}$  as  $t \rightarrow \infty$  in the moving half-frame  $[A + ct, \infty)$ . The proof of Theorem 3.2 is complete.  $\square$

292 *Proof of Theorem 3.3*

293 *Proof* For simplicity, we focus on a single neutral fraction and drop the superscript  $i$ . Using the fact that  
294  $\mathbf{B}(\mathbf{u}_t(y))\mathbf{v} \leq \mathbf{B}_0\mathbf{v}$  for all  $\mathbf{v} \geq 0$  we can use a comparison principle to show that a new sequence  $\mathbf{w}_t(x)$   
295 defined by

$$\mathbf{w}_{t+1}(x) = \int_{-\infty}^{\infty} [\mathbf{K}(x-y) \circ \mathbf{B}_0] \mathbf{w}_t(y) dy \quad (69)$$

296 is a super-solution to any neutral fraction  $\mathbf{v}_t(x)$  with the same initial condition  $\mathbf{w}_0(x) = \mathbf{v}_0(x)$ . By iterating  
297 we can write the solution to Equation (69) as the  $t$ -fold convolution

$$\mathbf{w}_t(x) = [\mathbf{K}(x-y) \circ \mathbf{B}_0]^{*t} \mathbf{w}_0(y). \quad (70)$$

298 Taking the bilateral Laplace transform

$$\mathcal{M}[\mathbf{w}_t(x)](s) = [\mathcal{M}[\mathbf{K}(x)](s) \circ \mathbf{B}_0]^t \mathcal{M}[\mathbf{w}_0(x)](s). \quad (71)$$

299 Since all of the dispersal kernels are Gaussian, we know that  $\mathcal{M}[\mathbf{K}(x)](s) = e^{\frac{\sigma^2 s^2}{2} + \mu s} \mathbf{1}$  where  $\mathbf{1}$  is a matrix  
300 of all ones. Then, we can see that

$$[\mathcal{M}[\mathbf{K}(x)](s) \circ \mathbf{B}_0]^t \mathcal{M}[\mathbf{w}_0(x)](s) = \left[ e^{\frac{\sigma^2 s^2}{2} + \mu s} \mathbf{1} \circ \mathbf{B}_0 \right]^t \mathcal{M}[\mathbf{w}_0(x)](s) \quad (72)$$

$$= \left[ e^{\frac{\sigma^2 s^2}{2} + \mu s} \mathbf{B}_0 \right]^t \mathcal{M}[\mathbf{w}_0(x)](s) \quad (73)$$

$$= e^{\frac{\sigma^2 t s^2}{2} + \mu t s} \mathbf{I} [\mathbf{B}_0]^t \mathcal{M}[\mathbf{w}_0(x)](s) \quad (74)$$

$$= [\mathbf{B}_0]^t \mathcal{M} \left[ \frac{1}{\sqrt{2\pi\sigma^2 t}} e^{-\frac{(x-\mu t)^2}{2\sigma^2 t}} \mathbf{I} \right] (s) \mathcal{M}[\mathbf{w}_0(x)](s) \quad (75)$$

$$= [\mathbf{B}_0]^t \mathcal{M} [(\mathbf{K}_t * \mathbf{w}_0)(x)](s) \quad (76)$$

301 where  $\mathbf{K}_t$  is a diagonal matrix with  $N(\mu t, \sigma^2 t)$  entries and  $\mathbf{I}$  is the identity matrix. Thus, we have

$$\mathcal{M}[\mathbf{w}_t(x)](s) = [\mathbf{B}_0]^t \mathcal{M} [(\mathbf{K}_t * \mathbf{w}_0)(x)](s). \quad (77)$$

302 Then applying the inverse transform yields

$$\mathbf{w}_t(x) = [\mathbf{B}_0]^t (\mathbf{K}_t * \mathbf{w}_0)(x) \quad (78)$$

$$= [\mathbf{B}_0]^t \int_{-\infty}^{\infty} \frac{1}{\sqrt{2\pi\sigma^2 t}} e^{-\frac{(x-y-\mu t)^2}{2\sigma^2 t}} \mathbf{w}_0(y) dy \quad (79)$$

303 In the moving half-frame  $[A + ct, \infty)$  with fixed  $A \in \mathbb{R}$ , consider the element  $x_0 + ct$  with  $c = c^* =$   
 304  $\sqrt{2\sigma^2 \ln(\lambda_1)} + \mu$  where  $\lambda_1$  is the dominant eigenvalue of  $\mathbf{B}_0$  as given by (12). By rewriting  $\mathbf{w}_t(x)$  in this  
 305 moving half-frame we have

$$\mathbf{w}_t(x_0 + ct) = [\mathbf{B}_0]^t \int_{-\infty}^{\infty} \frac{1}{\sqrt{2\pi\sigma^2 t}} e^{-\frac{(x_0+ct-y-\mu t)^2}{2\sigma^2 t}} \mathbf{w}_0(y) dy. \quad (80)$$

306 Expanding the exponent, yields

$$\frac{(x_0 + ct - y - \mu t)^2}{2\sigma^2 t} = \frac{(y - x_0)^2}{2\sigma^2 t} + \frac{(c - \mu)(x_0 - y)}{\sigma^2} + \frac{(c - \mu)^2}{2\sigma^2} t. \quad (81)$$

307 Thus,

$$\mathbf{w}_t(x_0 + ct) = \frac{[\mathbf{B}_0]^t}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(y-x_0)^2}{2\sigma^2 t}} e^{-\frac{(c-\mu)(x_0-y)}{\sigma^2}} e^{-\frac{(c-\mu)^2}{2\sigma^2} t} \mathbf{w}_0(y) dy \quad (82)$$

$$= \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t \frac{1}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(y-x_0)^2}{2\sigma^2 t}} e^{-\frac{(c-\mu)(x_0-y)}{\sigma^2}} e^{\left[ -\frac{(c-\mu)^2}{2\sigma^2} + \ln(\lambda_1) \right] t} \mathbf{w}_0(y) dy. \quad (83)$$

308 Since  $c = c^* = \sqrt{2\sigma^2 \ln(\lambda_1)} + \mu$ , we have that

$$\mathbf{w}_t(x_0 + ct) = \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t \frac{1}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(y-x_0)^2}{2\sigma^2 t}} e^{-\frac{(c-\mu)(x_0-y)}{\sigma^2}} \mathbf{w}_0(y) dy. \quad (84)$$

309 From Assumption A5'',  $\mathbf{w}_0(y) = (\mathbf{p}_0 \circ \mathbf{r}) e^{-\frac{c-\mu}{\sigma^2} y}$ . Thus,

$$\mathbf{w}_t(x_0 + ct) = \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t (\mathbf{p}_0 \circ \mathbf{r}) e^{-\frac{(c-\mu)}{\sigma^2} x_0} \frac{1}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(y-x_0)^2}{2\sigma^2 t}} dy \quad (85)$$

$$= \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t (\mathbf{p}_0 \circ \mathbf{r}) e^{-\frac{(c-\mu)}{\sigma^2} x_0}. \quad (86)$$

310 From Lemma B.2, see Appendix B for details, we know that

$$\lim_{t \rightarrow \infty} \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t = \mathbf{r} \boldsymbol{\ell} \quad (87)$$

311 where  $\mathbf{r}$  and  $\boldsymbol{\ell}$  are the right and left eigenvectors of  $\mathbf{B}_0$  corresponding to  $\lambda_1$  respectively where  $\boldsymbol{\ell}$  is normalized  
 312 by  $\langle \boldsymbol{\ell}^T, \mathbf{r} \rangle$ . Thus,

$$\lim_{t \rightarrow \infty} \mathbf{w}_t(x_0 + ct) = \lim_{t \rightarrow \infty} \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t (\mathbf{p}_0 \circ \mathbf{r}) e^{-\frac{(c-\mu)}{\sigma^2} x_0} \quad (88)$$

$$= \mathbf{r} \boldsymbol{\ell} (\mathbf{p}_0 \circ \mathbf{r}) e^{-\frac{(c-\mu)}{\sigma^2} x_0} \quad (89)$$

$$= e^{-\frac{(c-\mu)}{\sigma^2} x_0} \mathbf{r} p. \quad (90)$$

313 From the above calculations, we find that the super-solution approaches a proportion,  $p$ , of the traveling  
 314 wave for the linear equation where  $p = \boldsymbol{\ell} (\mathbf{p}_0 \circ \mathbf{r})$ . We now move onto our sub-solution. For any  $0 < \varepsilon \ll 1$ ,  
 315  $\boldsymbol{\delta}$  is chosen such that  $(1 - \varepsilon) \mathbf{B}_0 \boldsymbol{\delta} = \mathbf{B}(\boldsymbol{\delta}) \boldsymbol{\delta}$  and we define

$$(\mathbf{B}_{sub}(\mathbf{u}; \varepsilon))_{jl} := \begin{cases} (1 - \varepsilon) (\mathbf{B}(\mathbf{u}))_{jl} & \text{if } (\mathbf{B}(\mathbf{u}))_{jl} \text{ is constant} \\ \beta_{jl}(\mathbf{u}; \varepsilon) & \text{if } (\mathbf{B}(\mathbf{u}))_{jl} \text{ is non-constant,} \end{cases} \quad (91)$$

316 where

$$\beta_{jl}(\mathbf{u}; \varepsilon) := \begin{cases} (1 - \varepsilon) (\mathbf{B}_0)_{jl} & \text{for } 0 \leq \mathbf{u} < \boldsymbol{\delta} \\ (\mathbf{B}(\mathbf{u}))_{jl} & \text{for } \mathbf{u} \geq \boldsymbol{\delta}. \end{cases} \quad (92)$$

317 Then,

$$\mathbf{z}_{t+1}(x) = \int_{-\infty}^{\infty} [\mathbf{K}(x-y) \circ \mathbf{B}_{sub}(\mathbf{u}_t(y); \varepsilon)] \mathbf{z}_t(y) dy \quad (93)$$

318 with  $\mathbf{z}_0(x) = \mathbf{v}_0(x)$  is a sub-solution of  $\mathbf{v}_t(x)$  by the comparison principle since  $\mathbf{B}_{sub}(\mathbf{u}; \varepsilon)\mathbf{v} \leq \mathbf{B}(\mathbf{u})\mathbf{v}$  for  
 319 all  $\mathbf{v} \geq 0$ . Define  $c(\varepsilon) := \sqrt{2\sigma^2 \ln((1-\varepsilon)\lambda_1)} + \mu$  where  $(1-\varepsilon)\lambda_1$  is the dominant eigenvalue of the constant  
 320 matrix  $(1-\varepsilon)\mathbf{B}_0$ . In the moving half-frame  $[A + c(\varepsilon)t, \infty)$  with fixed  $A \in \mathbb{R}$ , choose  $x_0$  large such that  
 321  $\mathbf{u}_t(y)$  in (93) satisfies  $\mathbf{u}_t(y) < \delta$  for all  $t$  where  $y \in [x_0 + c(\varepsilon)t, \infty)$ . Then by the definition of  $\mathbf{B}_{sub}(\mathbf{u}; \varepsilon)$

$$\mathbf{z}_{t+1}(x_0 + c(\varepsilon)t) = \int_{-\infty}^{\infty} [\mathbf{K}(x_0 + c(\varepsilon)t - y) \circ (1-\varepsilon)\mathbf{B}_0] \mathbf{z}_t(y) dy. \quad (94)$$

322 By iterating we can write the solution to (94) as the  $t$ -fold convolution

$$\mathbf{z}_t(x_0 + c(\varepsilon)t) = [\mathbf{K}(x_0 + c(\varepsilon)t - y) \circ (1-\varepsilon)\mathbf{B}_0]^{*t} \mathbf{z}_0(y). \quad (95)$$

323 Since we assumed that all of the dispersal kernels are Gaussian, by repeating calculations done previously  
 324 we find that

$$\mathbf{z}_t(x_0 + c(\varepsilon)t) = [(1-\varepsilon)\mathbf{B}_0]^t \int_{-\infty}^{\infty} \frac{1}{\sqrt{2\pi\sigma^2 t}} e^{-\frac{(x_0 + c(\varepsilon)t - y - \mu t)^2}{2\sigma^2 t}} \mathbf{z}_0(y) dy \quad (96)$$

$$= \frac{[(1-\varepsilon)\mathbf{B}_0]^t}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(x_0 - y)^2}{2\sigma^2 t}} e^{-\frac{(c(\varepsilon) - \mu)(x_0 - y)}{\sigma^2}} e^{-\frac{(c(\varepsilon) - \mu)^2}{2\sigma^2} t} \mathbf{z}_0(y) dy \quad (97)$$

$$= \left[ \frac{(1-\varepsilon)\mathbf{B}_0}{(1-\varepsilon)\lambda_1} \right]^t \frac{1}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(x_0 - y)^2}{2\sigma^2 t}} e^{-\frac{(c(\varepsilon) - \mu)(x_0 - y)}{\sigma^2}} e^{\left[ -\frac{(c(\varepsilon) - \mu)^2}{2\sigma^2} + \ln((1-\varepsilon)\lambda_1) \right] t} \mathbf{z}_0(y) dy \quad (98)$$

$$= \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t \frac{1}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(x_0 - y)^2}{2\sigma^2 t}} e^{-\frac{(c(\varepsilon) - \mu)(x_0 - y)}{\sigma^2}} e^{\left[ -\frac{(c(\varepsilon) - \mu)^2}{2\sigma^2} + \ln((1-\varepsilon)\lambda_1) \right] t} \mathbf{z}_0(y) dy. \quad (99)$$

325 Since  $c(\varepsilon) = \sqrt{2\sigma^2 \ln((1-\varepsilon)\lambda_1)} + \mu$ ,

$$\mathbf{z}_t(x_0 + c(\varepsilon)t) = \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t \frac{1}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(x_0 - y)^2}{2\sigma^2 t}} e^{-\frac{(c(\varepsilon) - \mu)(x_0 - y)}{\sigma^2}} \mathbf{z}_0(y) dy. \quad (100)$$

326 Note that the integrand in (100) is nonnegative and integrable. Using Fatou's lemma we fix  $t$  and let  $\varepsilon \rightarrow 0$ ,  
 327 giving

$$\mathbf{z}_t(x_0 + ct) = \liminf_{\varepsilon \rightarrow 0} \mathbf{z}_t(x_0 + c(\varepsilon)t) \quad (101)$$

$$= \liminf_{\varepsilon \rightarrow 0} \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t \frac{1}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(x_0 - y)^2}{2\sigma^2 t}} e^{-\frac{(c(\varepsilon) - \mu)(x_0 - y)}{\sigma^2}} \mathbf{z}_0(y) dy \quad (102)$$

$$\geq \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t \frac{1}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} \liminf_{\varepsilon \rightarrow 0} e^{-\frac{(x_0 - y)^2}{2\sigma^2 t}} e^{-\frac{(c(\varepsilon) - \mu)(x_0 - y)}{\sigma^2}} \mathbf{z}_0(y) dy \quad (103)$$

$$= \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t \frac{1}{\sqrt{2\pi\sigma^2 t}} \int_{-\infty}^{\infty} e^{-\frac{(x_0 - y)^2}{2\sigma^2 t}} e^{-\frac{(c - \mu)(x_0 - y)}{\sigma^2}} \mathbf{z}_0(y) dy. \quad (104)$$

328 From Assumption A5'',  $\mathbf{z}_0(y) = (\mathbf{p}_0 \circ \mathbf{r}) e^{-\frac{(c-\mu)}{\sigma^2} y}$ . Thus, by the same calculations used in (85)-(86) for the  
 329 super-solution

$$\mathbf{z}_t(x_0 + ct) \geq \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t (\mathbf{p}_0 \circ \mathbf{r}) e^{-\frac{(c-\mu)}{\sigma^2} x_0}. \quad (105)$$

330 From Lemma B.2, see Appendix B for details, we see that

$$\lim_{t \rightarrow \infty} \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t = \mathbf{r}\ell, \quad (106)$$

331 where  $\mathbf{r}$  and  $\boldsymbol{\ell}$  are the right and left eigenvectors corresponding to  $\lambda_1$  respectively where the  $\boldsymbol{\ell}$  is normalized  
 332 by  $\langle \boldsymbol{\ell}^T, \mathbf{r} \rangle$ . Thus,

$$\lim_{t \rightarrow \infty} \mathbf{z}_t(x_0 + ct) \geq \lim_{t \rightarrow \infty} \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t (\mathbf{p}_0 \circ \mathbf{r}) e^{-\frac{(c-\mu)}{\sigma^2} x_0} \quad (107)$$

$$= \mathbf{r} \boldsymbol{\ell} (\mathbf{p}_0 \circ \mathbf{r}) e^{-\frac{(c-\mu)}{\sigma^2} x_0} \quad (108)$$

$$= e^{-\frac{(c-\mu)}{\sigma^2} x_0} \mathbf{r} \mathbf{p}. \quad (109)$$

333 Asymptotically, our sub-solution is bounded below by a proportion of the traveling wave for the linear  
 334 equation where  $p = \boldsymbol{\ell} (\mathbf{p}_0 \circ \mathbf{r})$ . Since our super-solution satisfies

$$\lim_{t \rightarrow \infty} \mathbf{w}_t(x_0 + ct) \leq e^{-\frac{(c-\mu)}{\sigma^2} x_0} \mathbf{r} \mathbf{p}, \quad (110)$$

335 and our sub-solution satisfies

$$\lim_{t \rightarrow \infty} \mathbf{z}_t(x_0 + ct) \geq e^{-\frac{(c-\mu)}{\sigma^2} x_0} \mathbf{r} \mathbf{p} \quad (111)$$

336 it follows that

$$\lim_{t \rightarrow \infty} \mathbf{v}_t(x_0 + ct) = e^{-\frac{(c-\mu)}{\sigma^2} x_0} \mathbf{r} \mathbf{p}. \quad (112)$$

The proof of Theorem 3.3 is complete.  $\square$

#### 337 4 Numerical simulations

338 In this section, we illustrate the theory of Section 3 with a numerical example. All simulations were done  
 339 using the fast Fourier transform technique (Cooley and Tukey 1965). This method is better than classical  
 340 quadrature because it speeds up the numerical process from  $O(n^2)$  to  $O(n \log(n))$ .

341 We begin with a two-stage population model of juveniles,  $J$ , and adults,  $A$ . The equations in this model  
 342 are given below,

$$\begin{aligned} J_{t+1}^i(x) &= \int_{-\infty}^{\infty} k(x-y) \zeta (1-m) J_t^i(y) dy + \int_{-\infty}^{\infty} k(x-y) f_0 e^{-\sum_{i=1}^n (J_t^i(y) + A_t^i(y))} A_t^i(y) dy, \\ A_{t+1}^i(x) &= \int_{-\infty}^{\infty} k(x-y) \zeta m J_t^i(y) dy + \int_{-\infty}^{\infty} k(x-y) \zeta A_t^i(y) dy, \end{aligned} \quad (113)$$

343 where

$$k(x-y) = \frac{1}{\sqrt{2\pi\sigma^2}} e^{-\frac{(x-y)^2}{2\sigma^2}}. \quad (114)$$

344 The demography in (113) follows a classical model for biological invasions (Neubert and Caswell 2000), but  
 345 we assume Gaussian dispersal to align with the assumptions in our theorems. In (113),  $\zeta$  is the probability  
 346 of survival to the next generation,  $m$  is the probability of maturation from a juvenile to an adult,  $f_0$  is the  
 347 number of juveniles produced by an adult in the absence of density-dependent effects. All individuals are  
 348 assumed to disperse according to a Gaussian dispersal kernel. The growth function for adults producing  
 349 juveniles is assumed to be a Ricker type growth function where the nonlinearity depends on the density of  
 350 both juveniles and adults. In the juvenile equation of (113), juveniles can remain juveniles if they survive  
 351 and do not mature and adults from location  $y$  can produce juveniles that disperse to location  $x$ . In the  
 352 adult equation of (113), juveniles become adults if they survive and mature, and adults remain adults if  
 353 they survive from the previous year.

354 Let

$$\mathbf{v}_t^i(x) = \begin{bmatrix} J_t^i(x) \\ A_t^i(x) \end{bmatrix}, \quad (115)$$

$$\mathbf{K}(x-y) = \begin{bmatrix} \frac{1}{\sqrt{2\pi\sigma^2}} e^{-\frac{(x-y)^2}{2\sigma^2}} & \frac{1}{\sqrt{2\pi\sigma^2}} e^{-\frac{(x-y)^2}{2\sigma^2}} \\ \frac{1}{\sqrt{2\pi\sigma^2}} e^{-\frac{(x-y)^2}{2\sigma^2}} & \frac{1}{\sqrt{2\pi\sigma^2}} e^{-\frac{(x-y)^2}{2\sigma^2}} \end{bmatrix}, \text{ and} \quad (116)$$

$$\mathbf{B}(\mathbf{u}_t(y)) = \begin{bmatrix} \zeta(1-m) f_0 e^{-\sum_{i=1}^n (J_t^i(y) + A_t^i(y))} \\ \zeta m & \zeta \end{bmatrix}. \quad (117)$$

355 Then we can write (113) in the matrix and vector notation provided in (4).

356 First, let us verify that the assumptions of Theorems 3.2 and 3.3 are satisfied. Recall that Assumptions  
 357 A1-A3 and A4' are the same for these two theorems. For Assumption A1, it is clear that our population  
 358 projection matrix,  $\mathbf{B}(\mathbf{u}_t(y))$ , is nonnegative from (117) since  $\zeta, m, f_0 > 0$ . We can calculate  $\mathbf{B}_0$  to be

$$\mathbf{B}_0 = \begin{bmatrix} \zeta(1-m) & f_0 \\ \zeta m & \zeta \end{bmatrix}. \quad (118)$$

359 Thus,  $\mathbf{B}_0$  is primitive. For Assumption A2, the dominant eigenvalue of  $\mathbf{B}_0$  is greater than one if

$$f_0 > \frac{(1-\zeta)(1-\zeta(1-m))}{\zeta m}. \quad (119)$$

360 For details of this calculation see Proposition 3.1 of Marculis and Lui (2016). Since  $e^{-\sum_{i=1}^n (J_i^i(y) + A_i^i(y))} \leq 1$   
 361 we have  $\mathbf{B}(\mathbf{u}_t(y))\mathbf{v} \leq \mathbf{B}_0\mathbf{v}$  for all  $\mathbf{v} \geq 0$  and Assumption A3 is satisfied. Even though our operator is not  
 362 order preserving because of the overcompensation in the Ricker function, Proposition 3.1 in Li et al. (2009)  
 363 suggests that the calculation for the spreading speed should still hold true. Assumption A4' is clear from  
 364 the definition of (116). Finally, if we assume our initial condition to decay faster than  $e^{-\frac{c-\mu}{\sigma^2}x}$ , then the  
 365 neutral fractions will satisfy Assumption A5' of Theorem 3.2 and we can see that (113) has a unique positive  
 366 steady state given by

$$J^* = \frac{1-\zeta}{\zeta m} A^* \text{ and } A^* = -\ln\left(\frac{(1-\zeta)(1-\zeta(1-m))}{f_0 \zeta m}\right), \quad (120)$$

367 see again Proposition 3.1 of Marculis and Lui (2016). In our numerical simulations the only neutral fraction  
 368 that does not decay faster than  $e^{-\frac{c-\mu}{\sigma^2}x}$  is the one at the leading edge because it was chosen to have an  
 369 initial form of the traveling wave solution with  $c = c^*$ . It should be mentioned here that since we are  
 370 solving this problem numerically it is solved on a finite domain and this is only an approximation to the  
 371 solution. Therefore, in the moving half-frame, the only neutral fractions that we see are the ones initially  
 372 at the leading edge. The neutral fractions at the leading edge do not satisfy the exact Assumption A5'' of  
 373 Theorem 3.3, but asymptotically they decay like  $e^{-\frac{c-\mu}{\sigma^2}x}$ . However, the asymptotic proportion calculated  
 374 from Theorem 3.3 agrees with the numerical simulation suggesting that this result should be able to extend  
 375 to a wider array of initial conditions.

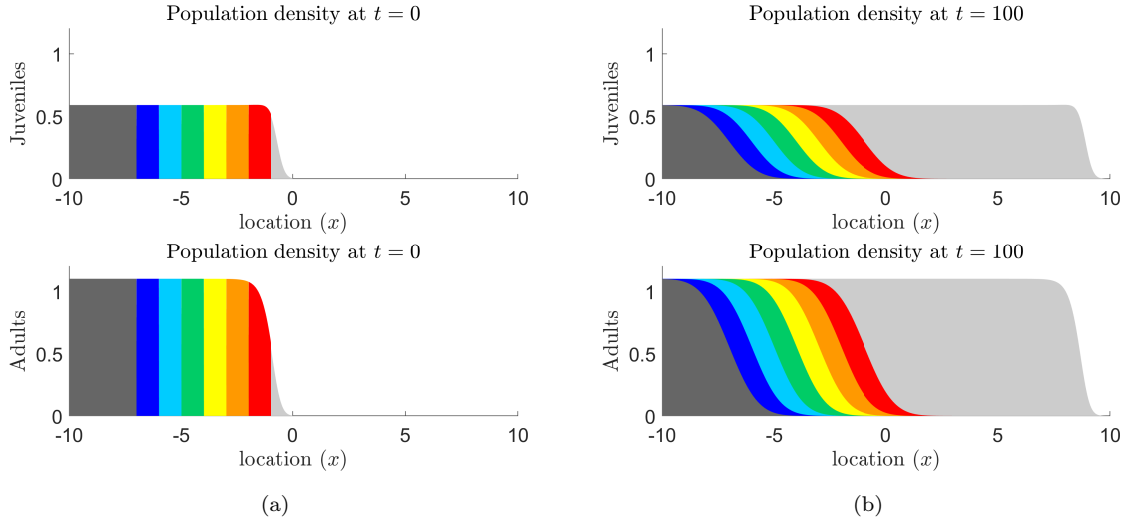
376 We provide some numerical simulations to see the neutral genetic patterns produced by (113). We begin  
 377 by running a simulation where the juvenile and adult populations have the same initial distribution as seen  
 378 in Figure 1(a). This simulation shows that the spread of both juveniles and adults is dominated by the  
 379 neutral fraction at the leading edge as seen in Figure 1(b). Switching the ordering of the neutral fractions  
 380 behind the leading edge does not affect the asymptotic behavior in the moving frame. This observation is  
 381 consistent with the founder effect. The simulations seen in Figure 1 agree with the results of Theorems 3.2  
 382 and 3.3.

383 For our next simulation, we consider the case where the distribution of the neutral fractions of juveniles  
 384 and adults do not appear in the same order. This is seen in Figure 2(a). Here we keep the same initial  
 385 distribution of juvenile individuals as in Figure 1(a), but the initial distribution of adult neutral fractions  
 386 is assorted differently. In Figure 2(a) we can see that initially the neutral fractions at the leading edge of  
 387 the juvenile and adult populations are light gray and red respectively. Figure 2(b) shows the distribution of  
 388 neutral fractions at  $t = 100$ . At the leading edge the spread is dominated by the light gray and red neutral  
 389 fractions. This simulation agrees with our theoretical results because Theorem 3.2 and 3.3 suggest that the  
 390 spread should be dominated by the neutral fractions that are initially at the leading edge of the population.  
 391 Again we see that the neutral fractions behind the leading edge do not contribute to the asymptotic spread.

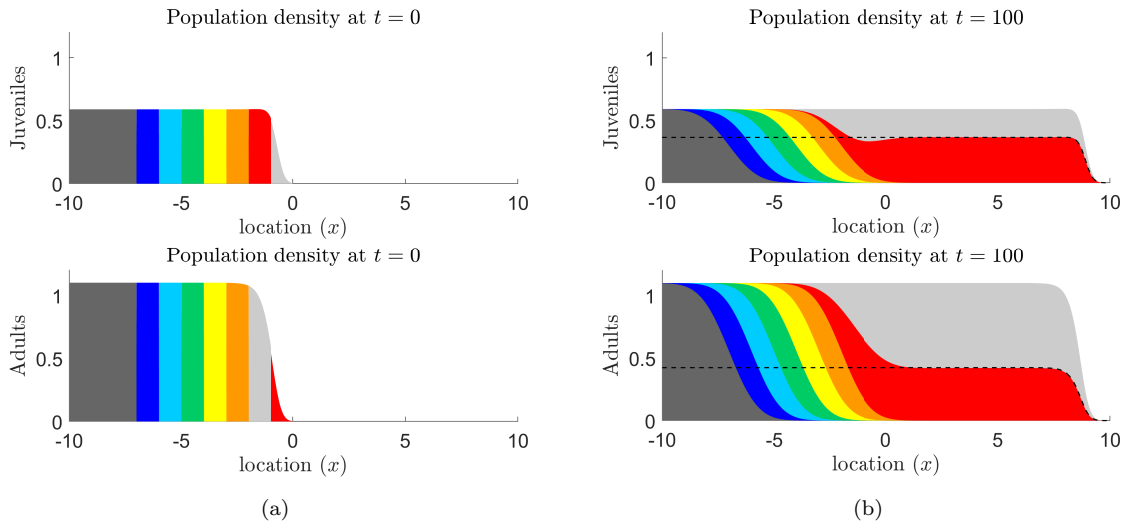
## 392 5 Discussion

393 The main objective of this work is to understand the effect that stage-structure has on the neutral genetic  
 394 composition of expanding populations as outlined in Section 1. We derived the model for the inside dyna-  
 395 mics of a stage-structured integrodifference equation in Section 2.1. Section 2.2 describes five of our main  
 396 assumptions related to demography and dispersal. Four of these assumptions are related to the population  
 397 projection matrix and the fifth is related to the form of the dispersal kernel.

398 The three main results of the paper are provided in Section 3, with their respective proofs in Section 3.3.  
 399 Theorem 3.1 is our first main result, which provides sufficient conditions for a neutral fraction to converge



**Fig. 1** Numerical realization of (113) for the parameter values  $\sigma^2 = 0.01$ ,  $\mu = 0$ ,  $\zeta = 0.7$ ,  $m = 0.8$ , and  $f_0 = 2.5$  for  $n = 8$  neutral fractions. In (a) the plots are the initial conditions for the juvenile and adult populations. Notice that the distribution of neutral fractions for juvenile and adult populations have the same order. In (b) we plot the densities of the juvenile and adult neutral fractions at  $t = 100$ .



**Fig. 2** Numerical realization of (113) for the parameter values  $\sigma^2 = 0.01$ ,  $\mu = 0$ ,  $\zeta = 0.7$ ,  $m = 0.8$ , and  $f_0 = 2.5$  for  $n = 8$  neutral fractions. For these parameters  $\mathbf{u}^* = (J^*, A^*) = (0.5900, 1.1013)$ . In (a) the plots are the initial conditions for the juvenile and adult population. Notice that the distribution of the first two neutral fractions is different for juveniles and adults. The plots in (b) are the densities of the juvenile and adult neutral fractions at  $t = 100$ . The neutral genetic pattern produced here is due to the difference in the initial distribution of neutral fractions for juveniles and adults. The dashed lines in (b) are calculated from Theorem 3.3, they represent the proportions of red juveniles and adults. Behind the leading edge the proportions are  $p^2 J^* = 0.3629$  for juveniles and  $p^1 A^* = 0.4238$  for adults.

400 uniformly to zero in the moving half-frame. The five assumptions that must be satisfied are as follows: the  
 401 population projection matrix must be nonnegative, the population projection matrix evaluated at zero must  
 402 be primitive and its dominant eigenvalue must be greater than one, the population projection matrix must  
 403 be maximal at the trivial steady state, all dispersal kernels must be thin-tailed, and the initial condition  
 404 must satisfy the decay assumption given in Lemma B.1. It should be noted that the Dirac delta function  
 405 is a thin-tailed dispersal kernel and thus we can consider cases where there is no dispersal between some  
 406 transitions making this theorem very general in terms of the dispersal assumptions.



407 The second main result is Theorem 3.2. Similar to Theorem 3.1, this theorem also shows conditions  
408 under which each neutral fraction converges uniformly to zero in the moving half-frame. The difference  
409 with this theorem is that we make a stronger assumption on the dispersal kernels in exchange for a weaker  
410 condition on the initial condition. In particular, we assume that all dispersal kernels are Gaussian with  
411 identical means and variances. Due to this assumption, we are then able to relax the decay condition on  
412 the initial condition of the population to be slightly weaker than is required for Theorem 3.1. The proof  
413 for Theorem 3.2 is more elegant than the proof for Theorem 3.1. However, this comes at some cost in the  
414 biological realism of the model since it is not common for all stages and transitions to disperse exactly via  
415 a Gaussian distribution.

416 The final result is given in Theorem 3.3. The first four assumptions of this theorem are the same as  
417 Theorem 3.2. The fifth assumption assumes that the initial condition decays according to the traveling  
418 wave ansatz for the linear equation. Under these assumptions, we are able to asymptotically calculate the  
419 proportion that each neutral fraction approaches in the moving frame. This proportion is dependent on the  
420 right and left eigenvectors of the population projection matrix evaluated at zero and the initial proportion  
421 of each neutral fraction at the leading edge. The proof relies on the construction of super- and sub-solutions  
422 to the system. The super-solution, as expected, is chosen to be the linearization of our operator while the  
423 sub-solution is defined in a piecewise manner to lie below the nonlinearities. Since all dispersal kernels were  
424 assumed to be identical Gaussian distributions, the proportion calculated by Theorem 3.3 does not apply  
425 when some stages and transitions do not disperse in the same way.

426 After completion of the mathematical results, we performed some numerical simulations in Section 4  
427 to compare our analytical results to a reasonable biological model. We chose to look at a classical two-  
428 stage juvenile adult model where dispersal occurs between all stages and transitions. The first simulation,  
429 in Figure 1, shows that the spread is dominated by the neutral fraction at the leading edge which is an  
430 extreme version of the founder effect. However, since we are working with a system of equations, it is  
431 possible for the initial distribution of neutral fractions in the juvenile and adult stages to be different. This  
432 is seen in Figure 2(a). As predicted from Theorem 3.2, in Figure 2(b), we see that all neutral fractions,  
433 except the ones at the leading edge of the juvenile and adult populations, converge uniformly to zero in  
434 the moving half-frame. The asymptotic proportions for the two neutral fractions that were initially at the  
435 leading edge of the juvenile and adult populations are given by the formula in Theorem 3.3 and plotted as  
436 the dashed line in Figure 2(b).

437 As expected, some of the same results obtained here are similar to those for the scalar population  
438 model. That is, Theorem 3.1 and Theorem 3.2 are equivalent to their scalar counterparts, Theorem 3 and  
439 Theorem 1 respectively, given in Marculis et al. (2017). However, Theorem 3.3 provides a new result for a  
440 special case of interacting neutral fractions at the leading edge. This is not possible in the scalar population  
441 model. From this theorem, we see the ability for multiple neutral fractions to contribute to the spread of  
442 the population. Contributions from multiple neutral fractions to the population spread are only possible  
443 in the scalar model when there is a strong Allee effect (Marculis et al. 2017). Although we would expect  
444 similar behavior from our stage-structured model, we are not able to analyze the inside dynamics of a  
445 stage-structured model with a strong Allee effect. This is due to the requirement that our results for the  
446 strong Allee effect in scalar systems rely on the operator being compact. For a system of equations the  
447 necessary theory is more complicated and we were unable to perform this analysis. In the special case  
448 where all dispersal kernels are Gaussian with the same mean and variance and all entries of the population  
449 projection matrix have the same strong Allee effect type per-capita growth function, then Theorem 2 given  
450 in Marculis et al. (2017) can be applied. However, such stringent assumptions would defeat the purpose for  
451 considering a stage-structured population model because all stages and transitions would grow and disperse  
452 in the same way, essentially reducing the stage-structured model to a scalar equation.

453 The interesting additional feature that the stage-structured population model offers over scalar models  
454 is the ability to have a different initial distribution of neutral fractions for each stage. This difference can  
455 lead to multiple neutral fractions driving the spread of the population. Here, we see these dynamics solely  
456 for the reason that the initial spatial distribution of neutral fractions is different for each stage.

457 Several assumptions about the integrodifference dynamics and dispersal kernels limit the applicability  
458 of the results in this paper. One limitation to the applicability of our work is seen in Assumption A3.  
459 Here we require that our population projection matrix is maximal at zero. This means that we are not  
460 considering any kind of demography with Allee effects. In order to prove the asymptotic proportion result  
461 seen in Theorem 3.3 we make some restrictive assumptions on the dispersal kernels and initial conditions  
462 in the model. Assumption A4' in Theorem 3.3 states that all dispersal kernels are Gaussian with the same  
463 mean and variance. This assumption may be unrealistic for many populations because the reason to use a  
464 stage-structured population model over a scalar population model is to include differences in demography

465 and dispersal between stages. Assumption  $A5''$  in Theorem 3.3 makes the assumption that the initial  
466 conditions are in the form of the traveling wave ansatz for the linear equation. It would be beneficial  
467 to generalize Theorem 3.3 for initial conditions that are in the form of the traveling wave solution. The  
468 numerical simulations show that we should be able to relax our sixth assumption in our in theorems to a  
469 more general class of initial conditions. These simulations are not only useful for verifying our mathematical  
470 results, but they also provide some insight into opportunities for further mathematical analysis.

## 471 A Asymptotic speed of propagation for a system

472 The following Proposition is taken from (Lui 1989a). Let  $\beta \in \mathbb{R}^n$  be a positive vector. We define

$$C = \{\mathbf{u} = (u^1, \dots, u^n) \mid \mathbf{0} \leq \mathbf{u}(x) \leq \beta, u^i(x) : \mathbb{R} \rightarrow [0, \beta^i] \\ \text{is piecewise continuous for } i = 1, \dots, n\}.$$

473 The operator  $\mathbf{Q}$  used in our analytical results is given by

$$\mathbf{Q}[\mathbf{u}] = \int_{-\infty}^{\infty} [\mathbf{K}(x-y) \circ \mathbf{B}(\mathbf{u}(y))] \mathbf{u}(y) dy. \quad (121)$$

474 **Proposition A.1** Let  $\mathbf{Q} = (Q^1, \dots, Q^n) : C \rightarrow C$  satisfy the following conditions:

- 475 (1)  $\mathbf{Q}[\mathbf{0}] = \mathbf{0}$ ,  $\mathbf{Q}[\beta] = \beta$ ,  $\mathbf{0}$  is unstable and  $\beta$  is stable with respect to  $\mathbf{Q}$ .  
476 (2)  $\mathbf{Q}$  is translation invariant and has no other fixed-point besides  $\mathbf{0}$  and  $\beta$  in  $C$ .  
477 (3)  $\mathbf{Q}$  is monotone or order-preserving in  $C$ ; that is, if  $\mathbf{u} \leq \mathbf{v}$  in  $C$ , then  $\mathbf{Q}[\mathbf{u}] \leq \mathbf{Q}[\mathbf{v}]$ .  
478 (4)  $\mathbf{Q}$  is continuous in the topology of uniform convergence on bounded subsets of  $\mathbb{R}$ .  
479 (5) Let

$$(\mathbf{M}[\mathbf{u}](x))_i = \sum_{j=1}^n \int_{-\infty}^{\infty} \mathbf{u}_j(x-y) m^{ij}(y) dy. \quad (122)$$

480 be the linearization of  $\mathbf{Q}$  at  $\mathbf{0}$ , where  $m^{ij}(y) \geq 0$  is an integrable function. We assume that

$$\mathbf{Q}[\mathbf{u}] \leq \mathbf{M}[\mathbf{u}] \quad \text{for all } \mathbf{u} \in C. \quad (123)$$

481 (6) The matrix  $\mathbf{B}(s) = (b^{ij}(s))$ , where

$$b^{ij}(s) = \int_{-\infty}^{\infty} e^{sy} m^{ij}(y) dy \quad (124)$$

482 is irreducible for  $0 < s < s^+$ .

483 Let  $\rho(s)$  be the dominant eigenvalue of  $\mathbf{B}(s)$  and let

$$c^* = \min_{0 < s < s^+} \frac{1}{s} \ln \rho(s). \quad (125)$$

484 Then  $c^*$  is the asymptotic speed of propagation of the operator  $\mathbf{Q}$  in the positive direction in the following sense. Let  $\mathbf{u}_0 \in C$ ,  
485  $\mathbf{u}_0$  is non-trivial and vanishes outside of a bounded interval in  $\mathbb{R}$ . Let  $\mathbf{u}_t$  be defined by  $\mathbf{u}_{t+1} = \mathbf{Q}[\mathbf{u}_t]$  for  $t = 0, 1, 2, \dots$   
486 Then for any small  $\varepsilon > 0$ ,

$$\lim_{t \rightarrow \infty} \min_{x \leq t(c^* - \varepsilon)} |\mathbf{u}_t(x) - \beta| = 0 \quad (126)$$

$$\text{and} \quad \lim_{t \rightarrow \infty} \max_{x \geq t(c^* + \varepsilon)} |\mathbf{u}_t(x)| = 0. \quad (127)$$

## 487 B Mathematical details

488 The purpose of this section is to provide the mathematical background needed to prove the theorems in Section 3. One  
489 tool that is used throughout all of our theorems is the reflected Bilateral Laplace transform.

490 **Definition 2** Let  $f : \mathbb{R} \rightarrow \mathbb{R}$  where  $f$  is piecewise continuous on every finite interval in  $\mathbb{R}$  and there exists a  $M \in \mathbb{R}^+$   
491 such that  $|f(x)| \leq M e^{-sx}$  for all  $x \in \mathbb{R}$  and  $0 < s < s^+$ . Then, the reflected bilateral Laplace transform and its inverse  
492 are defined to be

$$F(s) = \mathcal{M}[f(x)] := \int_{-\infty}^{\infty} f(x) e^{sx} dx, \quad \text{and} \quad (128)$$

$$f(x) = \mathcal{M}^{-1}[F(s)] := \frac{1}{2\pi i} \lim_{R \rightarrow \infty} \int_{\gamma - iR}^{\gamma + iR} F(s) e^{-sx} ds \quad (129)$$

493 for  $0 < s < s^+$ , where the integration in Equation (129) is over the vertical line,  $\text{Re}(s) = \gamma$  in the complex plane and  $\gamma$  is  
494 greater than the real parts of all singularities of  $F(s)$ .

495 By using the convolution theorem, the reflected bilateral Laplace transform can be used to write the solution to our  
496 model in terms of the initial condition. This theorem states that the reflected bilateral Laplace transform of a convolution  
497 is the product of the reflected bilateral Laplace transforms. That is,

$$\mathcal{M}[f(x) * h(x)](s) = F(s)H(s). \quad (130)$$

498 Note that the reflected bilateral Laplace transform of a probability density function is also referred to as its moment  
499 generating function.

500 Next, we provide results regarding vector and matrix analysis that are relevant to our subsequent analysis. First, it  
501 should be noted that when we write  $\mathbf{x} \geq \mathbf{y}$ , the inequality is element-wise. That is,  $x_i \geq y_i$  for each  $i$ . In a similar manner,  
502  $\mathbf{x} > \mathbf{y}$  means that  $x_i > y_i$  for each  $i$ . For the matrix analysis, the following definitions and proposition are needed:

503 **Definition 3** Let  $\lambda_1, \dots, \lambda_m$  be the eigenvalues of a matrix  $\mathbf{A}$ . Then its spectral radius  $\rho(\mathbf{A})$  is defined as:

$$\rho(\mathbf{A}) := \max_{i=1, \dots, m} |\lambda_i|. \quad (131)$$

504 In other words, the spectral radius of a matrix  $\mathbf{A}$  is the modulus of the largest eigenvalue.

505 **Definition 4** A matrix  $\mathbf{A}$  is called nonnegative,  $\mathbf{A} \geq \mathbf{0}$ , if  $a_{ij} \geq 0$  for all  $i, j$ .

506 Definition 4 states that a matrix is nonnegative if all elements of the matrix are greater than or equal to zero. Next, we  
507 consider primitive matrices.

508 **Definition 5** A nonnegative matrix  $\mathbf{A}$  is primitive if there is a positive integer  $k$  such that  $\mathbf{A}^k > \mathbf{0}$ .

509 Another important concept is that of the dominant eigenvalue of a matrix.

510 **Definition 6** Let  $\lambda_1, \dots, \lambda_m$  be the eigenvalues of an  $m \times m$  matrix  $\mathbf{A}$ . If  $|\lambda_1| > |\lambda_j|$  for  $j = 2, \dots, m$ , then  $\lambda_1$  is called  
511 the dominant eigenvalue of  $\mathbf{A}$ .

512 Next, we discuss the Perron-Frobenius theorem for nonnegative primitive matrices (Bapat and Raghavan 1997).

513 **Proposition B.1 (Perron-Frobenius theorem)** Let  $\mathbf{A} \geq \mathbf{0}$  be an  $m \times m$  primitive matrix. Then  $\mathbf{A}\mathbf{y} = \lambda_1\mathbf{y}$  for some  
514  $\lambda_1 > 0$ ,  $\mathbf{y} > \mathbf{0}$  where

- 515 (i) The eigenvalue  $\lambda_1$  is algebraically simple.
- 516 (ii) The eigenvalue  $\lambda_1$  is dominant. That is, for any other eigenvalue  $\mu$  of  $\mathbf{A}$ ,  $|\mu| < \lambda_1$ .
- 517 (iii) The only nonnegative eigenvectors of  $\mathbf{A}$  are positive scalar multiples of  $\mathbf{y}$ .

518 By the Perron-Frobenius theorem we know that the spectral radius of a nonnegative primitive matrix is equal to the  
519 dominant eigenvalue of that matrix;  $\rho(\mathbf{A}) = \lambda_1$ . In our analysis we also make use of the Jordan canonical form for square  
520 matrices. We use this decomposition because while a nonnegative primitive matrix is not necessarily diagonalizable, every  
521 square matrix can none-the-less be written in its Jordan canonical form.

522 **Definition 7** For any square matrix  $\mathbf{A}$ , there exists a matrix  $\mathbf{J}$  such that

$$\mathbf{A} = \mathbf{P}\mathbf{J}\mathbf{P}^{-1}, \quad (132)$$

523 where  $\mathbf{J}$  is the Jordan canonical form of  $\mathbf{A}$ . The Jordan canonical form is a block diagonal matrix

$$\mathbf{J} = \begin{bmatrix} \mathbf{J}_1 & \dots & \mathbf{0} \\ \vdots & \ddots & \vdots \\ \mathbf{0} & \dots & \mathbf{J}_b \end{bmatrix}, \quad (133)$$

524 where each  $\mathbf{J}_i$  is called a Jordan block of  $\mathbf{A}$ . For Jordan block  $i$ , the diagonal entries are  $\lambda_i$ , the superdiagonal entries are  
525 one, and all other entries are zero.

526 Next, we present two lemmas that were used in the proofs of the main theorems. The first lemma was used in Theorem  
527 3.1 and bounds our initial condition for each neutral fraction  $i$  for each stage  $j$ ,  $v_{j,0}^i(x)$ , sufficiently to establish the uniform  
528 convergence results for the neutral fractions.

529 **Lemma B.1** Let  $x \rightarrow v_{j,0}^i(x)$  satisfy  $x^2 v_{j,0}^i(x) e^{sx} \in L^1(\mathbb{R}) \cap L^\infty(\mathbb{R})$ , then for each  $s > 0$  there exists a positive constant  
530  $C_j$  such that

$$w_{j,0}^i(x) = \frac{C_j e^{-sx}}{1 + x^2} \quad (134)$$

531 bounds  $v_{j,0}^i(x)$  for all  $x \in \mathbb{R}$ . Moreover, the Fourier transform of  $w_{j,0}^i(x) e^{sx}$  with respect to  $x$  is in  $L^1(\mathbb{R})$  and is given by

$$C_j \pi e^{-|\omega|}. \quad (135)$$

532 For the proof of Lemma B.1, we refer the reader to Lemma 1 by Marculis et al. (2017).

533 We next provide a lemma that will be used in the proofs of the Theorems 3.2 and 3.3. In particular, we make use of  
534 the Jordan canonical form and the Perron-Frobenius theorem outlined above.

535 **Lemma B.2** Assume that the matrix  $\mathbf{B}_0$  is nonnegative and primitive. Let  $\lambda_1$  be the dominant eigenvalue of  $\mathbf{B}_0$ , then

$$\lim_{t \rightarrow \infty} \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t = \mathbf{r}\boldsymbol{\ell} \quad (136)$$

536 where  $\mathbf{r}$  and  $\boldsymbol{\ell}$  are the right and left eigenvectors corresponding to  $\lambda_1$  respectively with  $\boldsymbol{\ell}$  normalized by  $\langle \boldsymbol{\ell}^T, \mathbf{r} \rangle$  to account  
537 for the scaling in  $\mathbf{r}$ .

538 *Proof* Writing  $\mathbf{B}_0$  in terms of its Jordan canonical form, we have

$$\lim_{t \rightarrow \infty} \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t = \lim_{t \rightarrow \infty} \left[ \frac{\mathbf{P}\mathbf{J}\mathbf{P}^{-1}}{\lambda_1} \right]^t \quad (137)$$

$$= \lim_{t \rightarrow \infty} \frac{\mathbf{P}\mathbf{J}^t\mathbf{P}^{-1}}{\lambda_1^t}. \quad (138)$$

539 Since  $\mathbf{J}$  is block diagonal,

$$\mathbf{J}^t = \begin{bmatrix} \mathbf{J}_1^t & \dots & \mathbf{0} \\ \vdots & \ddots & \vdots \\ \mathbf{0} & \dots & \mathbf{J}_b^t \end{bmatrix}. \quad (139)$$

540 By the Perron-Frobenius theorem there exists a dominant eigenvalue  $\lambda_1$  of  $\mathbf{B}_0$  because  $\mathbf{B}_0$  is nonnegative and primitive .  
541 The first Jordan block is  $\mathbf{J}_1 = [\lambda_1]$  and  $\mathbf{J}_1^t = [\lambda_1^t]$ . For Jordan block  $j$  of size  $b_j \times b_j$  we have

$$\mathbf{J}_j^t = \begin{bmatrix} \lambda_j^t & \binom{t}{1}\lambda_j^{t-1} & \dots & \binom{t}{b_j-2}\lambda_j^{t-b_j+2} & \binom{t}{b_j-1}\lambda_j^{t-b_j+1} \\ 0 & \lambda_j^t & \dots & \binom{t}{b_j-3}\lambda_j^{t-b_j+3} & \binom{t}{b_j-2}\lambda_j^{t-b_j+2} \\ \vdots & \vdots & \ddots & \vdots & \vdots \\ 0 & 0 & \dots & \lambda_j^t & \binom{t}{1}\lambda_j^{t-1} \\ 0 & 0 & \dots & 0 & \lambda_j^t \end{bmatrix} \quad (140)$$

542 for  $t \geq b_j - 1$ . Since  $|\lambda_j| < \lambda_1$ , using L'Hôpital's rule, we have

$$\lim_{t \rightarrow \infty} \frac{\mathbf{J}_j^t}{\lambda_1^t} = \mathbf{0} \quad (141)$$

543 for  $j = 2, \dots, b$ . Returning to the Jordan canonical form,

$$\lim_{t \rightarrow \infty} \frac{\mathbf{J}^t}{\lambda_1^t} = \begin{bmatrix} 1 & \dots & 0 \\ \vdots & \ddots & \vdots \\ 0 & \dots & 0 \end{bmatrix}. \quad (142)$$

544 Hence from (138),

$$\lim_{t \rightarrow \infty} \frac{\mathbf{P}\mathbf{J}^t\mathbf{P}^{-1}}{\lambda_1^t} = \mathbf{P} \lim_{t \rightarrow \infty} \frac{\mathbf{J}^t}{\lambda_1^t} \mathbf{P}^{-1} \quad (143)$$

$$= \mathbf{P} \begin{bmatrix} 1 & \dots & 0 \\ \vdots & \ddots & \vdots \\ 0 & \dots & 0 \end{bmatrix} \mathbf{P}^{-1} \quad (144)$$

$$= \mathbf{r}\boldsymbol{\ell} \quad (145)$$

545 because  $\mathbf{r}$  is the first column vector of  $\mathbf{P}$  and  $\boldsymbol{\ell}$  is the first row vector of  $\mathbf{P}^{-1}$ . Therefore, from (138) and (145),

$$\lim_{t \rightarrow \infty} \left[ \frac{\mathbf{B}_0}{\lambda_1} \right]^t = \mathbf{r}\boldsymbol{\ell}. \quad (146)$$

The proof of Lemma B.2 is complete. □

## 546 References

- 547 Austerlitz F and Garnier-Géré P. Modelling the impact of colonisation on genetic diversity and differentiation of forest  
548 trees: interaction of life cycle, pollen flow and seed long-distance dispersal. *Heredity*, 90(4):282, 2003.  
549 Bapat RB and Raghavan TE. Nonnegative matrices and applications, volume 64. Cambridge university press, 1997.  
550 Bataillon TM, David JL, and Schoen DJ. Neutral genetic markers and conservation genetics: simulated germplasm col-  
551 lections. *Genetics*, 144(1):409–417, 1996.

552 Bateman AW, Buttenschön A, Erickson KD, and Marculis NG. Barnacles vs bullies: modelling biocontrol of the invasive  
553 european green crab using a castrating barnacle parasite. *Theor Ecol*, 10(3):305–318, 2017.

554 Bonnefon O, Garnier J, Hamel F, and Roques L. Inside dynamics of delayed travelling waves. *Math Mod Nat Phen*, 8:  
555 44–61, 2013.

556 Bonnefon O, Coville J, Garnier J, and Roques L. Inside dynamics of solutions of integro-differential equations. *Discret*  
557 *Contin Dyn Syst - Ser B*, 19(10):3057–3085, 2014.

558 Cooley JW and Tukey JW. An algorithm for the machine calculation of complex fourier series. *Math Comput*, 19(90):  
559 297–301, 1965.

560 Cullingham CI, Cooke JE, Dang S, Davis CS, Cooke BJ, and Coltman DW. Mountain pine beetle host-range expansion  
561 threatens the boreal forest. *Mol Ecol*, 20(10):2157–2171, 2011.

562 Davis MB and Shaw RG. Range shifts and adaptive responses to quaternary climate change. *Science*, 292(5517):673–679,  
563 2001.

564 Easterling MR, Ellner SP, and Dixon PM. Size-specific sensitivity: applying a new structured population model. *Ecology*,  
565 81(3):694–708, 2000.

566 Garnier J and Lewis MA. Expansion under climate change: the genetic consequences. *Bull Math Biol*, 78(11):2165–2185,  
567 2016.

568 Garnier J, Giletti T, Hamel F, and Roques L. Inside dynamics of pulled and pushed fronts. *J des Math Pures Appl*, 98  
569 (4):428–449, 2012.

570 Hallatschek O and Nelson DR. Gene surfing in expanding populations. *Theor Popul Biol*, 73(1):158–170, 2008.

571 Hastings A, Cuddington K, Davies KF, Dugaw CJ, Elmendorf S, Freestone A, Harrison S, Holland M, Lambrinos J,  
572 Malvadkar U, et al. The spatial spread of invasions: new developments in theory and evidence. *Ecol Lett*, 8(1):91–101,  
573 2005.

574 Hewitt G. The genetic legacy of the quaternary ice ages. *Nature*, 405(6789):907, 2000.

575 Holderegger R, Kamm U, and Gugerli F. Adaptive vs. neutral genetic diversity: implications for landscape genetics. *Landscape*  
576 *Ecol*, 21(6):797–807, 2006.

577 Howe HF and Smallwood J. Ecology of seed dispersal. *Annu Rev Ecol Syst*, 13(1):201–228, 1982.

578 Ibrahim KM, Nichols RA, and Hewitt GM. Spatial patterns of genetic variation generated by different forms of dispersal  
579 during range expansion. *Heredity*, 77(3):282, 1996.

580 Kot M, Lewis MA, and van den Driessche P. Dispersal data and the spread of invading organisms. *Ecology*, 77(7):  
581 2027–2042, 1996.

582 Lefkovich L. The study of population growth in organisms grouped by stages. *Biometrics*, pages 1–18, 1965.

583 Leslie PH. On the use of matrices in certain population mathematics. *Biometrika*, 33(3):183–212, 1945.

584 Levin LA. Recent progress in understanding larval dispersal: new directions and digressions. *Integr Comp Biol*, 46(3):  
585 282–297, 2006.

586 Lewis MA, Marculis NG, and Shen Z. Integrodifference equations in the presence of climate change: persistence criterion,  
587 travelling waves and inside dynamics. *J Math Biol*, 77(6-7):1649–1687, 2018.

588 Li B, Lewis MA, and Weinberger HF. Existence of traveling waves for integral recursions with nonmonotone growth  
589 functions. *J Math Biol*, 58(3):323–338, 2009.

590 Liebhold AM, Halverson JA, and Elmes GA. Gypsy moth invasion in north america: a quantitative analysis. *J Biogeogr*,  
591 pages 513–520, 1992.

592 Lubina JA and Levin SA. The spread of a reinvading species: range expansion in the california sea otter. *Am Nat*, 131(4):  
593 526–543, 1988.

594 Lui R. A nonlinear integral operator arising from a model in population genetics i. monotone initial data. *SIAM J Math*  
595 *Anal*, 13(6):913–937, 1982a.

596 Lui R. A nonlinear integral operator arising from a model in population genetics ii. initial data with compact support.  
597 *SIAM J Math Anal*, 13(6):938–953, 1982b.

598 Lui R. Existence and stability of travelling wave solutions of a nonlinear integral operator. *J Math Biol*, 16(3):199–220,  
599 1983.

600 Lui R. Biological growth and spread modeled by systems of recursions. i. mathematical theory. *Math Biosci*, 93(2):269–295,  
601 1989a.

602 Lui R. Biological growth and spread modeled by systems of recursions. ii. biological theory. *Math Biosci*, 93(2):297–311,  
603 1989b.

604 Lutscher F and Lewis MA. Spatially-explicit matrix models. *J Math Biol*, 48(3):293–324, 2004.

605 Marculis NG and Lui R. Modelling the biological invasion of *carcinus maenas* (the european green crab). *J Biol Dyn*, 10  
606 (1):140–163, 2016.

607 Marculis NG, Lui R, and Lewis MA. Neutral genetic patterns for expanding populations with nonoverlapping generations.  
608 *Bull Math Biol*, 79(4):828–852, 2017.

609 Moloney KA. A generalized algorithm for determining category size. *Oecologia*, 69(2):176–180, 1986.

610 Neubert MG and Caswell H. Demography and dispersal: calculation and sensitivity analysis of invasion speed for structured  
611 populations. *Ecology*, 81(6):1613–1628, 2000.

612 Pluess AR. Pursuing glacier retreat: genetic structure of a rapidly expanding *larix decidua* population. *Mol Ecol*, 20(3):  
613 473–485, 2011.

614 Roques L, Hosono Y, Bonnefon O, and Boivin T. The effect of competition on the neutral intraspecific diversity of invasive  
615 species. *J Math Biol*, 71(2):465–489, 2015.

616 Roques L, Garnier J, Hamel F, and Klein EK. Allee effect promotes diversity in traveling waves of colonization. *Proc Natl*  
617 *Acad Sci*, 109(23):8828–8833, 2012.

618 Vandermeer J. Choosing category size in a stage projection matrix. *Oecologia*, 32(1):79–84, 1978.

619 Veit RR and Lewis MA. Dispersal, population growth, and the allee effect: dynamics of the house finch invasion of eastern  
620 north america. *Am Nat*, 148(2):255–274, 1996.

621 Weinberger HF. Asymptotic behavior of a model in population genetics. In *Nonlinear Partial Differential Equations and*  
622 *Applications*, pages 47–96. Springer, 1978.

623 Weinberger HF. Long-time behavior of a class of biological models. *SIAM J Math Anal*, 13(3):353–396, 1982.