

BILEL ELBETCH 🕩 (Saïda)

## Effect of dispersal in single-species discrete diffusion systems with source-sink patches

Abstract A multi-patch source-sink model with and without intraspecific competition in the sink patches is considered. First, we study the dynamics of the model when the matrix of migration is irreducible and reducible. We show that, there is a threshold number of source patches such that the population potentially becomes extinct below the threshold and established above the threshold. Next, used the theory of singular perturbation and theorem of Tikhonov, in the case of perfect mixing, i.e. when the diffusion rate tends to infinity, we calculate the equilibrium of the model and we give a good approximations of the solutions in this case. Second, we determine, in some particular cases, the conditions under which fragmentation and the existence of sink patches can lead to a total equilibrium population greater or smaller than the sum of the carrying capacities of the source patches. Finally, we study the effect of the rapid growth of the population in source patches and the rapid death of the population in sink patches on the dynamics of the total equilibrium population and on the coexistence of the populations in the patches.

2010 Mathematics Subject Classification: Primary: 37N25; Secondary: 34D23, 34D15, 92D25..

*Key words and phrases:* Population Dynamics, Logistic equation, diffusion rate, Slow-fast systems, Tikhonov's theorem, Perfect mixing..

1. Introduction Population dynamics is a wide field of mathematics, which contains many problems, for example fragmentation of population and the effect of migration in the general dynamics of population. Bibliographies can be found in the work of Levin [29, 30] and Holt [26]. There are ecological situations that motivate the representation of space as a finite set of patches connected by migrations, for instance an archipelago with bird population and predators. It is an example of insular bio-geography. A reference work on mathematical models is the book of Levin et al. [31], whereas Hanski and Gaggiotti [24] give a more ecological account of the subject. The standard question in this type of biomathematical problems, is to study the effect of migration on the general population dynamics, and the consequences of fragmentation on the persistence or extinction of the population.

An ecological model is used to describe population changes in two habitats, both occupied by the same species. One habitat is of high quality and allows a population to increase, leading to a surplus. This is the source. The other habitat is of low quality, leading to a deficit that ordinarily would lead to the habitat being abandoned. This is the sink. The source-sink model was first proposed by biologist Crick 1970. In population ecology, the source-sink model is used to describe how variation in habitat quality may affect the population growth or decline of organisms.

In 2019, Wu et al. [39] studied the following two-patch source-sink model:

$$\begin{cases} \frac{dx_1}{dt} = r_1 x_1 \left( 1 - \frac{x_1}{K_1} \right) + D(x_2 - s x_1), \\ \frac{dx_2}{dt} = r_2 x_2 \left( -1 - \frac{x_2}{K_2} \right) + D(s x_1 - x_2), \end{cases}$$
(1)

where  $x_1$  and  $x_2$  represent population densities of the species in patch 1 and 2, respectively. The parameters  $r_i > 0$  and  $K_i > 0$  are respectively the intrinsic growth rate and the carrying capacity of patch *i*. Parameter *D* represents the dispersal intensity while the parameter *s* reflects the dispersal asymmetry. The authors show that the dispersal asymmetry can lead to either an increased total size of the population in two patches, a decreased total size with persistence in the patches, or even extinction in both patches. They show also that for a large growth rate of the species in the source and a fixed dispersal intensity:

- If the asymmetry is small, the population would persist in both patches and reach a density higher than that without dispersal and the population approaches its maximal density at an appropriate asymmetry.
- If the asymmetry is intermediate, the population persists in both patches but reaches a density less than that without dispersal.
- If the asymmetry is large, the population goes to extinction in both patches.

Arino et al. [4] also studied a source-sink model of n patches, where the source patch follows a logistic growth rate, and the sink patch follows exponential decay, i.e the model

$$\begin{cases} \frac{dx_i}{dt} = r_i x_i \left(1 - \frac{x_i}{K_i}\right) + D \sum_{j=1}^n \gamma_{ij} x_j, \quad i = 1, \dots, s, \\ \frac{dx_i}{dt} = -r_i x_i + D \sum_{j=1}^n \gamma_{ij} x_j, \quad i = s+1, \dots, n, \end{cases}$$
(2)

where  $x_i$  represent population densities of the species in the patch *i*. The parameter *D* represents the dispersion rate of the population,  $\gamma_{ij} \ge 0$  denote the flux between patches *j* and *i* for  $i \ne j$ . We denote  $\Gamma$  the matrix  $\Gamma := (\gamma_{ij})_{n \times n}$  with

$$\gamma_{ii} = -\sum_{j=1, j \neq i}^{n} \gamma_{ji}.$$
(3)

For system (2), the authors proved the existence of a threshold number of source patches such that the population potentially becomes extinct below the threshold and persistent above the threshold.

Elbetch et al. [14, 15] have answered in the particular case of the n-source 0-sink patch model (i.e System (2) for n = s), which reads:

$$\frac{dx_i}{dt} = r_i x_i \left(1 - \frac{x_i}{K_i}\right) + D \sum_{j=1}^n \gamma_{ij} x_j, \qquad i = 1, \dots, n,$$
(4)

to the following important question:

QUESTION 1.1 Is it possible, depending on the migration rate D, that the total equilibrium population  $X_T^*(D) = \sum_i x_i^*(D)$ , where  $(x_1^*(D), \ldots, x_n^*(D))$  is the positive equilibrium of (4) be larger than the sum of the capacities  $\sum_i K_i$ ?

Note that, System (4) is studied also by Elbetch et al. [14] and Takeuchi [36] in the case when the matrix  $\Gamma$  is symmetric, i.e. the flux between patches jand i for  $i \neq j$  is equal in both directions. We recall that, when the matrix of migration  $\Gamma$  is irreducible, System (4) admits a unique positive equilibrium which is globally asymptotically stable (GAS), see [3, Theorem 2.2], [4, Theorem 1] or [14, Theorem 6.1], when  $D \to \infty$ , this equilibrium tends to

$$\frac{\sum_i \delta_i r_i}{\sum_i \delta_i^2 \alpha_i} (\delta_1, \dots, \delta_n),$$

where  $\alpha_i = \frac{r_i}{K_i}$  and  $(\delta_1, \ldots, \delta_n)^T$  is the vector which generate the vector space ker  $\Gamma$ . For the existence, uniqueness, and positivity of  $\delta$  see Remark **3.1.** Question 1.1 is of ecological importance since the answer gives the conditions under which dispersal is either beneficial or detrimental to total equilibrium population. Note that, this last question has been studied by many researches (see [1, 2, 5, 8, 9, 10, 11, 14, 15, 17, 18, 20, 21, 28, 41], [13] for effect of nonlinear diffusion on the total biomass, and [22, 23] for susceptibleinfected-susceptible (SIS) patch-model). Elbetch et al. [14] proved that, if all the patches do not differ with respect to the intrinsic growth rate (i.e.,  $r_1 = \ldots = r_n$ ), then the effect of migration is always detrimental. In the case when  $(K_1, \ldots, K_n)^T \in \ker \Gamma$  (if the matrix  $\Gamma$  is symmetric, the condition  $(K_1,\ldots,K_n)^T \in \ker \Gamma$  means that the patches do not differ with respect to the carrying capacity), migration has no effect on the total equilibrium population. An example when the effect of migration is always beneficial, is in the case when  $\Gamma$  is symmetric and all the patches do not differ with respect to the parameter  $\alpha = r/K$  quantifying intraspecific competition (i.e.,  $\alpha_1 = \ldots, \alpha_n$ ) (see also [15, Prop. 4.2] for another example when  $\Gamma$  is non symmetric).

It was shown by Arditi et al. [1, Proposition 2, page 54], for 2-source, 0-sink patch model, that only three situations can occur: the case where the total

equilibrium population is always greater than the sum of carrying capacities, the case where it is always smaller, and a third case, where the effect of migration is beneficial for lower values of the migration coefficient D and detrimental for the higher values. More precisely, it was shown in [1] that, if n = 2 in (4), the following trichotomy holds

- If  $X_T^*(+\infty) > K_1 + K_2$  then  $X_T^*(D) > K_1 + K_2$  for all D > 0.
- If  $\frac{dX_T^*}{dD}(0) > 0$  and  $X_T^*(+\infty) < K_1 + K_2$ , then there exists  $D_0 > 0$  such that  $X_T^*(D) > K_1 + K_2$  for  $0 < D < D_0$ ,  $X_T^*(D) < K_1 + K_2$  for  $D > D_0$  and  $X_T^*(D_0) = K_1 + K_2$ .
- If  $\frac{dX_T^*}{dD}(0) < 0$ , then  $X_T^*(D) < K_1 + K_2$  for all D > 0.

Therefore, the condition  $X_T^*(D) = K_1 + K_2$  holds only for D = 0 and at most for one positive value  $D = D_0$ . The value  $D_0$  exists if and only if  $\frac{dX_T^*}{dD}(0) > 0$ and  $X_T^*(+\infty) < K_1 + K_2$ .

In [14, Section 5.2], Elbetch et al. have considered Model (4) for n = 3 with  $\Gamma$  symmetric, and shown by numerical simulations the following situations, which do not exist in the two-patch model:

- The case where  $\frac{dX_T^*}{dD}(0) < 0$  and  $X_T^*(+\infty) > K_1 + K_2 + K_3$ .
- The case where  $\frac{dX_T^*}{dD}(0) > 0$  and  $X_T^*(+\infty) > K_1 + K_2 + K_3$  and there exist values of D for which  $X_T^*(D) < K_1 + K_2 + K_3$ .
- The case where  $\frac{dX_T^*}{dD}(0) < 0$  and  $X_T^*(+\infty) < K_1 + K_2 + K_3$  and there exist values of D for which  $X_T^*(D) > K_1 + K_2 + K_3$ .

Therefore the equality  $X_T^*(D) = K_1 + K_2 + K_3$  can occur for two positive values of D, not only for a unique positive value as in the two-patch case.

In [15, Section 6], Elbetch et al. have reconsidered the three-patch model with  $\Gamma$  not symmetric. The novelty when  $\Gamma$  is not symmetric is the existence of three positive values of migration rate solution of the following equation:

Total equilibrium population = Sum of three carrying capacities,

i.e. the following situation hold:

• The case where  $\frac{dX_T^*}{dD}(0) > 0$  and  $X_T^*(+\infty) < K_1 + K_2 + K_3$ , and there exists three values  $0 < D_1 < D_2 < D_3$  for which we have:

$$X_T^*(D) = \begin{cases} > K_1 + K_2 + K_3 & \text{for } D \in ]0, D_1[\cup]D_2, D_3[, \\ < K_1 + K_2 + K_3 & \text{for } D \in ]D_1, D_2[\cup]D_3, \infty[. \end{cases}$$

For more details and the proof of some previous numerical results, see the recent work of Elbetch [12], where it is studied the model (4) under the assumption that some growth rates are much larger than the other. Recently, Yu et al. [40] considered a consumer-resource patch model, where the consumer moves between multiple source-sink patches with both resource and toxicant given by the following system:

$$\begin{cases}
\frac{dx_i}{dt} = N_{0i} - \mu_i x_i - \frac{r_i x_i y_i}{\Gamma_i (k_i + x_i)}, & i = 1, \dots, n, \\
\frac{dy_i}{dt} = y_i \left(\frac{r_i x_i}{k_i + x_i} - m_i - g_i y_i\right) - D\left(s_i y_i - \frac{1}{2}s_{i-1} y_{i-1} - \frac{1}{2}s_{i+1} + y_{i+1}\right),
\end{cases} (5)$$

where  $i = i \mod n$ , and "mod" means modula [42]. For example,  $y_0 = y_n$ and  $y_{n+1} = y_1$ . Variable  $x_i$  represents the nutrient concentration and  $y_i$  is the consumer's population density in patch *i*. Parameter  $N_{0i}$  represents the nutrient input,  $\mu_i$  is the dilution rate of nutrient,  $\Gamma_i$  is the yield, or fraction of nutrient per unit biomass. Parameter  $r_i$  represents the consumer's maximal growth rate with infinite resource,  $k_i$  is the half saturation coefficient,  $m_i$ is the mortality rate, and  $g_i$  is the density-dependent loss rate. Parameter D represents the diffusion rate, while  $s_i$  is the asymmetry in diffusion. Note that, when  $s_i = 1$  for all *i*, the diffusion is symmetric. Yu et al. [40] showed the global stability of positive equilibria in System (5). They have shown also that diffusion could make the consumer persists in sinks, even make it reaches total population abundance larger than if non-diffusing. It is also shown that under certain conditions, diffusion could make the total abundance less than if non-diffusing, even make the consumer go into extinction in all patches.

An important result proven by Yu et al. [40] is that when toxicants are distributed homogeneously, asymmetric diffusion always makes the total abundance less than if non-diffusing. For general information on the effect of asymmetric diffusion, toxicant distribution, and geographic pattern of patches on the total population abundance of the consumer, and also in the continuous and discrete cases of (5), the reader is referred to the work of Yu et al. [40] and Zhang et al. [42].

Our aim of the present paper, is to study the effect of the migration on the total population with the assumption that some patches among the n patches are sinks. Thus we generalize some results of [14, 15] for n-source, 0-sink patch model to s-source, (n - s)-sink patch model and also we extend the results proved by Wu et al. [39] for 1-source, 1-sink patch model.

The paper is organized as follows. In Section 2, some proprieties of 1source, 1-sink patch model (1) have been recalled as a function of the two parameters  $\gamma_1$  and  $\gamma_2$  (see Theorem 2.2). Two-patch model with the growth (resp. death) rate much larger than the death (resp. growth) rate is considered. In both last cases, we compare the total equilibrium population with the capacity (see Theorems 2.6 and 2.10). In Section 3, multi-patch source-sink model with intraspecific competition in the sink patches is being described. We prove that there exists a threshold number of source patches such that the population becomes extinct below the threshold and persist above the threshold (Theorem 3.3). The behavior of the model for large migration rate is studied (Theorem 3.7). Total population abundance is analyzed also in some homogeneous and heterogeneous particular case (Propositions 3.11, 3.12, and 3.16). The following both cases: death rates are much larger than the growth rates and growth rates are much larger than the death rates are considered. In Section 4, multi-patch source-sink model without intraspecific competition in the sink patches is considered. In Appendix A, we give some properties of the total equilibrium population. In Appendix B, we give some background concepts and preliminaries results which are used in the analysis of the global stability of our model.

2. Some preliminary results for the two-patch model with sourcesink patches In this section, we consider the 2-patch system with sourcesink dynamics given by:

$$\begin{cases} \frac{dx_1}{dt} = a_1 x_1 \left( 1 - \frac{x_1}{L_1} \right) + D \left( \gamma_2 x_2 - \gamma_1 x_1 \right), \\ \frac{dx_2}{dt} = a_2 x_2 \left( -1 - \frac{x_2}{L_2} \right) + D \left( \gamma_1 x_1 - \gamma_2 x_2 \right), \end{cases}$$
(6)

where  $x_1$  and  $x_2$  represent population densities of the species in patch 1 and 2, respectively. Patch 1 is assumed to be the source but patch 2 is the sink, i.e.  $a_1, a_2 > 0$ . The parameters  $\alpha_i := a_i/L_i$  are the intraspecific competition degree. Parameter D represents the dispersal intensity. We denote  $\gamma_2$  the migration rate from source patch 2 to the sink patch 1 and  $\gamma_1$  from sink patch 1 to source patch 2, the dispersal is symmetric if  $\gamma_1 = \gamma_2$ . This system is studied in [39]. We recall some essential results of [39] as function of the parameters  $\gamma_1$  and  $\gamma_2$ . First of all, let's start by recalling the global dynamics of System (6).

**2.1. Global dynamics** We consider the following regions in the set of parameters  $\gamma_1$  and  $\gamma_2$ , denoted  $\mathcal{D}_0$ ,  $\mathcal{D}_1$  and  $\mathcal{D}_2$  depicted in Figure 1 and defined by:

$$\begin{cases}
\mathcal{D}_{0} = \left\{ (\gamma_{1}, \gamma_{2}) : \quad \gamma_{2} \geq \frac{a_{2}}{a_{1}} \gamma_{1} \right\}, \\
\mathcal{D}_{1} = \left\{ (\gamma_{1}, \gamma_{2}) : \quad \frac{a_{2}}{a_{1}} \gamma_{1} < \gamma_{2} < \frac{a_{2}D}{a_{1}a_{2} + Da_{1}} \gamma_{1} \right\}, \\
\mathcal{D}_{2} = \left\{ (\gamma_{1}, \gamma_{2}) : \quad \gamma_{2} \geq \frac{a_{2}D}{a_{1}a_{2} + Da_{1}} \gamma_{1} \right\}.
\end{cases}$$
(7)

The global dynamic of System (6) is described as follows.

THEOREM 2.1 (PROP. 5.5 IN [39]) Consider Model (6). Then, if  $(\gamma_1, \gamma_2) \in \mathcal{D}_0 \cup \mathcal{D}_1$ , System (6) admits unique equilibrium in  $\mathbb{R}^2 \setminus \{0\}$  denoted  $E^*(D)$ , which is GAS, and if  $(\gamma_1, \gamma_2) \in \mathcal{D}_2$ , then the origin is GAS.



Figure 1: Global stability of Model (6). In  $\mathcal{D}_0$  and  $\mathcal{D}_1$  the system has unique equilibrium  $E^*(D)$  which is GAS. In the region  $\mathcal{D}_2$ , the system has the origin as unique equilibrium which is GAS.

**2.2. Total population abundance** In this section, we recall the comparison given in [39, Proposition 5.11], between the total equilibrium population

$$X_T^*(D) = x_1^*(D) + x_2^*(D), \qquad E^*(D) = (x_1^*(D), x_2^*(D)),$$

of (6) and carrying capacity  $L_1$ , by analyzing the stable positive equilibrium  $E^*(D)$ . Note that, when there is no dispersal (i.e., D = 0), the total equilibrium population is  $X_T^*(0) = L_1$ . We consider the regions in the set of the parameters  $\gamma_1$  and  $\gamma_2$ , denoted  $\mathcal{L}_0$ ,  $\mathcal{L}_1$ ,  $\mathcal{L}_2$ ,  $\mathcal{L}_3$  and  $\mathcal{L}_4$ , depicted in Figure 2 and defined by:

$$\begin{cases} \text{If } a_{2} \geq a_{1} \text{ then } \begin{cases} \mathcal{L}_{0} = \left\{ (\gamma_{1}, \gamma_{2}) : \frac{\gamma_{2}}{\gamma_{1}} < \frac{a_{2}}{a_{1}} \right\}, \\ \mathcal{L}_{1} = \left\{ (\gamma_{1}, \gamma_{2}) : \frac{\gamma_{2}}{\gamma_{1}} \geq \frac{a_{2}}{a_{1}} \right\}, \end{cases} \end{cases}$$

$$(8)$$

$$\text{If } a_{2} < a_{1} \text{ then } \begin{cases} \mathcal{L}_{2} = \left\{ (\gamma_{1}, \gamma_{2}) : \frac{\gamma_{2}}{\gamma_{1}} \leq \frac{a_{2}}{a_{1}} \right\}, \\ \mathcal{L}_{3} = \left\{ (\gamma_{1}, \gamma_{2}) : \frac{a_{2}}{a_{1}} < \frac{\gamma_{2}}{\gamma_{1}} < \frac{L_{2}(a_{1}-a_{2})}{a_{2}(L_{1}+L_{2})} \right\}, \\ \mathcal{L}_{4} = \left\{ (\gamma_{1}, \gamma_{2}) : \frac{\gamma_{2}}{\gamma_{1}} \geq \frac{L_{2}(a_{1}-a_{2})}{a_{2}(L_{1}+L_{2})} \right\}. \end{cases}$$

THEOREM 2.2 The total equilibrium population described by (6) satisfies the following properties

1. If  $a_2 \ge a_1$ , let  $\mathcal{L}_0$  and  $\mathcal{L}_1$  be defined by (8). Then we have:



Figure 2: Qualitative properties of source-sink model (6). In  $\mathcal{L}_0$  and  $\mathcal{L}_1$  the effect is detrimental with extinction in two patches for  $\mathcal{L}_0$  and persistence for  $\mathcal{L}_1$ . In  $\mathcal{L}_2$  and  $\mathcal{L}_3$ , the effect is beneficial for  $D < D_0$  and detrimental for  $D > D_0$  with persistence of the population in the region  $\mathcal{L}_2$  and extinction in the region  $\mathcal{L}_3$ . In  $\mathcal{L}_4$ , patchiness has a beneficial effect on the total equilibrium population.

• if  $(\gamma_1, \gamma_2) \in \mathcal{L}_0$  then  $X_T^*(D) \leq L_1$  for all  $D \geq 0$ . More over, there is  $D^* = \frac{\gamma_2 a_1 a_2}{\gamma_1 a_2 - \gamma_2 a_1}$ , such that:

$$\begin{cases} 0 < X_T^*(D) \le L_1 & \text{ If } D < D^*, \\ X_T^*(D) = 0 & \text{ If } D \ge D^*. \end{cases}$$
(9)

• if  $(\gamma_1, \gamma_2) \in \mathcal{L}_1$  then  $0 < X_T^*(D) \le L_1$  for all  $D \ge 0$ .

2. If  $a_2 < a_1$ , let  $\mathcal{L}_2, \mathcal{L}_3$  and  $\mathcal{L}_4$  be defined by (8). Then we have:

• if  $(\gamma_1, \gamma_2) \in \mathcal{L}_2$  then  $X_T^*(D) > L_1$  for  $D < D_0$  and  $X_T^*(D) < L_1$ for all  $D > D_0$ , where

$$D_0 = \frac{(a_1 - a_2)(L_1 + L_2)}{(\gamma_2(a_2 - a_1) + \gamma_1\alpha_1(L_1 + L_2))(\alpha_1^{-1} + \alpha_2^{-1})}, \quad with \ \alpha_i = a_i/L_i$$
(10)

Moreover, there is  $D^* \ge D_0$  such that  $X_T^*(D) = 0$  for all  $D \ge D^*$ .

• if  $(\gamma_1, \gamma_2) \in \mathcal{L}_3$  then we have

$$\begin{cases} X_T^*(D) \ge L_1 & \text{If } D \le D^*, \\ 0 < X_T^*(D) < L_1 & \text{If } D > D^*. \end{cases}$$
(11)

• if  $(\gamma_1, \gamma_2) \in \mathcal{L}_4$ , then  $X_T^*(D) \ge L_1$  for all  $D \ge 0$ .

PROOF All the results were established by Wu at al. [39, Proposition 5.11]. Note that, the explicit expression (10) of  $D_0$  was not given in [39].

In biological terms, the results of the previous theorem for 1-source 1-sink patch shows that, the dispersal asymmetry can lead to an increased total size of the species in two patches, a decreased total size with persistence in the patches, and even extinction in both patches. Comparing these results with that of Arditi et al. [1, 2] for 1-source 1-source patch model, we deduce that the existence of a sink patch among the two patches, can cause an extinction of the total population in the two patches.

In the case of perfect mixing (i.e  $D \to \infty$ ), we have the following result [39, Proposition 5.10]:

**PROPOSITION 2.3** We have:

$$X_{T}^{*}(\infty) := \lim_{D \to \infty} X_{T}^{*}(D) = \begin{cases} (\gamma_{1} + \gamma_{2}) \frac{\gamma_{2}a_{1} - \gamma_{1}a_{2}}{\gamma_{2}^{2}a_{1}/L_{1} + \gamma_{1}^{2}a_{2}/L_{2}} & if \quad \gamma_{1}/\gamma_{2} < a_{1}/a_{2}, \\ 0 \quad otherwise. \end{cases}$$
(12)

Wu et al. [39] proved that large dispersal intensity (i.e.,  $D \to \infty$ ), the intermediate asymmetry  $\gamma_1/\gamma_2$  can lead to population density higher than that without dispersal, and extremely small asymmetry is still favorable, while extremely large asymmetry is unfavorable: (i) When the dispersal asymmetry is small, the species can approach a density larger than that without dispersal, while it reaches its maximum value at an intermediate asymmetry  $\frac{\gamma_1}{\gamma_2} = \frac{a_1-a_2}{2\alpha_2(L_1+L_2)}$ . (ii) When  $\gamma_1/\gamma_2$  is extremely large, the species goes to extinction in both patches. Mathematically speaking, we can rewrite the following result [39, Proposition 5.10]:

PROPOSITION 2.4 [39, Proposition 5.10] Assume that  $\frac{\gamma_1}{\gamma_2} < \frac{a_1}{a_2}$ . Consider the total equilibrium population for  $D \to \infty$  given by (12). We have:

$$X_T^*(+\infty) \begin{cases} > L_1 & if \quad \frac{\gamma_1}{\gamma_2} < \frac{a_1 - a_2}{\alpha_2(L_1 + L_2)}, \\ = L_1 & if \quad \frac{\gamma_1}{\gamma_2} = \frac{a_1 - a_2}{\alpha_2(L_1 + L_2)}, \\ < L_1 & if \quad \frac{\gamma_1}{\gamma_2} > \frac{a_1 - a_2}{\alpha_2(L_1 + L_2)}. \end{cases}$$
(13)

Moreover,  $X_T^*(+\infty)$  approaches its maximum value

$$\gamma_2^2 a_2 L_1 \frac{L_1 + L_2}{4(\gamma_2^2 a_1 L_2 + \gamma_1^2 a_2 L_1)} \left(\frac{a_1 - a_2}{\alpha_2(L_1 + L_2)}\right)^2$$

 $at \frac{\gamma_1}{\gamma_2} = \frac{a_1 - a_2}{2\alpha_2(L_1 + L_2)}.$ 

In the remainder of this section, we present our first result in this work. More precisely, we study the effect of the rapid growth of the population in the source patch and rapid death of the population in the sink patch on the dynamics of the total equilibrium population and on the coexistence of the population in both patches. Note that, these situations were not examined in [4, 39]. Here, we study the total equilibrium population as a function of the migration rate in the case where the growth (resp. death) rate is much larger than the death (resp. growth) rate. In particular, we explicitly calculate the total equilibrium in the both situations, its derivative in the absence of the migration, its limit for large migration rate and we compare the total equilibrium population with the carrying capacity of the source patch. First, we start by the following situation:

**2.3. The death rate is much larger than the growth rate** In this part, we consider the two-patch model (6) and we assume that the death rate  $a_2$  is much larger than the growth rate  $a_1$ . One can write the model in the following way:

$$\begin{cases} \frac{dx_1}{dt} = a_1 x_1 \left( 1 - \frac{x_1}{L_1} \right) + D \left( \gamma_2 x_2 - \gamma_1 x_1 \right), \\ \frac{dx_2}{dt} = \frac{a_2}{\epsilon} x_2 \left( -1 - \frac{x_2}{L_2} \right) + D \left( \gamma_1 x_1 - \gamma_2 x_2 \right), \end{cases}$$
(14)

where  $\epsilon$  is assumed to be a small positive number. First, we have the following result:

THEOREM 2.5 Let  $(x_1(t, \epsilon), x_2(t, \epsilon))$  be the solution of System (14) with initial condition  $(x_1^0, x_2^0)$  satisfying  $x_i^0 \ge 0$  for i = 1, 2. Let z(t) be the solution of the differential equation

$$\frac{dx_1}{dt} = a_1 x_1 \left( 1 - \frac{x_1}{L_1} \right) - D\gamma_1 x_1 =: \varphi(x_1),$$
(15)

with initial condition  $z(0) = x_1^0$ . Then, when  $\epsilon \to 0$ , we have

$$x_1(t,\epsilon) = z(t) + o_{\epsilon}(1),$$
 uniformly for  $t \in [0, +\infty)$  (16)

and, for any  $t_0 > 0$ , we have

$$x_2(t,\epsilon) = o_{\epsilon}(1), \quad uniformly for \quad t \in [t_0, +\infty).$$
 (17)

PROOF When  $\epsilon \to 0$ , System (14) is a slow-fast system, with one slow variable,  $x_1$ , and one fast variable,  $x_2$ . Tikhonov's theorem [32, 37, 38] prompts us to consider the dynamics of the fast variables in the time scale  $\tau = \frac{1}{\epsilon}t$ . One obtains

$$\frac{dx_2}{d\tau} = a_2 x_2 \left( -1 - \frac{x_2}{L_2} \right) + \epsilon D(\gamma_1 x_1 - \gamma_2 x_2).$$
(18)

In the limit  $\epsilon \to 0$ , we find the *fast dynamics* 

$$\frac{dx_2}{d\tau} = a_2 x_2 \left(-1 - \frac{x_2}{L_2}\right). \tag{19}$$

The slow manifold is given by the equilibrium of System (19), i.e.  $x_2 = 0$ , which is locally asymptotically stable (LAS) in the positive axis. When  $\epsilon$  goes to zero, Tikhonov's theorem ensures that after a fast transition toward the slow manifold, the solutions of (14) converge to the solutions of the *reduced* model (15), obtained by replacing  $x_2 = 0$  into the dynamics of the slow variable.

If  $a_1 - D\gamma_1 \leq 0$ , then, the differential equation (15) admits  $x_1^*(D, 0^+) = 0$  for all D, as equilibrium, which is LAS. If  $a_1 - D\gamma_1 > 0$ , then, the differential equation (15) admits as a positive equilibrium

$$x_1^*(D,0^+) := \frac{L_1(a_1 - D\gamma_1)}{a_1}.$$
(20)

As  $\varphi(x_1) > 0$  for all  $0 \le x_1 < x_1^*(D, 0^+)$  and  $\varphi(x_1) < 0$  for all  $x_1 > x_1^*(D, 0^+)$ then, the equilibrium  $x_1^*(D, 0^+)$  is LAS in the positive axis, so, the approximation given by Tikhonov's theorem holds for all  $t \ge 0$  for the slow variable and for all  $t \ge t_0 > 0$  for the fast variable, where  $t_0$  is as small as we want. Therefore, let z(t) be the solution of the reduced model (15) of initial condition  $z(0) = x_1^0$ , then, when  $\epsilon \to 0$ , we have Approximations (16) and (17).

We have the following result which gives the conditions for which patchiness is beneficial or detrimental in System (14) when  $\epsilon$  goes to zero.

THEOREM 2.6 Consider the total equilibrium population  $x_1^*(D, 0^+)$  of System (14) when  $\epsilon \to 0$ , given by (20). Then,  $0 < x_1^*(D, 0^+) < L_1$  for  $D < \frac{\gamma_1}{a_1}$ , and  $x_1^*(D, 0^+) = 0$  for  $D \ge \frac{\gamma_1}{a_1}$ .

**2.4. The growth rate is much larger than the death rate** In this part, we consider the two-patch model (6) and we assume that the growth rate  $a_1$  is much larger than the death rate  $a_2$ . On can write the model in the following way:

$$\begin{cases} \frac{dx_1}{dt} = \frac{a_1}{\epsilon} x_1 \left( 1 - \frac{x_1}{L_1} \right) + D\left( \gamma_2 x_2 - \gamma_1 x_1 \right), \\ \frac{dx_2}{dt} = a_2 x_2 \left( -1 - \frac{x_2}{L_2} \right) + D\left( \gamma_1 x_1 - \gamma_2 x_2 \right), \end{cases}$$
(21)

where  $\epsilon$  is assumed to be a small positive number. We prove the following result:

THEOREM 2.7 Let  $(x_1(t, \epsilon), x_2(t, \epsilon))$  be the solution of System (21) with initial condition  $(x_1^0, x_2^0)$  satisfying  $x_i^0 \ge 0$  for i = 1, 2. Let z(t) be the solution of the differential equation

$$\frac{dx_2}{dt} = a_2 x_2 \left( -1 - \frac{x_2}{L_2} \right) + D(\gamma_1 L_1 - \gamma_2 x_2) =: \psi(x_2), \tag{22}$$

with initial condition  $z(0) = x_2^0$ . Then, when  $\epsilon \to 0$ , we have

$$x_2(t,\epsilon) = z(t) + o_{\epsilon}(1), \qquad uniformly \text{ for } t \in [0,+\infty)$$
(23)

and, for any  $t_0 > 0$ , we have

$$x_1(t,\epsilon) = L_1 + o_{\epsilon}(1), \quad uniformly for \quad t \in [t_0, +\infty).$$
(24)

PROOF When  $\epsilon \to 0$ , System (21) is a *slow-fast* system, with one *slow variable*,  $x_2$ , and one *fast variable*,  $x_1$ . Tikhonov's theorem [32, 37, 38] prompts us to consider the dynamics of the fast variables in the time scale  $\tau = \frac{1}{\epsilon}t$ . One obtains

$$\frac{dx_1}{d\tau} = a_1 x_1 \left( 1 - \frac{x_1}{L_1} \right) + \epsilon D(-\gamma_1 x_1 + \gamma_2 x_2). \tag{25}$$

In the limit  $\epsilon \to 0$ , we find the *fast dynamics* 

$$\frac{dx_1}{d\tau} = a_1 x_1 \left( 1 - \frac{x_1}{L_1} \right). \tag{26}$$

The slow manifold is given by the equilibrium of System (19), i.e.  $x_1 = L_1$ , which is LAS in the positive axis. When  $\epsilon$  goes to zero, Tikhonov's theorem ensures that after a fast transition toward the slow manifold, the solutions of (21) converge to the solutions of the *reduced model* (22), obtained by replacing  $x_1 = L_1$  into the dynamics of the slow variable.

The differential equation (22) admits as a positive equilibrium

$$x_{2}^{*}(D,0^{+}) := -\frac{L_{2}}{2} - \frac{DL_{2}}{2a_{2}}\gamma_{2} + \frac{1}{2a_{2}}\sqrt{L_{2}^{2}\gamma_{2}^{2}D^{2} + (2a_{2}L_{2}^{2}\gamma_{2} + 4a_{2}L_{2}L_{1}\gamma_{1})D + a_{2}^{2}L_{2}^{2}}$$
(27)

As  $\psi(x_2) > 0$  for all  $0 \le x_2 < x_2^*(D, 0^+)$  and  $\psi(x_2) < 0$  for all  $x_2 > x_2^*(D, 0^+)$  then, the equilibrium  $x_2^*(D, 0^+)$  is LAS in the positive axis, so, the approximation given by Tikhonov's theorem holds for all  $t \ge 0$  for the slow variable and for all  $t \ge t_0 > 0$  for the fast variable, where  $t_0$  is as small as we want. Therefore, let z(t) be the solution of the reduced model (22) with initial condition  $z(0) = x_0^0$ , then, when  $\epsilon \to 0$ , we have Approximations (23) and (24).

As a corollary of the previous theorem, we have the following result which give the limit of the total equilibrium population  $X_T^*(D, 0^+)$  of System (21) when  $\epsilon$  goes to zero:

COROLLARY 2.8 We have:  

$$X_T^*(D, 0^+) := L_1 - \frac{L_2}{2} - \frac{DL_2}{2a_2}\gamma_2 + \frac{1}{2a_2}\sqrt{L_2^2\gamma_2^2D^2 + (2a_2L_2^2\gamma_2 + 4a_2L_2L_1\gamma_1)D + a_2^2L_2^2}.$$
(28)

In the following proposition, we calculate the derivative of  $X_T^*(D, 0^+)$  at D = 0 and the formula of perfect mixing (i.e. when  $D \to \infty$ ) of the total equilibrium population defined by (28).

PROPOSITION 2.9 Consider the total equilibrium population (28). Then,

$$\frac{dX_T^*}{dD}(0,0^+) = \frac{\gamma_1 L_1}{a_2},\tag{29}$$

and

$$X_T^*(+\infty, 0^+) = \frac{\gamma_1 + \gamma_2}{\gamma_2} L_1.$$
 (30)

PROOF The derivative of the total equilibrium population  $X_T^*(D, 0^+)$  defined by (28) with respect to D is:

$$\frac{dX_T^*}{dD}(D,0^+) = -\frac{L_2\gamma_2}{2a_2} + 1/4 \frac{2a_2L_2^2\gamma_2 + 2DL_2^2\gamma_2^2 + 4a_2L_2\gamma_1L_1}{a_2\sqrt{a_2^2L_2^2 + 2a_2L_2^2}D\gamma_2 + D^2L_2^2\gamma_2^2 + 4a_2DL_2\gamma_1L_1}.$$
(31)

In particular, the derivative of the total equilibrium population at D = 0 is given by the formula (29).

By taking the limit of (28) when  $D \to \infty$ , we get that the total equilibrium population  $X_T^*(D, 0^+)$  tends to (30).

We have the following result which gives the conditions for which patchiness is beneficial or detrimental in model (21) when  $\epsilon$  goes to zero.

THEOREM 2.10 Consider the total equilibrium population  $X_T^*(D, 0^+)$  given by (28). Then,  $X_T^*(D, 0^+) \ge L_1$ , for all  $D \ge 0$ .

PROOF First, we try to solve the equation  $X_T^*(D, 0^+) = L_1$  with respect to D, the solutions of this last equation give the points of intersection between the curve of the total equilibrium population  $D \mapsto X_T^*(D, 0^+)$  and the straight line  $D \mapsto L_1$ . For any  $D \ge 0$ , we have

$$\begin{aligned} X_T^*(D, 0^+) &= L_1 \Longleftrightarrow \frac{1}{2a_2} \sqrt{L_2^2 \gamma_2^2 D^2 + (2a_2 L_2^2 \gamma_2 + 4a_2 L_2 L_1 \gamma_1) D + a_2^2 L_2^2} = \frac{L_2}{2} + \frac{DL_2}{2a_2} \gamma_2 \\ &\iff \sqrt{L_2^2 \gamma_2^2 D^2 + (2a_2 L_2^2 \gamma_2 + 4a_2 L_2 L_1 \gamma_1) D + a_2^2 L_2^2} = a_2 L_2 + \gamma_2 DL_2 \\ &\iff 4a_2 \gamma_1 L_1 L_2 D = 0 \\ &\iff D = 0. \end{aligned}$$

Therefore, since  $\frac{dX_T^*}{dD}(0,0^+) > 0$ , the curve of the total equilibrium population intersects the straight line  $D \mapsto L_1 + L_2$  in a unique point which is  $(0, L_1)$ . Therefore,  $X_T^*(D, 0^+) \ge L_1$ , for all  $D \ge 0$ .

Biologically speaking, from Sections 2.4 and 2.3, we conclude that, the rapid increase in the population in the source patch results in persistence in the both patches with increased total size population, and the rapid decrease in the sink population results in extinction in both patches.

3. Multi-patch source-sink model with intraspecific competition in the sink patches In this section, we consider the model of n patches, with s source patches and n - s sink patches given by:

$$\frac{dx_i}{dt} = x_i(a_i - \alpha_i x_i) + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j - \gamma_{ji} x_i), \qquad i = 1, \dots, n,$$
(32)

where  $x_i$  represent population densities of in the patch *i*. Without loss of generality, the *s* first patches are assumed to be the source (i.e.  $a_i > 0$  for all i = 1, ..., s) and the other n - s patches, assumed to bet the sink (i.e.  $a_i < 0$  for all i = s + 1, ..., n). The parameter  $\alpha_i$  is positive for all *i* and represent the intraspecific competition for the *i*-th patch. The parameter *D* represents the dispersion rate of the population,  $\gamma_{ij} \ge 0$  denote the flux between patches *j* and *i* for  $i \neq j$ . If  $\gamma_{ij} = 0$  then there is non direct flux from *j* to *i* and if  $\gamma_{ij} > 0$  there is a flux of population from patch *j* to patch *i*. We assume that, there exists intraspecific competition in n - s sink patches, i.e.  $\alpha_i > 0$  for all i = s + 1, ..., n. If we denote:

$$a_i = \begin{cases} r_i & \text{if } i = 1, \dots, s, \\ -r_i & \text{if } i = s+1, \dots, n \end{cases}$$

where  $r_i > 0$  for all *i*, and  $K_i = r_i/\alpha_i$  for all i = 1, ..., n, then System (32) can be written as:

$$\begin{cases} \frac{dx_i}{dt} = r_i x_i \left( 1 - \frac{x_i}{K_i} \right) + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j - \gamma_{ji} x_i), & i = 1, \dots, s, \\ \frac{dx_i}{dt} = r_i x_i \left( -1 - \frac{x_i}{K_i} \right) + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j - \gamma_{ji} x_i), & i = s+1, \dots, n, \end{cases}$$
(33)

where the parameters  $r_i > 0$  is the both growth rate in the case of source (i.e  $i \leq s$ ) patches and the death rate in the case of sink (i.e.  $s + 1 \leq i \leq n$ ),  $K_i > 0$  is the carrying capacity of source patches. System (33) can be written:

$$\begin{cases} \frac{dx_i}{dt} = r_i x_i \left(1 - \frac{x_i}{K_i}\right) + D \sum_{j=1}^n \gamma_{ij} x_j, & i = 1, \dots, s, \\ \frac{dx_i}{dt} = r_i x_i \left(-1 - \frac{x_i}{K_i}\right) + D \sum_{j=1}^n \gamma_{ij} x_j, & i = s+1, \dots, n, \end{cases}$$
(34)

where the term  $\gamma_{ii}$  accounts for the flux out of patch *i* and takes the form (3). We denote by  $\Gamma$  the matrix  $\Gamma := (\gamma_{ij})_{n \times n}$ . We call  $\Gamma$  the movement matrix of System (33). Its columns sum to 0 since the diagonal elements  $\gamma_{ii}$  are defined by (3) in such a way that each row sums to 0 and  $\Gamma$  is cooperative matrix (see Definition B.3). If  $\Gamma$  is irreducible, then 0 is a simple eigenvalue of  $\Gamma$ (see [8] or Lemma 4.1 and Lemma 1 of Elbetch et al. [14, 15]), and all nonzero eigenvalues of  $\Gamma$  have negative real parts, i.e., the stability modulus of a matrix  $\Gamma$  equals to zero. Moreover, the kernel of the matrix  $\Gamma$  is generated by a positive vector (see Lemma 2 in [4]). In all of this paper, we denote by  $\delta := (\delta_1, \ldots, \delta_n)^T$  this positive vector. Note that, if the matrix  $\Gamma$  is symmetric, then ker  $\Gamma$  is generated by  $\delta = (1, \ldots, 1)^T$ . The matrix

$$\Gamma_0 := \Gamma - \operatorname{diag}(\gamma_{11}, \dots, \gamma_{nn}) \tag{35}$$

which is the same as the matrix  $\Gamma$ , except that the diagonal elements are 0, is called the connectivity matrix. It is the adjacency matrix of the weighted directed graph  $\mathcal{G}$ , which has exactly *n* vertices (the patches), and there is an arrow from patch *j* to patch *i* precisely when  $\gamma_{ij} > 0$ , with weight  $\gamma_{ij}$  assigned to the arrow.

REMARK 3.1 For the existence, uniqueness, and positivity of  $\delta$  see Lemma 1 of Cosner et al. [8], Lemma 4.1 and Lemma 1 of Elbetch et al. [14, 15]. On the other hand, it is shown in Guo et al. [23, Lemma 2.1] and Gao and Dong [21, Lemma 3.1] that the vector  $(\Gamma_{11}^*, \ldots, \Gamma_{nn}^*)^T$  is a right eigenvector of  $\Gamma$  associated with the zero eigenvalue, where,  $\Gamma_{ii}^*$  is the cofactor of the *i*-th diagonal entry of  $\Gamma$ , and  $sgn(\Gamma_{ii}^*) = (-1)^{n-1}$ . For two patches we have  $\delta = (\gamma_{12}, \gamma_{21})^T$ , and for three patches we have  $\delta = (\delta_1, \delta_2, \delta_3)^T$ , where

$$\begin{cases} \delta_1 = \gamma_{12}\gamma_{13} + \gamma_{12}\gamma_{23} + \gamma_{32}\gamma_{13}, \\ \delta_2 = \gamma_{21}\gamma_{13} + \gamma_{21}\gamma_{23} + \gamma_{31}\gamma_{23}, \\ \delta_3 = \gamma_{21}\gamma_{32} + \gamma_{31}\gamma_{12} + \gamma_{31}\gamma_{32}. \end{cases}$$
(36)

In Lemma 2.1 Guo et al. [23] gives explicit formulas of the components of the vector  $\delta$ , with respect of the coefficients of  $\Gamma$  as follow:

$$\delta_k = \sum_{T \in \mathcal{T}_k} \prod_{(i,j) \in E(T)} \gamma_{ij}, \qquad k = 1, \dots, n,$$
(37)

where  $\mathcal{T}_k$  is the set of all directed trees of *n* vertices rooted at the *k*-th vertex, and E(T) denotes the set of arcs in a directed tree *T*.

System (33) can be also rewritten in matrix form as follow:

$$\begin{cases} \dot{X}_{s} = \operatorname{diag}(r_{1} - \frac{r_{1}}{K_{1}}x_{1}, \cdots, r_{s} - \frac{r_{s}}{K_{s}}x_{s})X_{s} + D(\Gamma_{ss}X_{s} + \Gamma_{sp}X_{p}), \\ \dot{X}_{p} = \operatorname{diag}(-r_{s+1} - \frac{r_{s+1}}{K_{s+1}}x_{s+1}, \cdots, -r_{n} - \frac{r_{n}}{K_{n}}x_{n})X_{p} \\ + D(\Gamma_{ps}X_{s} + \Gamma_{pp}X_{p}), \end{cases}$$
(38)

where  $X_s = (x_1, \ldots, x_s)^T$ ,  $X_p = (x_{s+1}, \ldots, x_p)^T$  and the matrices  $\Gamma_{ss}$ ,  $\Gamma_{pp}$ ,  $\Gamma_{sp}$ ,  $\Gamma_{ps}$ describe the flux within and between source and sink patches. They are obtained by writing the matrix  $\Gamma$  in block form as

$$\Gamma = \begin{bmatrix} \Gamma_{ss} & \Gamma_{sp} \\ \Gamma_{ps} & \Gamma_{pp} \end{bmatrix}.$$
(39)

Model (33) is studied in [39] for two patches, i.e n = 2 and s = 1.

**3.1. Global dynamics** In this part, our goal is to study the dynamics of System (33). Note that, in the absence of migration, i.e. the case where D = 0, System (33) admits  $(K_1, \ldots, K_s, 0, \ldots, 0)$  as a non trivial equilibrium point, which furthermore is GAS, and the origin as trivial equilibrium which is unstable. The problem is whether or not, the equilibrium continues to exist and to be GAS for any D > 0. The Jacobian matrix of System (33) evaluated at x = 0 is given by:

$$\mathbb{J}_s(0) = \operatorname{diag}(r_1, \dots, r_s, -r_{s+1}, \dots, -r_n) + D\Gamma,$$

$$\tag{40}$$

which is the same as the matrix obtained by Arino et al. [4, Equation 7] for System (2). The matrix  $\mathbb{J}_s(0)$  is cooperative. We have the following result which was proven also by Arino et al. [4]:

LEMMA 3.1 Consider the matrix  $\mathbb{J}_s(0)$ . Then, if s = 0,  $S(\mathbb{J}_0(0)) < 0$ , and if s = n,  $S(\mathbb{J}_n(0)) > 0$ , where S denote the stability modulus defined by (111).

PROOF If s = 0, then the matrix  $\mathbb{J}_0(0)$  becomes

$$\mathbb{J}_0(0) = \operatorname{diag}(-r_1, \dots, -r_n) + D\Gamma.$$
(41)

Let  $u = (1, \cdots, 1)^T$ . We have

$$\mathbb{J}_0(0)^T u = (-r_1, \cdots, -r_n)^T \le \lambda u, \quad \text{where} \quad \lambda = \max\{-r_1, \cdots, -r_n\} < 0.$$

Therefore, since  $\mathbb{J}_0(0)$  is a cooperative matrix, according to Lemma B.6, we have

$$S(\mathbb{J}_0(0)) = S(\mathbb{J}_0(0)^T) \le \lambda < 0.$$

If s = n, then the matrix  $\mathbb{J}_n(0)$  becomes

$$\mathbb{J}_n(0) = \operatorname{diag}(r_1, \dots, r_n) + D\Gamma.$$
(42)

Let  $u = (1, \cdots, 1)^T$ . We have

$$\mathbb{J}_n(0)^T u = (r_1, \cdots, r_n)^T \ge \lambda u, \quad \text{where} \quad \lambda = \min\{r_1, \cdots, r_n\} > 0.$$

Therefore, since  $\mathbb{J}_n(0)$  is a cooperative matrix, according to Lemma B.6, we have

$$S(\mathbb{J}_n(0)) = S(\mathbb{J}_n(0)^T) \ge \lambda > 0.$$

This completes the proof of the lemma.

We have also the following result:

LEMMA 3.2 The stability modulus of the matrix  $\mathbb{J}_s(0)$  is a non-decreasing function of s. Moreover, if the matrix of movement  $\Gamma$  is irreducible, then  $\mathbb{J}_s(0)$  is an increasing function of s.

**PROOF** See proof of Proposition 6 in [4].

The dynamics of System (33) in the case where  $\Gamma$  is reducible, is given as follows:

THEOREM 3.3 Consider System (33). Assume that  $\Gamma$  is reducible. Then, there exists a unique interval  $\mathcal{I} \subset ]0, n[\subset \mathbb{R}, such that:$ 

- If  $s < \min \mathcal{I}$ , then the origin is locally asymptotically stable (LAS), and
- if  $s > \max \mathcal{I}$ , then the origin is unstable.

PROOF We have  $S(\mathbb{J}_0(0)) < 0$  and  $S(\mathbb{J}_n(0)) > 0$  by Lemma 3.1. Moreover, the function  $s \mapsto S(\mathbb{J}_s(0))$  is non-decreasing by Lemma 3.2 and continues with respect to s (see [27, Theorem 2.4.9.2]). So by the intermediate value theorem, there exists an interval  $\mathcal{I}$ , possibly reduced to a single point, such that  $S(\mathbb{J}_s(0)) = 0$  for all  $s \in \mathcal{I}$ . Criteria for local asymptotic stability and instability of equilibria completes proof of the theorem.

Our goal in the remainder of this section is to study the dynamics of System (33) in the case when the matrix  $\Gamma$  is irreducible. First, it is clear that the solutions of (33) exist for all  $t \geq 0$  and remain non negative for non negative initial conditions. Thus, the positive cone  $\mathbb{R}^n_+$  is invariant under the flow of System (33). To establish the boundedness of solutions, we have the following result:

PROPOSITION 3.4 For any non negative initial condition, the solutions of System (33) remain non negative and bounded. Moreover, the set

$$\Lambda = \left\{ (x_1, \dots, x_n) \in \mathbb{R}^n : 0 \le \sum_{i=1}^n x_i \le \frac{\xi_2^*}{\xi_1^*} \right\}$$
(43)

is positively invariant and is a global attractor for (33), where  $\xi_1^* = \min_{1 \le i \le n} r_i$ and  $\xi_2^* = \sum_{s=1}^s r_i K_i$ .

PROOF To show that all solutions are bounded, we consider the quantity defined by  $X_T(t) = \sum_{i=1}^n x_i(t)$ . So, we have

$$\dot{X}_{T}(t) = \sum_{i=1}^{s} r_{i} x_{i}(t) \left( 1 - \frac{x_{i}(t)}{K_{i}} \right) + \sum_{i=s+1}^{n} r_{i} x_{i}(t) \left( -1 - \frac{x_{i}(t)}{K_{i}} \right).$$
(44)

For all  $r_i, K_i \in \mathbb{R}^*_+$ , we have the following inequality:

$$r_i x_i \left(1 - \frac{x_i}{K_i}\right) \le r_i (K_i - x_i), \quad i = 1, \dots, s,$$
  
$$r_i x_i \left(-1 - \frac{x_i}{K_i}\right) \le -r_i x_i, \quad i = s + 1, \dots, n.$$
(45)

Substituting (45) into (44), we get

$$\dot{X}_T(t) \le -\xi_1^* X_T(t) + \xi_2^* \qquad \forall t \ge 0,$$

which gives

$$X_T(t) \le \left(X_T(0) - \frac{\xi_2^*}{\xi_1^*}\right) e^{-\xi_1^* t} + \frac{\xi_2^*}{\xi_1^*}, \quad \text{for all } t \ge 0.$$
(46)

Hence,

$$X_T(t) \le \max\left(X_T(0), \frac{\xi_2^*}{\xi_1^*}\right), \quad \text{for all } t \ge 0$$

Therefore, the solutions of System (33) are positively bounded and defined for all  $t \ge 0$ . From (46) it can be deduced that the set  $\Lambda$  is positively invariant and it is a global attractor for System (33).

We have the result:

THEOREM 3.5 Consider System (33). Assume that the matrix  $\Gamma$  (or equivalently, the connectivity matrix  $\Gamma_0$ ) is irreducible, then, there exists a unique point  $\mathcal{I}^* \in ]0, n[$ , such that:

- If  $s < \mathcal{I}^*$ , then the origin is GAS, and
- if s > I<sup>\*</sup>, then the model has a unique equilibrium point E<sup>\*</sup>(D), which is GAS in the interior of the positive cone ℝ<sup>n</sup> \ {0}.

PROOF If the matrix  $\Gamma$  is irreducible, then the interval  $\mathcal{I}$  is reduced to a single point  $\mathcal{I}^*$ , such that: if  $s < \mathcal{I}^*$ , then  $S(\mathbb{J}_s(0)) < 0$ , and if  $s > \mathcal{I}^*$ , then  $S(\mathbb{J}_s(0)) > 0$ . According to [33, Theorem 1], if  $S(\mathbb{J}_s(0)) < 0$ , the origin is GAS. If  $S(\mathbb{J}_s(0)) > 0$ , then, System (33) is persistent for any D > 0, that is, any solution x(t) satisfies  $\liminf_{t\to\infty} x_i(t) > 0$ , for all i, and furthermore, since all the solutions to (33) are bounded, there exists a positive equilibrium point. We denote by  $(x_1^*(D), \ldots, x_n^*(D))$  an equilibrium of (33). Now, define the map:

$$\Upsilon_i: ]0, +\infty[ \to \mathbb{R}, \qquad \Upsilon_i(\xi) = \Psi_i(\xi E^*(D)),$$

where  $\Psi = (\Psi_1, \ldots, \Psi_n)$  denotes the vector field associated to (33). We have,

$$\Upsilon_i(\xi) = \frac{r_i}{K_i} \, (x_i^*(D))^2 \, \xi(1-\xi), \qquad i = 1, \dots, n.$$

Therefore, according to Theorem B.9, we conclude the proof of theorem.

As a corollary of the previous theorem we obtain the following result which proven in [14, Theorem 6.1]:

COROLLARY 3.6 If s = n, System (33) has a unique equilibrium point in the interior of the positive cone, which is GAS.



Figure 3: The assumption that the matrix  $\Gamma$  is irreducible, implies that the species can reach any i-th patch from any j-patch. For one-source, one-sink patch model, the matrix  $\Gamma$  is irreducible if and only if  $\gamma_{12}$  and  $\gamma_{21}$  are positives. For three-patch model, under the irreducibility hypothesis on the matrix  $\Gamma$ , there are five possible cases, modulo permutation of the three patches. The two graphs  $\mathcal{G}_1$  and  $\mathcal{G}_2$  for which the migration matrix may be symmetric, if  $\gamma_{ij} = \gamma_{ji}$ . For the remaining cases, the graphs  $\mathcal{G}_3, \mathcal{G}_4$  and  $\mathcal{G}_5$ , cannot be symmetrical.

In all of this work, we denote  $E^*(D)$  the unique equilibrium in the interior of the positive cone of System (33) if it exists, and  $X_T^*(D)$ , the total equilibrium population:

$$X_T^*(D) = \sum_{i=1}^n x_i^*(D), \qquad E^*(D) = (x_1^*(D), \dots, x_n^*(D)).$$
(47)

## 3.2. The behavior of the model for large migration rate

In this section, our aim is to study the behavior of System (33) for large migration rate, i.e. when  $D \to \infty$ . We use the theory of singular perturbations and Tikhonov's theorem [32, 37, 38] to obtain a better understanding of the

behavior of the system in the case of perfect mixing. We have the following result:

THEOREM 3.7 Let  $(x_1(t, D), \ldots, x_n(t, D))$  be the solution of System (33) with initial condition  $(x_1^0, \cdots, x_n^0)$  satisfying  $x_i^0 \ge 0$  for  $i = 1 \cdots n$ . Let Y(t) be the solution of the equation

$$\frac{dX}{dt} = rX\left(1 - \frac{X}{\left(\sum_{i=1}^{n} \delta_i\right)K}\right),\tag{48}$$

where  $(\delta_1, ..., \delta_n)^T$  the vector which generate the vector space ker  $\Gamma$ ,  $\alpha_i = r_i/K_i$ and

$$r = \frac{\sum_{i=1}^{s} \delta_{i} r_{i} - \sum_{i=s+1}^{n} \delta_{i} r_{i}}{\sum_{i=1}^{n} \delta_{i}}, \quad K = \frac{\sum_{i=1}^{s} \delta_{i} r_{i} - \sum_{i=s+1}^{n} \delta_{i} r_{i}}{\sum_{i=1}^{n} \delta_{i}^{2} \alpha_{i}}.$$
 (49)

Then, when  $D \to \infty$ , we have

$$\sum_{i=1}^{n} x_i(t, D) = Y(t) + o_D(1) \qquad \text{uniformly for } t \in [0, +\infty)$$
(50)

and, for any  $t_0 > 0$ , we have

$$x_i(t,D) = \frac{\delta_i}{\sum_{i=1}^n \delta_i} Y(t) + o_D(1) \qquad i = 1, \dots, n, \text{ uniformly for } t \in [t_0, +\infty).$$
(51)

PROOF Let  $X(t, D) = \sum_{i=1}^{n} x_i(t, D)$ . We rewrite System (33) using the variables  $(X, x_1, \dots, x_{n-1})$ , and get:

$$\begin{cases} \frac{dX}{dt} = \sum_{i=1}^{s} r_i x_i \left(1 - \frac{x_i}{K_i}\right) - \sum_{i=s+1}^{n} r_i x_i \left(1 + \frac{x_i}{K_i}\right), \\ \frac{dx_i}{dt} = r_i x_i \left(1 - \frac{x_i}{K_i}\right) + D \sum_{j=1, j \neq i}^{n} (\gamma_{ij} x_j - \gamma_{ji} x_i), \quad i = 1, \dots, s, \\ \frac{dx_i}{dt} = r_i x_i \left(-1 - \frac{x_i}{K_i}\right) + D \sum_{j=1, j \neq i}^{n} (\gamma_{ij} x_j - \gamma_{ji} x_i), \quad i = s+1, \dots, n-1 \end{cases}$$

$$(52)$$

This system is actually a system in the variables  $(X, x_1, \dots, x_{n-1})$ , since, whenever  $x_n$  appears in the right hand side of (52), it should be replaced by

$$x_n = X - \sum_{i=1}^{n-1} x_i.$$
 (53)

When  $D \to \infty$ , (52) is a *slow-fast* system, with one *slow variable*, X, and n-1 fast variables,  $x_i$  for  $i = 1 \dots n-1$ . As suggested by Tikhonov's theorem

[32, 37, 38], we consider the dynamics of the fast variables in the time scale  $\tau = Dt$ . We get

$$\begin{cases} \frac{dx_i}{d\tau} = \frac{1}{D} r_i x_i \left( 1 - \frac{x_i}{K_i} \right) + \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j - \gamma_{ji} x_i), & i = 1, \dots, s, \\ \frac{dx_i}{d\tau} = \frac{1}{D} r_i x_i \left( -1 - \frac{x_i}{K_i} \right) + \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j - \gamma_{ji} x_i), & i = s+1, \dots, n-1. \end{cases}$$

$$\tag{54}$$

where  $x_n$  is given by (53). In the limit  $D \to \infty$ , we find the *fast dynamics* 

$$\frac{dx_i}{d\tau} = \sum_{j=1, j\neq i}^n (\gamma_{ij}x_j - \gamma_{ji}x_i), \qquad i = 1, \cdots, n-1.$$

This is an (n-1)-dimensional linear differential system. According to [15, Lemma B.1], this system admits unique LAS equilibrium given by

$$\left(\frac{\delta_1}{\sum_{i=1}^n \delta_i} X, \dots, \frac{\delta_{n-1}}{\sum_{i=1}^n \delta_i} X\right)^T.$$

Thus, the slow manifold of System (52) is given by

$$x_i = \frac{\delta_i}{\sum_{i=1}^n \delta_i} X, \qquad i = 1, \dots, n-1.$$
 (55)

As this manifold is LAS, Tikhonov's theorem ensures that after a fast transition toward the slow manifold, the solutions of (52) are approximated by the solutions of the *reduced model*, which is obtained by replacing (55) into the dynamics of the slow variable, that is:

$$\begin{aligned} \frac{dX}{dt} &= \sum_{i=1}^{s} r_i \frac{X}{\sum_{i=1}^{n} \delta_i} \delta_i \left( 1 - \frac{X}{\left(\sum_{i=1}^{n} \delta_i\right) K_i} \delta_i \right) + \sum_{i=s+1}^{n} r_i \frac{X}{\sum_{i=1}^{n} \delta_i} \delta_i \left( -1 - \frac{X}{\left(\sum_{i=1}^{n} \delta_i\right) K_i} \delta_i \right) \\ &= rX \left( 1 - \frac{X}{\left(\sum_{i=1}^{n} \delta_i\right) K} \right), \end{aligned}$$

where r and K are defined in (49). Therefore, the reduced model is (48). If  $\sum_{i=1}^{s} \delta_i r_i > \sum_{i=s+1}^{n} \delta_i r_i$ , (48) admits

$$X^* = \left(\sum_{i=1}^n \delta_i\right) K = \left(\sum_{i=1}^n \delta_i\right) \frac{\sum_{i=1}^s \delta_i r_i - \sum_{i=s+1}^n \delta_i r_i}{\sum_{i=1}^n \delta_i^2 \alpha_i}$$

as a positive equilibrium point, which is LAS in the positive axis, and if  $\sum_{i=1}^{s} \delta_i r_i \leq \sum_{i=s+1}^{n} \delta_i r_i$ , (48) admits the origin as unique equilibrium point, which is LAS. The approximation given by Tikhonov's theorem holds for all  $t \geq 0$  for the slow variable and for all  $t \geq t_0 > 0$  for the fast variables, where  $t_0$  is as small as we want. Therefore, letting Y(t) be the solution of the reduced model (48) with initial condition  $Y(0) = X(0, D) = \sum_{i=1}^{n} x_i^0$ , then, then  $D \to \infty$ , we have Approximations (50) and (51).

Note that, in the case of perfect mixing, Approximation (50) shows that:

• If  $\sum_{i=1}^{s} \delta_i r_i > \sum_{i=s+1}^{n} \delta_i r_i$ , then the total population behaves like the unique logistic equation (48) and then, when t and D tend to  $\infty$ , the total population  $\sum x_i(t, D)$  tends toward

$$\left(\sum_{i=1}^{n} \delta_i\right) K = \left(\sum_{i=1}^{n} \delta_i\right) \frac{\sum_{i=1}^{s} \delta_i r_i - \sum_{i=s+1}^{n} \delta_i r_i}{\sum \delta_i^2 \alpha_i}$$

• If  $\sum_{i=1}^{s} \delta_i r_i \leq \sum_{i=s+1}^{n} \delta_i r_i$ , then Equation (48) has the origin as unique equilibrium, and then, when t and D tend to  $\infty$ , the total population  $\sum x_i(t, D)$  tends toward 0.

Approximation (51) shows that, with the exception of a thin initial boundary layer, where the density population  $x_i(t, D)$  quickly jumps from its initial condition  $x_i^0$  to the average  $\delta_i X_0 / \sum_{i=1}^n \delta_i$ , each patch of the *n*-patch source-sink model behaves like the following equation:

$$\frac{du}{dt} = \begin{cases} ru\left(1 - \frac{u}{\delta_i K}\right) & \text{if} \quad \sum_{i=1}^s \delta_i r_i > \sum_{i=s+1}^n \delta_i r_i, \\ \tilde{r}u\left(-1 - \frac{u}{\delta_i \tilde{K}}\right) & \text{otherwise,} \end{cases}$$

where  $\tilde{r} = -r, \tilde{K} = -K, r$  and K are given in (49).

Hence, when t and D tend to  $\infty$ , the density population  $x_i(t, D)$  tends toward  $K = \delta_i \frac{\sum_{i=1}^s \delta_i r_i - \sum_{i=s+1}^n \delta_i r_i}{\sum_{i=1}^{\delta_i^2} \alpha_i}$  if  $\sum_{i=1}^s \delta_i r_i > \sum_{i=s+1}^n \delta_i r_i$ , and  $x_i(t, D)$ tends toward 0 if  $\sum_{i=1}^s \delta_i r_i \le \sum_{i=s+1}^n \delta_i r_i$ .

According to the previous theorem, we obtain the limit  $E^*(\infty)$  of  $E^*(D)$ when  $D \to \infty$ :

COROLLARY 3.8 We have:

$$\lim_{D \to +\infty} E^*(D) = \begin{cases} \frac{\sum_{i=1}^s \delta_i r_i - \sum_{i=s+1}^n \delta_i r_i}{\sum_{i=1}^n \delta_i^2 \alpha_i} (\delta_1, \dots, \delta_n), & \text{if } \sum_{i=1}^s \delta_i r_i > \sum_{i=s+1}^n \delta_i r_i, \\ 0 & \text{otherwise,} \end{cases}$$
(56)

where  $\alpha_i = r_i/K_i$  and  $(\delta_1, \ldots, \delta_n)$  the vector which generate the kernel of  $\Gamma$ . Moreover, if the matrix  $\Gamma$  is symmetric, then:

$$\lim_{D \to +\infty} E^*(D) = \begin{cases} \frac{\sum_{i=1}^{s} r_i - \sum_{i=s+1}^{n} r_i}{\sum_{i=1}^{n} \alpha_i} (1, \dots, 1), & \text{if } \sum_{i=1}^{s} r_i > \sum_{i=s+1}^{n} r_i, \\ 0 & \text{otherwise.} \end{cases}$$
(57)

As a second corollary of the previous theorem we obtain the following result which describes the total equilibrium population for perfect mixing: COROLLARY 3.9 We have

$$X_T^*(+\infty) = \begin{cases} \sum_{i=1}^n \delta_i \frac{\sum_{i=1}^s \delta_i r_i - \sum_{i=s+1}^n \delta_i r_i}{\sum_{i=1}^n \delta_i^2 \alpha_i} & \text{if } \sum_{i=1}^s \delta_i r_i > \sum_{i=s+1}^n \delta_i r_i, \\ 0, & \text{otherwise.} \end{cases}$$
(58)

Moreover, if the matrix  $\Gamma$  is symmetric, then:

$$X_T^*(+\infty) = \begin{cases} n \frac{\sum_{i=1}^{s} r_i - \sum_{i=s+1}^{n} r_i}{\sum_{i=1}^{n} \alpha_i}, & \text{if } \sum_{i=1}^{s} r_i > \sum_{i=s+1}^{n} r_i, \\ 0, & \text{otherwise.} \end{cases}$$
(59)

PROOF The sum of the *n* components of the point  $E^*(\infty)$  immediately gives Formula (58).

In the case n = 2 and s = 1, one has  $\delta_1 = \gamma_{12}$  and  $\delta_2 = \gamma_{21}$ . Therefore (58) becomes

$$X_T^*(+\infty) = \begin{cases} (\gamma_{12} + \gamma_{21}) \frac{\gamma_{12}r_1 - \gamma_{21}r_2}{\gamma_{12}^2 \alpha_1 + \gamma_{21}^2 \alpha_2} & if & \gamma_{21}/\gamma_{12} < r_1/r_2, \\ 0 & otherwise. \end{cases}$$

which is the formula given by Wu et al. [39, Equation 5.8]. In the case n = s = 2, Formula (58) becomes

$$X_T^*(+\infty) = (\gamma_{12} + \gamma_{21}) \frac{\gamma_{12}r_1 + \gamma_{21}r_2}{\gamma_{12}^2\alpha_1 + \gamma_{21}^2\alpha_2},$$

which is the formula given by Arditi et al. [2, Equation 7] and by Poggiale et al. [35, page 362].

In the case of the multi-patch logistic model with asymmetric migration, i.e. the model (33) with s = n, Formula (58) becomes

$$X_T^*(+\infty) = \left(\sum_{i=1}^n \delta_i\right) \frac{\sum_{i=1}^n \delta_i r_i}{\sum_{i=1}^n \delta_i^2 \alpha_i},$$

which is the formula given by Elbetch et al. [15, Equation 13].

As a corollary of the previous theorem we obtain the following result which describes the behavior of System (33) for perfect mixing and symmetrical dispersal:

COROLLARY 3.10 Assume that the matrix  $\Gamma$  is symmetric. Let  $(x_1(t, D), \ldots, x_n(t, D))$ be the solution of System (33) with initial condition  $(x_1^0, \cdots, x_n^0)$  satisfying  $x_i^0 \ge 0$  for  $i = 1 \cdots n$ . Let Y(t) be the solution of the equation

$$\frac{dX}{dt} = rX\left(1 - \frac{X}{nK}\right),\tag{60}$$

where

$$r = \frac{\sum_{i=1}^{s} r_i - \sum_{i=s+1}^{n} r_i}{n}, \quad K = \frac{\sum_{i=1}^{s} r_i - \sum_{i=s+1}^{n} r_i}{\sum_{i=1}^{n} \alpha_i} \quad and \quad \alpha_i = r_i / K_i.$$

Then, when  $D \to \infty$ , we have

$$\sum_{i=1}^{n} x_i(t, D) = Y(t) + o_D(1) \qquad \text{uniformly for } t \in [0, +\infty)$$

and, for any  $t_0 > 0$ , we have

$$x_i(t,D) = \frac{Y(t)}{n} + o_D(1) \qquad i = 1, \dots, n, \text{ uniformly for } t \in [t_0, +\infty).$$

PROOF If  $\Gamma$  is symmetric, one has  $\delta_i = 1$  for all *i*. Therefore, Formulas (48), (49), and Approximations (50), (51) for  $\delta_i = 1$ , give the proof of the corollary.

**3.3. Total population size** In this section, Our aim is to compare the total equilibrium population with the sum of carrying capacities  $K_1 + \ldots + K_s$ , when the migration rate D varies from zero to infinity. First, we start with the following case.

**3.3.1. Homogeneous source-sink system** Let we consider a source-sink patch model (33) when the patches have a homogeneous structure in the sense that the growth rates and death rates are independent of the patch. In the next proposition, we show that, if the growth rates in source patches, and the death rates in the sink patches, are equal, then the total equilibrium population is smaller than the sum of carrying capacities. Moreover, under some conditions, we can have a persistence or extinction in all the patches. Mathematically speaking, we have the result:

PROPOSITION 3.11 Consider System (33). If  $r_1 = \ldots = r_n$ , then  $X_T^*(D) \leq \sum_{i=1}^s K_i$  for all  $D \geq 0$ , and  $\frac{dX_T^*}{dD}(0) = 0$ . Moreover,

- If  $\sum_{i=1}^{s} \delta_i r_i \leq \sum_{i=s+1}^{n} \delta_i r_i$ , then there is  $D^* > 0$  such that,  $X_T^*(D) > 0$  for  $D < D^*$ , and  $X_T^*(D) = 0$ , for  $D \geq D^*$ .
- If  $\sum_{i=1}^{s} \delta_i r_i > \sum_{i=s+1}^{n} \delta_i r_i$ , then  $X_T^*(D) > 0$  for all  $D \ge 0$ .

PROOF If the equilibrium  $E^*(D)$  exist, then it is a solution of the algebraic system:

$$\begin{cases} 0 = r_i x_i \left( 1 - \frac{x_i}{K_i} \right) + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j - \gamma_{ji} x_i), & i = 1, \dots, s, \\ 0 = r_i x_i \left( -1 - \frac{x_i}{K_i} \right) + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j - \gamma_{ji} x_i), & i = s+1, \dots, n. \end{cases}$$

The sum of these equations shows that  $E^*(D)$  satisfies the following equation

$$\sum_{i=1}^{s} r_i x_i \left( 1 - \frac{x_i}{K_i} \right) - \sum_{i=s+1}^{n} r_i x_i \left( 1 + \frac{x_i}{K_i} \right) = 0.$$

Therefore  $E^*(D)$  belongs to the ellipsoid:

$$\mathbb{E}_{s}^{n-1} := \left\{ x \in \mathbb{R}^{n} : \Theta(x) := \sum_{i=1}^{s} r_{i} x_{i} \left( 1 - \frac{x_{i}}{K_{i}} \right) - \sum_{i=s+1}^{n} r_{i} x_{i} \left( 1 + \frac{x_{i}}{K_{i}} \right) = 0 \right\}.$$
(61)

Note that, this ellipsoid is independent of the migration terms D and  $\gamma_{ij}$ . It depends on the number of the sources and sinks. The ellipsoid  $\mathbb{E}_s^{n-1}$  passes through the points O,  $(K_1, \ldots, K_s, 0, \ldots, 0)$  and  $(0, \ldots, 0, -K_{s+1}, \ldots, -K_n)$ .

The equation of the tangent space to the ellipsoid  $\mathbb{E}_s^{n-1}$ , defined by (61), at point  $\mathcal{A}_s = (K_1, \ldots, K_s, 0, \ldots, 0)$  is given by

$$\sum_{i=1}^{s} (x_i - K_i) \frac{\partial \Theta}{\partial x_i} (\mathcal{A}_s) + \sum_{i=s+1}^{n} x_i \frac{\partial \Theta}{\partial x_i} (\mathcal{A}_s) = 0,$$
(62)

where  $\Theta$  is given by Equation (61). Since  $\frac{\partial \Theta}{\partial x_i}(\mathcal{A}_s) = -r_i$  for all  $i = 1, \ldots, n$ , Equation (62) can be written as follows:

$$\sum_{i=1}^{n} r_i x_i = \sum_{i=1}^{s} r_i K_i.$$
(63)

If we take  $r_1 = \ldots = r_n$ , in Equation (63), we get that the equation of the tangent plane to  $\mathbb{E}_s^{n-1}$  at the point  $\mathcal{A}_s$  is

$$\sum_{i=1}^n x_i = \sum_{i=1}^s K_i.$$

By the convexity of ellipsoid  $\mathbb{E}_s^{n-1}$ , any point of  $\mathbb{E}_s^{n-1}$  lies in the half-space defined by the inequality  $\sum_{i=1}^n x_i \leq \sum_{i=1}^s K_i$ . Therefore  $E^*(D)$  satisfies

$$\sum_{i=1}^{n} x_i^*(D) \le \sum_{i=1}^{s} K_i \quad \text{for all} \quad D \ge 0.$$

Now, according to the formula of perfect mixing (58), we can see immediately that  $X_T^*(+\infty) = 0$  if and only if  $\sum_{i=1}^s \delta_i r_i = \sum_{i=s+1}^n \delta_i r_i$ .

If  $r_1 = \ldots = r_n =: r$ , then the formula of the derivative (108) at D = 0 becomes

$$\frac{dX_T^*}{dD}(0) = \frac{1}{r} (1, \dots, 1) \Gamma (K_1, \dots, K_s, 0, \dots, 0)^T = 0.$$

This completes the proof of the proposition.

In the case when s = n, the previous proposition becomes: if  $r_1 = \ldots = r_s$ , then  $0 < X_T^*(D) \le \sum_i K_i$ , which is [14, Prop. 3.1 and Prop. 6.2].

**3.3.2.** Heterogeneous source-sink system In the next proposition we give sufficient and necessary conditions for the total equilibrium population not to depend on the migration rate. More precisely, we show that, the only situation where the total equilibrium population is independent with respect to dispersal, is when all the patches are sources and the vector of the carrying capacities lies in the vector space ker  $\Gamma$ . That is, if there is at least one sink patch, or we have n sources and the vector of the carrying capacities does not belong to the vector space ker  $\Gamma$ , then the total equilibrium population depends on the dispersion.

PROPOSITION 3.12 The equilibrium  $E^*(D)$  does not depend on D if and only if, s = n and  $(K_1, \ldots, K_n) \in \ker \Gamma$ . In this case  $E^*(D) = (K_1, \ldots, K_n)$  for all D > 0.

PROOF The equilibrium  $E^*(D) := (X_s^*(D); X_p^*(D))$ , where  $X_s^*(D) = (x_1^*(D), \ldots, x_s^*(D))$  and  $X_p^*(D) = (x_{s+1}^*(D), \ldots, x_n^*(D))$ , is the unique positive solution of the system (38), i.e:

$$\begin{cases}
0 = \operatorname{diag}\left(r_{1} - \frac{r_{1}}{K_{1}}x_{1}^{*}(D), \cdots, r_{s} - \frac{r_{s}}{K_{s}}x_{s}^{*}(D)\right) X_{s}^{*}(D) \\
+ D\left(\Gamma_{ss}X_{s}^{*}(D) + \Gamma_{sp}X_{p}^{*}(D)\right), \\
0 = \operatorname{diag}\left(-r_{s+1} - \frac{r_{s+1}}{K_{s+1}}x_{s+1}^{*}(D), \cdots, -r_{n} - \frac{r_{n}}{K_{n}}x_{n}^{*}(D)\right) X_{p}^{*}(D) \\
+ D\left(\Gamma_{ps}X_{s}^{*}(D) + \Gamma_{pp}X_{p}^{*}(D)\right).
\end{cases}$$
(64)

Suppose that the equilibrium  $E^*(D)$  does not depend on D. The derivative of (64) with respect to D gives:

$$\begin{cases} 0 = \Gamma_{ss}X_s^*(D) + \Gamma_{sp}X_p^*(D) \\ 0 = \Gamma_{ps}X_s^*(D) + \Gamma_{pp}X_p^*(D) \end{cases} \Leftrightarrow \begin{bmatrix} \Gamma_{ss} & \Gamma_{sp} \\ \Gamma_{ps} & \Gamma_{pp} \end{bmatrix} \begin{bmatrix} X_s^* \\ X_p^* \end{bmatrix} = 0 \Leftrightarrow \Gamma E^*(D) = 0.$$
(65)

Replacing Equation (65) in (64), we get  $E^*(D) = (K_1, \ldots, K_s, 0, \ldots, 0)$ . From Equation (65), we conclude that  $(K_1, \ldots, K_s, 0, \ldots, 0) \in \ker \Gamma$ . Since the vector space ker  $\Gamma$  is generated by a positive vector, then  $(K_1, \ldots, K_s, 0, \ldots, 0) \in \ker \Gamma$  is hold if and only if, s = n.

Now, suppose that s = n and  $(K_1, \ldots, K_n) \in \ker \Gamma$ , then  $(K_1, \ldots, K_n)$  satisfies Equation (64), for all  $D \ge 0$ . So,  $E^*(D) = (K_1, \ldots, K_n)$ , for all  $D \ge 0$ , which proves that the total equilibrium population is independent of the migration rate D.

It is also clear that when all the patches are sources and the vector of the carrying capacities lies in the vector space ker  $\Gamma$ , we obtain the results proved by Elbetch et al. [14, Prop. 3.2] and [15, Prop. 4.5].

**3.3.3. Two blocks of identical source and sink patches** We consider the model of source-sink patches (33) and we assume that we have one block of identical source patches and one block of identical sink patches. We denote by  $I_{so} = \{1, \ldots, s\}$  and  $I_{si} = \{s + 1, \ldots, n\}$  for the block of the source patches and sink patches respectively such that  $I_{so} \cup I_{si} = \{1, \ldots, n\}$ . The source patches being identical means that they have the same growth rate  $r_i$  and carrying capacity  $K_i$ . Therefore, we have

$$r_1 = \ldots = r_s =: r_{so}, \quad K_1 = \ldots = K_s =: K_{so}.$$
 (66)

The same for the sink patches, we suppose that:

$$r_{s+1} = \ldots = r_n =: r_{si}, \quad K_{s+1} = \ldots = K_n =: K_{si}.$$
 (67)

First, we give some definitions:

DEFINITION 3.13 Let the flux

$$\Gamma_{iI_{si}} = \sum_{j \in I_{si}} \gamma_{ij}, \Gamma_{jI_{so}} = \sum_{i \in I_{so}} \gamma_{ij}, \Gamma_{I_{so}I_{si}} = \sum_{i \in I_{so}, j \in I_{si}} \gamma_{ij}, \text{and} \quad \Gamma_{I_{si}I_{so}} = \sum_{i \in I_{so}, j \in I_{si}} \gamma_{ji}$$

- For  $i \in I_{so}$ ,  $\Gamma_{iI_{si}}$  is the flux from block  $I_{si}$  to patch *i*, i.e. the sum of the migration rates  $\gamma_{ij}$  from patches  $j \in I_{si}$  to patch *i*.
- For  $j \in I_{si}, \Gamma_{jI_{so}}$  is the flux from block  $I_{so}$  to patch j, i.e. the sum of the migration rates  $\gamma_{ji}$  from patches  $i \in I_{so}$  to patch j.
- $\Gamma_{I_{so}I_{si}}$  is the flux from block  $I_{si}$  to block  $I_{so}$ , i.e. the sum of the migration rates  $\gamma_{ij}$  from patches  $j \in I_{si}$ , to patches  $i \in I_{so}$ .
- $\Gamma_{I_{si}I_{so}}$  is the flux from block  $I_{so}$  to block  $I_{si}$ , i.e. the sum of the migration rates  $\gamma_{ji}$  from patches  $i \in I_{so}$ , to patches  $j \in I_{si}$ .

For each patch *i* we denote by  $\mathcal{T}_i$  the sum of all migration rates  $\gamma_{ji}$  from patch *i* to another patch  $j \neq i$  (i.e. the outgoing flux of patch i) minus the sum of the migration rates  $\gamma_{ik}$  from patch *k* to patch *i*, where *k* belongs to the same block as *i*. Hence, we have:

If 
$$i \in I_{so}$$
,  $\mathcal{T}_i = \sum_{j \in I_{si}} \gamma_{ji} + \sum_{k \in I_{so} \setminus \{i\}} (\gamma_{ki} - \gamma_{ik})$   
If  $j \in I_{si}$ ,  $\mathcal{T}_j = \sum_{i \in I_{so}} \gamma_{ij} + \sum_{k \in I_{si} \setminus \{j\}} (\gamma_{kj} - \gamma_{jk})$ 

We make the following assumption on the migration rates

$$\Gamma_{1I_{si}} = \dots = \Gamma_{sI_{si}}, \qquad \Gamma_{(s+1)I_{so}} = \dots = \Gamma_{nI_{so}}$$
  
$$\mathcal{T}_1 = \dots = \mathcal{T}_s, \qquad \qquad \mathcal{T}_{s+1} = \dots = \mathcal{T}_n$$
(68)

If Conditions (68) are satisfied, then, according to [15, Lemma 4.6], for all  $i \in I_{so}$  and  $j \in I_{si}$  one has

$$\Gamma_{iI_{si}} = \Gamma_{I_{so}I_{si}}/s, \quad \Gamma_{jI_{so}} = \Gamma_{I_{si}I_{so}}/\overline{s}, \quad \mathcal{T}_{i} = \Gamma_{I_{si}I_{so}}/s, \quad \mathcal{T}_{j} = \Gamma_{I_{so}I_{si}}/\overline{s}.$$
(69)

where  $\overline{s} = n - s$ ,  $\Gamma_{I_{so}I_{si}}$  and  $\Gamma_{I_{si}I_{so}}$  are defined in Definition 3.13.

We consider the following regions in the set of parameters  $\Gamma_{I_{si}I_{so}}$  and  $\Gamma_{I_{so}I_{si}}$ , denoted by  $\mathcal{Z}_0, \mathcal{Z}_1$  and  $\mathcal{Z}_2$  depicted in Figure 4 and defined by:

$$\begin{cases} \mathcal{Z}_{0} = \left\{ (\Gamma_{I_{si}I_{so}}, \Gamma_{I_{so}I_{si}}) : & \Gamma_{I_{so}I_{si}} \geq \frac{r_{si}}{r_{so}} \Gamma_{I_{si}I_{so}} \right\}, \\ \mathcal{Z}_{1} = \left\{ (\Gamma_{I_{si}I_{so}}, \Gamma_{I_{so}I_{si}}) : & \frac{r_{si}}{r_{so}} \Gamma_{I_{si}I_{so}} < \Gamma_{I_{so}I_{si}} < \frac{r_{si}D}{r_{so}r_{si} + Dr_{so}} \Gamma_{I_{si}I_{so}} \right\}, \\ \mathcal{Z}_{2} = \left\{ (\Gamma_{I_{si}I_{so}}, \Gamma_{I_{so}I_{si}}) : & \Gamma_{I_{so}I_{si}} \geq \frac{r_{si}D}{r_{so}r_{si} + Dr_{so}} \Gamma_{I_{si}I_{so}} \right\}. \end{cases}$$

$$(70)$$



Figure 4: The regions  $\mathcal{Z}_0, \mathcal{Z}_1$  and  $\mathcal{Z}_2$  in the set of parameters  $\Gamma_{I_{si}I_{so}}$  and  $\Gamma_{I_{so}I_{si}}$ .

We have the following result

THEOREM 3.14 Consider the regions  $Z_0, Z_1$  and  $Z_2$  depicted in Figure 4 and defined by (70). Assume that Conditions (66),(67) and (68) are satisfied. If  $(\Gamma_{I_{si}I_{so}}, \Gamma_{I_{so}I_{si}}) \in Z_2$ , then System (33) admits the origin as unique equilibrium point, which is GAS, and if  $(\Gamma_{I_{si}I_{so}}, \Gamma_{I_{so}I_{si}}) \in Z_0 \cup Z_1$ , System (33) admits unique equilibrium point in the interior of the positive cone, which is of the form

$$x_1 = x_1^*, \dots, x_s = x_{so}^*, \quad x_{s+1} = x_n^*, \dots, x_n = x_{si}^*$$

where  $(x_{so}^*, x_{si}^*)$  is the interior equilibrium point of the 2-patch source -sink model

$$\begin{cases} \frac{dx_{so}}{dt} = sr_{so}x_{so}\left(1 - \frac{x_{so}}{K_{so}}\right) + D\left(\Gamma_{I_{so}I_{si}}x_{si} - \Gamma_{I_{si}I_{so}}x_{so}\right), \\ \frac{dx_{si}}{dt} = \bar{s}r_{si}x_{si}\left(-1 - \frac{x_{si}}{K_{si}}\right) + D\left(\Gamma_{I_{si}I_{so}}x_{so} - \Gamma_{I_{so}I_{si}}x_{si}\right), \end{cases}$$
(71)

with specific growth rates  $sr_{so}$  and death rate  $\overline{s}r_{si}$ , carrying capacities  $K_{so}$  for the source patch, parameter  $K_{si}$  due to the intraspecific competition in the sink patch and migration rates  $\Gamma_{I_{si}I_{so}}$  from source patch to the sink patch and  $\Gamma_{I_{so}I_{si}}$  from the sink patch to the source patch.

PROOF Assume that Conditions (66) and (67) are satisfied. Then, if the interior equilibrium point of (33) exist, it is the unique positive solution of the set of algebraic equations

$$\begin{cases} r_{so}x_{i}\left(1-\frac{x_{i}}{K_{so}}\right)+D\sum_{k=1,k\neq i}^{n}(\gamma_{ik}x_{k}-\gamma_{ki}x_{i})=0, & i=1,\cdots,s, \\ r_{si}x_{j}\left(-1-\frac{x_{j}}{K_{si}}\right)+D\sum_{k=1,k\neq j}^{n}(\gamma_{jk}x_{k}-\gamma_{kj}x_{j})=0, & j=s+1,\cdots,n. \end{cases}$$
(72)

We consider the following set of algebraic equations obtained from (72) by replacing  $x_i = x_{so}$  for i = 1, ..., s and  $x_i = x_{si}$  for i = s + 1, ..., n:

$$\begin{cases} r_{so}x_{so}\left(1-\frac{x_{so}}{K_{so}}\right)+D\left(\Gamma_{iI_{si}}x_{si}-\mathcal{T}_{i}x_{so}\right)=0, & i=1,\cdots,s,\\ r_{si}x_{n}\left(-1-\frac{x_{si}}{K_{si}}\right)+D\left(\Gamma_{jI_{so}}x_{so}-\mathcal{T}_{j}x_{si}\right)=0, & j=s+1,\cdots,n. \end{cases}$$

$$\tag{73}$$

Now, using Condition (68), together with (69), we see that System (73) is equivalent to the set of two algebraic equations:

$$\begin{cases} r_{so}x_{so}\left(1-\frac{x_{so}}{K_{so}}\right) + D\left(\frac{\Gamma_{I_{so}I_{si}}}{s}x_{si} - \frac{\Gamma_{I_{si}I_{so}}}{s}x_{so}\right) = 0, \\ r_{si}x_{si}\left(-1-\frac{x_{si}}{K_{si}}\right) + D\left(\frac{\Gamma_{I_{si}I_{so}}}{\overline{s}}x_{so} - \frac{\Gamma_{I_{so}I_{si}}}{\overline{s}}x_{si}\right) = 0. \end{cases}$$
(74)

We first notice that if  $x_{so} = x_{so}^*$ ,  $x_{si} = x_{si}^*$  is a positive solution of (74) then  $x_i = x_{so}^*$  for  $i = 1, \ldots, s$  and  $x_i = x_{si}^*$  for  $i = s + 1, \ldots, n$  is a positive solution of (72). According to Proposition 2.1, if  $(\Gamma_{I_{si}I_{so}}, \Gamma_{I_{so}I_{si}}) \in \mathbb{Z}_2$ , then System (74) has the origin as unique equilibrium point, which is GAS, and if  $(\Gamma_{I_{si}I_{so}}, \Gamma_{I_{so}I_{si}}) \in \mathbb{Z}_0 \cup \mathbb{Z}_1$ , System (74) has unique equilibrium point in the interior of the positive cone.

As a corollary of the previous theorem:

COROLLARY 3.15 Assume that Conditions (66), (67) and (68) are satisfied. Then the total equilibrium population  $X_T^*(D) = sx_{so}^*(D) + \bar{s}x_{si}^*(D)$  of (33) behaves like the total equilibrium population of the 2-patch source-sink model

$$\begin{cases}
\frac{dy_{so}}{dt} = r_{so}y_{so}\left(1 - \frac{y_{so}}{sK_{so}}\right) + D\left(\gamma_{2}y_{si} - \gamma_{1}y_{so}\right), \\
\frac{dy_{si}}{dt} = r_{si}y_{si}\left(-1 - \frac{y_{si}}{\overline{s}K_{si}}\right) + D\left(\gamma_{1}y_{so} - \gamma_{2}y_{si}\right).
\end{cases}$$
(75)

with specific growth rate  $r_{so}$ , death rate  $r_{si}$ , carrying capacities  $sK_{so}$ , parameter  $K_{si}$  due to the intraspecific competition in the sink patch and migration rates  $\gamma_1 = \frac{\Gamma_{I_{si}I_{so}}}{s}, \ \gamma_2 = \frac{\Gamma_{I_{so}I_{si}}}{\overline{s}}.$ 

PROOF The equilibrium point  $(x_{so}^*, x_{si}^*)$  is the positive solution of the following system:

$$\begin{cases} sr_{so}x_{so}\left(1-\frac{x_{so}}{K_{so}}\right) + D\left(\Gamma_{I_{so}I_{si}}x_{si}-\Gamma_{I_{si}I_{so}}x_{so}\right) = 0,\\ \overline{s}r_{si}x_{si}\left(-1-\frac{x_{si}}{K_{si}}\right) + D\left(\Gamma_{I_{si}I_{so}}x_{so}-\Gamma_{I_{so}I_{si}}x_{si}\right) = 0. \end{cases}$$
(76)

Therefore  $(y_{so}^* = sx_{so}^*, y_{si}^* = \overline{s}x_{si}^*)$  is the solution of the set of equations

$$\begin{cases} r_{so}y_{so}\left(1-\frac{y_{so}}{sK_{so}}\right) + D\left(\gamma_{2}y_{si}-\gamma_{1}y_{so}\right) = 0, \\ r_{si}y_{si}\left(-1-\frac{y_{si}}{\overline{s}K_{si}}\right) + D\left(\gamma_{1}y_{so}-\gamma_{2}y_{si}\right) = 0, \end{cases}$$
(77)

obtained from (76) by using the change of variables  $y_{so} = sx_{so}, y_{si} = \overline{s}x_{si}$ .

We can describe Conditions for which, under the conditions (66), (67) and (68), patchiness is beneficial or detrimental in System (33). We consider the regions in the set of the parameters  $\Gamma_{I_{so}I_{si}}$  and  $\Gamma_{I_{si}I_{so}}$ , denoted by  $\mathcal{L}_0$ ,  $\mathcal{L}_1$ ,  $\mathcal{L}_2$ ,  $\mathcal{L}_3$  and  $\mathcal{L}_4$ , depicted in Fig. 5 and defined by:

$$\begin{cases}
\text{If } r_{si} \ge r_{so} \text{ then} \begin{cases}
\mathcal{L}_{0} = \begin{cases} (\Gamma_{I_{so}I_{si}}, \Gamma_{I_{si}I_{so}}) : \frac{\Gamma_{I_{si}I_{so}}}{\Gamma_{I_{so}I_{si}}} < \frac{r_{si}}{r_{so}} \\
\mathcal{L}_{1} = \begin{cases} (\Gamma_{I_{so}I_{si}}, \Gamma_{I_{si}I_{so}}) : \frac{\Gamma_{I_{si}I_{so}}}{\Gamma_{I_{so}I_{si}}} \ge \frac{r_{si}}{r_{so}} \end{cases} \\
\end{cases}$$

$$\begin{cases}
\text{If } r_{si} < r_{so} \text{ then} \\
\mathcal{L}_{2} = \begin{cases} (\Gamma_{I_{so}I_{si}}, \Gamma_{I_{si}I_{so}}) : \frac{\Gamma_{I_{so}I_{si}}}{\Gamma_{I_{si}I_{so}}} \le \frac{r_{so}}{r_{si}} \end{cases} \\
\mathcal{L}_{3} = \begin{cases} (\Gamma_{I_{so}I_{si}}, \Gamma_{I_{si}I_{so}}) : \frac{r_{si}}{r_{so}} < \frac{\Gamma_{I_{si}I_{so}}}{\Gamma_{I_{so}I_{si}}} < \frac{K_{si}(r_{so} - r_{si})}{r_{so}(K_{so} + K_{si})} \\
\mathcal{L}_{4} = \begin{cases} (\Gamma_{I_{so}I_{si}}, \Gamma_{I_{sI}I_{so}}) : \frac{\Gamma_{I_{si}I_{so}}}{\Gamma_{I_{so}I_{si}}} \ge \frac{K_{si}(r_{so} - r_{si})}{r_{so}(K_{so} + K_{si})} \\
\end{cases}
\end{cases}$$

$$\end{cases}$$

$$\begin{cases}
(r_{1so}I_{si}, \Gamma_{I_{sI}I_{so}}) : \frac{\Gamma_{I_{si}I_{so}}}{\Gamma_{I_{so}I_{si}}} \ge \frac{K_{si}(r_{so} - r_{si})}{r_{so}(K_{so} + K_{si})} \\
\end{cases}$$

$$\end{cases}$$

$$\end{cases}$$



Figure 5: Qualitative properties of source-sink model (33) under Conditions (66),(67) and (68). In  $\mathcal{L}_0$  and  $\mathcal{L}_1$  the effect is detrimental with extinction in two patches for  $\mathcal{L}_0$  and persistence for  $\mathcal{L}_1$ . In  $\mathcal{L}_4$ , patchiness has a beneficial effect on total equilibrium population. In  $\mathcal{L}_2$  and  $\mathcal{L}_3$ , the effect is beneficial for  $D < D_0$  and detrimental for  $D > D_0$  with persistence of the population in the region  $\mathcal{L}_2$  and extinction in the region  $\mathcal{L}_3$ .

PROPOSITION 3.16 Assume that Conditions (66),(67) and (68) are satisfied, then, the total equilibrium population  $X_T^*(D) = sx_{so}^*(D) + \bar{s}x_{si}^*(D)$  of (33) satisfies the following properties

- 1. If  $r_{si} \ge r_{so}$ , let  $\mathcal{L}_0$  and  $\mathcal{L}_1$  be defined by (78) and depicted in Figure 5. Denote  $D^* = \frac{\Gamma_{I_{si}I_{so}}r_{so}r_{si}}{\Gamma_{I_{so}I_{si}}r_{si} - \Gamma_{I_{si}I_{so}}r_{so}}$ . Then we have:
  - if  $(\Gamma_{I_{so}I_{si}}, \Gamma_{I_{si}I_{so}}) \in \mathcal{L}_0$  then  $X_T^*(D) \leq sK_{so}$  for all  $D \geq 0$ . More precisely,

$$\begin{cases} 0 < X_T^*(D) \le sK_{so} & If \quad D < D^*, \\ X_T^*(D) = 0 & If \quad D \ge D^*. \end{cases}$$
(79)

- if  $(\Gamma_{I_{so}I_{si}}, \Gamma_{I_{si}I_{so}}) \in \mathcal{L}_1$  then  $0 < X_T^*(D) \le sK_{so}$  for  $D \ge 0$ .
- 2. If  $r_{si} < r_{so}$ , let  $\mathcal{L}_2, \mathcal{L}_3$  and  $\mathcal{L}_4$  be defined by (78) and depicted in Figure 5. Then we have:
  - if  $(\Gamma_{I_{so}I_{si}}, \Gamma_{I_{si}I_{so}}) \in \mathcal{L}_2$  then  $X_T^*(D) > sK_{so}$  for  $D < D_0$  and

**PROOF** The result is a consequence of Theorem 2.2 and Corollary 3.15.

**3.4. Death rates are much larger than the growth rates** In this part, we consider the multi-patch source-sink model (33) and we assume that the death rates of the sink patches are much larger than the growth rates of the source patches. Under this assumption, one can write the model in the matrix form as follow:

$$\begin{cases} \dot{X}_s = \operatorname{diag}\left(r_1 - \frac{r_1}{K_1}x_1, \cdots, r_s - \frac{r_s}{K_s}x_s\right)X_s + D\left(\Gamma_{ss}X_s + \Gamma_{sp}X_p\right), \\ \dot{X}_p = \frac{1}{\epsilon}\operatorname{diag}\left(-r_{s+1} - \frac{r_{s+1}}{K_{s+1}}x_{s+1}, \cdots, -r_n - \frac{r_n}{K_n}x_n\right)X_p + D\left(\Gamma_{ps}X_s + \Gamma_{pp}X_p\right), \end{cases}$$

$$\tag{82}$$

where  $\epsilon$  is assumed to be a small positive number. We have the following result:

THEOREM 3.17 Let  $(x_1(t, \epsilon), \ldots, x_n(t, \epsilon))$  be the solution of System (82) with initial condition  $(x_1^0, \ldots, x_n^0)$  satisfying  $x_i^0 \ge 0$  for  $i = 1, \ldots, n$ . Let  $u(t) = (u_1(t), \ldots, u_s(t))$  be the solution of the following differential system

$$\dot{X}_s = \operatorname{diag}\left(r_1 - \frac{r_1}{K_1}x_1, \dots, r_s - \frac{r_s}{K_s}x_s\right)X_s + D\Gamma_{ss},\tag{83}$$

with initial condition  $u(0) = (x_1^0, \ldots, x_s^0)$ ,  $X_s = (x_1, \ldots, x_s)^T$  and  $\Gamma_{ss}$  is the sub matrix of  $\Gamma$  defined by (39). Then, when  $\epsilon \to 0$ , we have

$$x_i(t,\epsilon) = u_i(t) + o_{\epsilon}(1), \quad i = 1, \dots, s \qquad uniformly \text{ for } t \in [0,T], \quad (84)$$

and

$$x_i(t,\epsilon) = o_\epsilon(1), \quad i = s+1,\dots,n,$$
(85)

uniformly for  $t \in [t_0, T]$ , where  $0 < t_0 < T$  are arbitrary but fixed and independent of  $\epsilon$ . If the solution  $u_s(t)$  of the reduced problem converges to an asymptotically stable equilibrium, then we can put  $T = +\infty$  in Approximations (84) and (85). PROOF When  $\epsilon \to 0$ , System (82) is a *slow-fast* system, with  $x_1, \ldots, x_s$  are slow variables, and  $x_{s+1}, \ldots, x_n$  fast variable. Tikhonov's theorem [32, 37, 38] prompts us to consider the dynamics of the fast variables in the time scale  $\tau = \frac{1}{D}t$ . One obtains

$$\dot{X}_p = \operatorname{diag}\left(-r_{s+1} - \frac{r_{s+1}}{K_{s+1}}x_{s+1}, \cdots, -r_n - \frac{r_n}{K_n}x_n\right)X_p + \epsilon D\left(\Gamma_{ps}X_s + \Gamma_{pp}X_p\right)$$
(86)

In the limit  $\epsilon \to 0$ , we find the fast dynamics

$$\dot{X}_p = \text{diag}\left(-r_{s+1} - \frac{r_{s+1}}{K_{s+1}}x_{s+1}, \cdots, -r_n - \frac{r_n}{K_n}x_n\right)X_p.$$
(87)

The slow manifold is given by the equilibrium of System (87), i.e  $X_p = 0$ , which is LAS in the positive axis. When  $\epsilon$  goes to zero, Tikhonov's theorem ensures that after a fast transition toward the slow manifold, the solutions of (82) converge to the solutions of the *reduced model* (83), obtained by replacing  $X_p = 0$  into the dynamics of the slow variable. Approximations (84) and (85) follow from Tikhonov's Theorem. Recall that when the reduced problem (83) has an asymptotically stable equilibrium, then these approximations hold for all t > 0 and not only on a compact interval [0, T]. Recall also that there is a boundary layer for the fast variables, that is Approximations (85) hold only for  $t > t_0$  where  $t_0 > 0$  can be arbitrarily small but fixed.

For the dynamics of the reduced model (83), we have the following result:

THEOREM 3.18 Consider System (83). Let A be the matrix defined by

$$A := \operatorname{diag}(r_1, \ldots, r_s) + D\Gamma_{ss}.$$

Assume that the matrix  $\Gamma_{ss}$  is irreducible, then we have:

- if  $S(A) \leq 0$ , the origin is GAS for (83), and
- if S(A) > 0, the reduced model has a GAS positive equilibrium.

PROOF As the matrix  $\Gamma_{ss}$  is irreducible, then the matrix A is also. Note that, the matrix A is the Jacobian matrix of the reduced model (83) evaluated at  $X_s = 0$ . According to [33, Corollary 1], we conclude the complete proof.

REMARK 3.2 Under the assumption that the matrix  $\Gamma_{ss}$  is irreducible, Approximations (84) and (85) hold for all t > 0 and not only on a compact interval [0, T].

Note that, we can write the reduced model (83) as follow:

$$\frac{dx_i}{dt} = r_i x_i \left(1 - \frac{x_i}{K_i}\right) - \gamma_i x_i + D \sum_{j=1}^s \gamma_{ij} x_j, \dots i = 1, \dots, s, \qquad (88)$$

where  $\gamma_i = D \sum_{j=s+1}^n \gamma_{ji}$  for  $i = 1, \ldots, s$ . We denote  $\tilde{\Gamma} := (\gamma_{ij})_{1 \le i,j \le s}$  the matrix which represent the migration between the source patches. In [20, Theorem 2.1], Gao have considered System (88) and proved under the assumptions  $\gamma_i > 0$ ,  $(K_1, \ldots, K_s) \in \ker \tilde{T}$  and  $\tilde{\Gamma}$  is irreducible, that, if  $\mathcal{R}_0 \le 0$ , the origin is GAS for (83), and if  $\mathcal{R}_0 > 0$ , there exist unique positive equilibrium which is GAS, where  $\mathcal{R}_0$  is the basic reproduction number of the reduced model (83) defined as:

$$\mathcal{R}_0 = \rho(FV^{-1})$$
, with  $F = \operatorname{diag}(r_1, \ldots, r_s)$  and  $V = \operatorname{diag}(\gamma_1, \ldots, \gamma_s) - D\tilde{\Gamma}$ ,

where  $\rho$  is the spectral radius.

**3.5. Growth rates are much larger than the death rates** In this part, we consider the multi-patch source-sink model (33) and we assume that the growth rates of the source patches are much larger than the death rates of the sink patches. Under this assumption, one can write the model in the matrix form as follow:

$$\begin{cases} \dot{X}_{s} = \frac{1}{\epsilon} \operatorname{diag} \left( r_{1} - \frac{r_{1}}{K_{1}} x_{1}, \cdots, r_{s} - \frac{r_{s}}{K_{s}} x_{s} \right) X_{s} \\ + D \left( \Gamma_{ss} X_{s} + \Gamma_{sp} X_{p} \right), \\ \dot{X}_{p} = \operatorname{diag} \left( -r_{s+1} - \frac{r_{s+1}}{K_{s+1}} x_{s+1}, \cdots, -r_{n} - \frac{r_{n}}{K_{n}} x_{n} \right) X_{p} \\ + D \left( \Gamma_{ps} X_{s} + \Gamma_{pp} X_{p} \right), \end{cases}$$

$$\tag{89}$$

where  $\epsilon$  is assumed to be a small positive number. We have the following result:

THEOREM 3.19 Let  $(x_1(t, \epsilon), \ldots, x_n(t, \epsilon))$  be the solution of System (82) with initial condition  $(x_1^0, \ldots, x_n^0)$  satisfying  $x_i^0 \ge 0$  for  $i = 1, \ldots, n$ . Let  $u(t) = (u_1(t), \ldots, u_p(t))$  be the solution of the differential system

$$\dot{X}_p = \operatorname{diag}\left(-r_{s+1} - \frac{r_{s+1}}{K_{s+1}}x_{s+1}, \dots, -r_n - \frac{r_n}{K_n}x_n\right)X_p + D\left(\Gamma_{ps}\mathcal{K}_s + \Gamma_{pp}X_p\right),$$
(90)

with initial condition  $(x_{s+1}^0, \ldots, x_n^0)$ ,  $X_p = (x_{s+1}, \ldots, x_n)^T$ ,  $\mathcal{K}_s = (K_1, \ldots, K_s)^T$ ,  $\Gamma_{pp}$  and  $\Gamma_{ps}$  are the sub matrices of  $\Gamma$  defined by (39). Then, when  $\epsilon \to 0$ , we have

$$x_i(t,\epsilon) = K_i + o_{\epsilon}(1), \quad i = 1, \dots, s \qquad uniformly \text{ for } t \in [0,T], \qquad (91)$$

and

$$x_i(t,\epsilon) = u_i(t) + o_\epsilon(1), \quad i = s + 1, \dots, n,$$
 (92)

uniformly for  $t \in [t_0, T]$ , where  $0 < t_0 < T$  are arbitrary but fixed and independent of  $\epsilon$ . If the solution  $u_p(t)$  of the reduced problem converges to an asymptotically stable equilibrium, then we can put  $T = +\infty$  in Approximations (91) and (92). **PROOF** The proof is the same as Theorem 3.17.

Our next goal is to prove the global stability of the reduced model (90) under the hypothesis that the two matrices  $\Gamma_{pp}$  and  $\Gamma$  are irreducible. First, we start by the following lemma:

LEMMA 3.20 Assume that the matrix  $\Gamma$  is irreducible. The reduced model (90) does not have the origin as equilibrium.

PROOF We suppose that the origin is an equilibrium of (90), then  $\Gamma_{ps}\mathcal{K}_s = 0$ , which is equivalent to  $\Gamma_{ps} = 0$ . So, we obtain a contradiction since  $\Gamma$  is irreducible.

THEOREM 3.21 Assume that the two matrices  $\Gamma_{pp}$  and  $\Gamma$  are irreducible. The reduced model (90) has unique equilibrium point in the interior of the positive cone  $\mathbb{R}^{n-s}_+ \setminus \{0\}$  which is GAS.

PROOF To show the global stability of the reduced model (90) in this case, we use the result of Hirsch [25] recalled in Theorem B.10.

The Jacobian matrix of the reduced model (90) is given by

$$G(X_p) := -\text{diag}\left(r_{s+1} + 2\frac{r_{s+1}}{K_{s+1}}x_{s+1}, \dots, r_n + 2\frac{r_n}{K_n}x_n\right) + D\Gamma_{pp},$$

which is irreducible because  $\Gamma_{pp}$  is also. Moreover, if  $G(X_p) \leq G(Y_p)$  then  $\operatorname{diag}(-r_i - 2\alpha_i x_i) \leq \operatorname{diag}(-r_i - 2\alpha_i y_i)$  which gives  $x_i \geq y_i$  for all *i*, i.e  $X_p \geq Y_p \geq 0$ . All solutions are bounded and the reduced model (90) does not admits the origin as equilibrium by Lemma 3.20. Hence, the reduced model (90) is globally stable according to Hirsch [25].

4. Multi-patch source-sink model without intraspecific competition in the sink patches In this section, we assume that, there is no intraspecific competition in n-s sink patches, i.e  $\alpha_i = 0$  for all  $i \ge s+1$  in System (33). Under this assumption, System (33) is rewritten as follows:

$$\begin{cases}
\frac{dx_i}{dt} = r_i x_i \left(1 - \frac{x_i}{K_i}\right) + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j - \gamma_{ji} x_i), & i = 1, \dots, s, \\
\frac{dx_i}{dt} = -r_i x_i + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j - \gamma_{ji} x_i), & i = s+1, \dots, n.
\end{cases}$$
(93)

System (93) was studied by Arino et al. [4] for n patches connected by migration terms and also by Wu et al. [39] for two patches. The global dynamics of (93) is given in [4, Theorem 1]. In all of this section, we denote  $\mathcal{E}^*(D)$  the positive equilibrium of (93) if it exists, and  $\mathcal{X}_T^*(D)$ , the total equilibrium population.

**4.1. The large migration rate** We have the following result which is a consequence of Theorem **3.8**.

COROLLARY 4.1 We have:

$$\lim_{D \to +\infty} \mathcal{E}^*(D) = \begin{cases} \frac{\sum_{i=1}^s \delta_i r_i - \sum_{i=s+1}^n \delta_i r_i}{\sum_{i=1}^s \delta_i^2 \alpha_i} (\delta_1, \dots, \delta_n), & \text{if } \sum_{i=1}^s \delta_i r_i > \sum_{i=s+1}^n \delta_i r_i, \\ 0 & \text{otherwise,} \end{cases}$$
(94)

where  $\alpha_i = r_i/K_i$ . Moreover, if the matrix  $\Gamma$  is symmetric, then:

$$\lim_{D \to +\infty} \mathcal{E}^*(D) = \begin{cases} \frac{\sum_{i=1}^s r_i - \sum_{i=s+1}^n r_i}{\sum_{i=1}^s \alpha_i} (1, \dots, 1), & \text{if } \sum_{i=1}^s r_i > \sum_{i=s+1}^n r_i, \\ 0 & \text{otherwise.} \end{cases}$$
(95)

PROOF Just replace  $\alpha_i = 0$  for i = s + 1, ..., n in Theorem 3.8.

According to the previous corollary, we obtain the formula of the total equilibrium population for perfect mixing:

$$\mathcal{X}_{T}^{*}(+\infty) = \begin{cases} \sum_{i=1}^{n} \delta_{i} \frac{\sum_{i=1}^{s} \delta_{i} r_{i} - \sum_{i=s+1}^{n} \delta_{i} r_{i}}{\sum_{i=1}^{s} \delta_{i}^{2} \alpha_{i}} & \text{if } \sum_{i=1}^{s} \delta_{i} r_{i} > \sum_{i=s+1}^{n} \delta_{i} r_{i}, \\ 0 & \text{otherwise.} \end{cases}$$

$$(96)$$

Moreover, if the matrix  $\Gamma$  is symmetric, then:

$$\mathcal{X}_{T}^{*}(+\infty) = \begin{cases} n \frac{\sum_{i=1}^{s} r_{i} - \sum_{i=s+1}^{n} r_{i}}{\sum_{i=1}^{s} \alpha_{i}}, & \text{if} & \sum_{i=1}^{s} r_{i} > \sum_{i=s+1}^{n} r_{i}, \\ 0 & \text{otherwise} & . \end{cases}$$

$$(97)$$

4.2. Derivative of the total equilibrium population In this section, our aim is to calculate the derivative of the total equilibrium population of System (93) at D = 0. First, we start by giving the following result:

LEMMA 4.2 Consider System (93). The total equilibrium population  $\mathcal{X}_T^*$  satisfies the following relation:

$$\mathcal{X}_{T}^{*}(D) = \sum_{i=1}^{s} K_{i} + D\left(\sum_{i=1}^{s} \sum_{i=1, j \neq i}^{n} \frac{\gamma_{ij} x_{j}^{*}(D) - \gamma_{ji} x_{i}^{*}(D)}{\alpha_{i} x_{i}^{*}(D)} + \sum_{i=s+1}^{n} \sum_{i=1, j \neq i}^{n} \frac{\gamma_{ij} x_{j}^{*}(D) - \gamma_{ji} x_{i}^{*}(D)}{r_{i}}\right).$$
(98)

PROOF If System (93) has unique equilibrium  $\mathcal{E}^*(D)$  in the interior of the positive cone, then it satisfies the following system:

$$\begin{cases} 0 = r_i x_i^*(D) \left( 1 - \frac{x_i^*(D)}{K_i} \right) + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)), & i = 1, \dots, s, \\ 0 = -r_i x_i^*(D) + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)), & i = s+1, \dots, n. \end{cases}$$
(99)

Dividing the first s equations in (99) by  $\alpha_i x_i^*(D)$ , and the last n-s equations by  $r_i$ , one obtain

$$\begin{cases} x_i^*(D) = K_i + D \sum_{i=1, j \neq i}^n \frac{\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)}{\alpha_i x_i^*(D)} & i = 1, \dots, s, \\ x_i^*(D) = D \sum_{i=1, j \neq i}^n \frac{\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)}{r_i} & i = s + 1, \dots, n. \end{cases}$$
(100)

Taking the sum of these expressions gives (98).

PROPOSITION 4.3 The derivative of the total equilibrium population  $\mathcal{X}_T^*$  at D = 0, is given by:

$$\frac{d\mathcal{X}_T^*}{dD}(0) = \left(\frac{1}{r_1}, \dots, \frac{1}{r_n}\right) \Gamma\left(K_1, \dots, K_s, 0, \dots, 0\right)^T.$$
(101)

**PROOF** Using matrix notation, the relation (99) is written as follow:

$$\mathcal{X}_{T}^{*}(D) = \sum_{i=1}^{s} K_{i} + D\left(\frac{1}{\alpha_{1}x_{i}^{*}(D)}, \dots, \frac{1}{\alpha_{s}x_{s}^{*}(D)}, \frac{1}{r_{s+1}}, \dots, \frac{1}{r_{n}}\right) \Gamma\left(x_{1}^{*}(D), \dots, x_{n}^{*}(D)\right)^{T}.$$
(102)

By differentiating Equation (102) at D = 0, we get:

$$\frac{d\mathcal{X}_T^*}{dD}(0) = \left(\frac{1}{\alpha_1 x_i^*(0)}, \dots, \frac{1}{\alpha_s x_s^*(0)}, \frac{1}{r_{s+1}}, \dots, \frac{1}{r_n}\right) \Gamma\left(x_1^*(0), \dots, x_n^*(0)\right)^T,$$
(103)

which gives (101), since  $x_i^*(0) = K_i$  for all i = 1, ..., s, and  $x_i^*(0) = 0$  for all i = s + 1, ..., n.

4.3. Comparison between results on (33) and the results on (93) In this part, our aim is to compare the results on (33) and the results on (93). We focus on two results on the total equilibrium population, the formulas of perfect mixing and the derivatives of the total equilibrium population at D = 0. We have the following result:

COROLLARY 4.4 Consider Systems (33) and (93) with the total equilibrium population  $X_T^*(D)$  and  $\mathcal{X}_T^*(D)$  respectively. Then,

$$\frac{dX_T^*}{dD}(0) = \frac{d\mathcal{X}_T^*}{dD}(0), \quad \text{and} \quad \mathcal{X}_T^*(+\infty) - X_T^*(+\infty) \begin{cases} > 0, & \text{if} \quad \sum_{i=1}^s \delta_i r_i > \sum_{i=s+1}^n \delta_i r_i, \\ = 0 & \text{otherwise.} \end{cases}$$

5. Conclusion The goal of this paper was to generalize to a multi-patch source-sink model the results obtained in [39] for a two-patch source-sink model and also the results of [14, 15] for muti-patch logistic model i.e. n-source 0-sink patch model. The diffusion between patches is modeled by a cooperative matrix. When this last matrix is irreducible, the system has a unique equilibrium, which furthermore is globally asymptotically stable (see Subsection 3.1).

In Subsection 3.2 we considered the particular case of perfect mixing, i.e. when the diffusion rate goes to infinity, that is, individuals may travel freely between patches. As in [39] for two-patch model and [14, 15], we compute the total equilibrium population in that case as a function of the number of the source patches, and, by perturbation arguments, we proved that the dynamics in this ideal case provides a good approximation for the case when the diffusion rate is large.

In Subsection 3.3 we considered the total equilibrium population in the n patches. We gave a complete solution in the case when the source and sink patches are partitioned into two blocks of identical patches ( source patches are identical and sink patches also). Our results mirror those of [39], which deals with the two-patch source-sink case (see Section 2). As shown in Proposition 3.11, diffusion could make total equilibrium population small than the sum of carrying capacities and also the extinction in both patches.

In Subsections 3.4 and 3.5, we study the total equilibrium population of the source-sink patch model (33) as a function of the diffusion rate in the case where the growth (resp. death) rate is much larger than the death (resp. growth) rate.

Some important questions remain open: Is there a way to make connections between the sources and sinks that increases the total equilibrium population? Mathematically speaking, are there conditions on the parameters of the model in which:

$$\sum_{i=1}^{s} \sum_{i=1, j \neq i}^{n} \frac{\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)}{\alpha_i x_i^*(D)} + \sum_{i=s+1}^{n} \sum_{i=1, j \neq i}^{n} \frac{\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)}{\alpha_i (K_i + x_i^*(D))}$$

is positive for all positive diffusion rate D? Anther problem, for example, for three-patch logistic model (one-source two-sink, two-source one-sink), is it possible to give a complete comparison between the total equilibrium population and the sum of the carrying capacities. I think this question is difficult and requires a lot of work and mathematical tools.

**A. Derivative of the total equilibrium population of** (33) First, we start by the following result:

LEMMA A.1 Consider System (33). The total equilibrium population  $X_T^*$  sat-

isfies the following relation:

$$X_T^*(D) = \sum_{i=1}^s K_i + D\left(\sum_{i=1}^s \sum_{i=1, j \neq i}^n \frac{\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)}{\alpha_i x_i^*(D)} + \sum_{i=s+1}^n \sum_{i=1, j \neq i}^n \frac{\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)}{\alpha_i (K_i + x_i^*(D))}\right).$$
(104)

PROOF If System (33) has unique equilibrium  $E^*(D)$  in the interior of the positive cone, then it satisfies the following system:

$$\begin{cases} 0 = r_i x_i^*(D) \left( 1 - \frac{x_i^*(D)}{K_i} \right) + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)), \quad i = 1, \dots, s, \\ 0 = r_i x_i^*(D) \left( -1 - \frac{x_i^*(D)}{K_i} \right) + D \sum_{j=1, j \neq i}^n (\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)), \quad i = s+1, \dots, n. \end{cases}$$

$$(105)$$

Dividing the first s equations in (105) by  $\alpha_i x_i^*(D)$ , and the last n-s equations by  $\alpha_i(K_i + x_i^*(D))$ , one obtain

$$\begin{cases} x_i^*(D) = K_i + D \sum_{i=1, j \neq i}^n \frac{\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)}{\alpha_i x_i^*(D)} & i = 1, \dots, s, \\ x_i^*(D) = D \sum_{i=1, j \neq i}^n \frac{\gamma_{ij} x_j^*(D) - \gamma_{ji} x_i^*(D)}{\alpha_i (K_i + x_i^*(D))} & i = s + 1, \dots, n. \end{cases}$$

$$(106)$$

Taking the sum of these expressions gives (104).

REMARK A.1 Using matrix notation, Relation (105) is written as follow:

$$X_T^*(D) = \sum_{i=1}^s K_i + Dv^T(D)\Gamma(x_1^*(D), \dots, x_n^*(D))^T, \qquad (107)$$

where 
$$v^T(D) = \left(\frac{1}{\alpha_1 x_i^*(D)}, \dots, \frac{1}{\alpha_s x_s^*(D)}, \frac{1}{\alpha_{s+1}(K_{s+1} + x_{s+1}^*(D))}, \dots, \frac{1}{\alpha_n(K_n + x_n^*(D))}\right)$$
.

PROPOSITION A.2 The derivative of the total equilibrium population at D = 0, is given by:

$$\frac{dX_T^*}{dD}(0) = \left(\frac{1}{r_1}, \dots, \frac{1}{r_n}\right) \Gamma\left(K_1, \dots, K_s, 0, \dots, 0\right)^T.$$
(108)

**PROOF** By differentiating Equation (107) at D = 0, we get:

$$\frac{dX_T^*}{dD}(0) = v^T(0)\Gamma\left(x_1^*(0), \dots, x_n^*(0)\right)^T.$$
(109)

which gives (108), since  $x_i^*(0) = K_i$  for all  $i = 1, \ldots, s$ , and  $x_i^*(0) = 0$  for all  $i = s + 1, \ldots, n$ .

For s = n, (108) becomes

$$\frac{dX_T^*}{dD}(0) = \left(\frac{1}{r_1}, \dots, \frac{1}{r_n}\right) \Gamma\left(K_1, \dots, K_n\right)^T,\tag{110}$$

which is the formula [14, Equation 28]. Note that, Formula (108) shows that, the derivative of the total equilibrium population at D = 0 is depend on growth and death rates  $r_i$ , the carrying capacities  $K_i$  for all  $i = 1, \ldots, s$ , and the sub matrices  $\Gamma_{ss}, \Gamma_{ps}$  of the matrix  $\Gamma$ . It is independents of the parameters  $K_i$  with  $i = s + 1, \ldots, n$ , and the sub matrix  $\Gamma_{pp}, \Gamma_{sp}$  of the matrices  $\Gamma$ 

**B.** Background concepts and preliminaries results In this section, our goal is to recall some concepts and results which we need in this work. Proofs of some results are given here and the others we refer interested readers to references.

DEFINITION B.1 The kernel of a matrix A is defined by ker  $A = \{v \in \mathbb{R}^n; Av = 0\}.$ 

DEFINITION B.2 A matrix  $A = (a_{ij})$  is called reducible, if there is a matrix of permutation  $P = (p_{ij})$  (i.e  $p_{ij} = 0$  or  $p_{ij} = 1$ ) such that:

$$P^T A P = \left[ \begin{array}{cc} A_{11} & A_{12} \\ 0 & A_{22} \end{array} \right],$$

where  $A_{11}$  and  $A_{22}$  are two square sub-matrices of A. We say that A is irreducible if A is non-reducible.

DEFINITION B.3 A matrix  $A = (a_{ij})$  is called cooperative if  $a_{ij} \ge 0$  for all  $i \ne j$ .

DEFINITION B.4 The stability modulus of a matrix A is given by

$$S(A) = \max \{ \operatorname{Re}(\lambda) : \lambda \text{ is an eigenvalue of } A \}, \qquad (111)$$

and the spectral radius of A is

$$\rho(A) = \max\{|\lambda|: \lambda \text{ is an eigenvalue of } A\}.$$
(112)

We have the following result [34, Lemma 8]:

LEMMA B.5 Let A be a non negative matrix. Let  $u \in \mathbb{R}^n_+$  be a non-zero vector and  $\lambda \in \mathbb{R}_+$  be a real number. If  $Au \ge \lambda u$  then  $\rho(A) \ge \lambda$ . If for a strictly positive vector u we have  $Au \le \lambda u$  then  $\rho(A) \le \lambda$ . PROOF If  $Au \geq \lambda u$  then, since A is non negative,  $A^k u \geq \lambda^k u$  for all k. Therefore  $||A^k|| \geq \lambda^k$  for any matricial norm. Using the Gelfand formula  $\rho(A) = \lim_{k \to \infty} ||A^k||^{\frac{1}{k}}$ , we obtain that  $\rho(A) \geq \lambda$ . The second statement is a simple consequence of the representation (2) in [34].

We have also the following result [7, Lemma 8]:

LEMMA B.6 Let A be a cooperative matrix. Let  $u \in \mathbb{R}^n$  and  $\lambda \in \mathbb{R}$ . If  $Au \geq \lambda u$ then  $S(A) \geq \lambda$ . If for a strictly positive vector u we have  $Au \leq \lambda u$  then  $S(A) \leq \lambda$ .

PROOF Let A be a cooperative matrix, there exists h > 0 such that A + hI, where I is the identity matrix, is non negative. Let u and  $\lambda$  be such that  $Au \ge \lambda u$ . Since  $S(A + hI)u \ge (\lambda + h)u$ , using Lemma B.5, we deduce that  $\rho(A+hI) \ge \lambda + h$ . According to the Perron-Frobenius Theorem [19, Theorem 3, page 66], we have

$$S(A+hI) = \rho(A+hI).$$

Therefore we have  $S(A+hI) \ge \lambda + h$ . Using S(A+hI) = S(A) + h, we obtain  $S(A) \ge \lambda$ . By the same method, we prove the second statement.

Let we consider the autonomous system:

$$\dot{x} = \Psi(x),\tag{113}$$

where  $\dot{x}$  denote the derivative of  $x, \Psi = (\Psi_1, \dots, \Psi_n)$  is  $\mathcal{C}^1$  on a domain  $\mathbb{R}^n_+$ .

DEFINITION B.7 Consider System (113). Let  $x(t, x_0)$  is a trajectory, and  $x_0$  is the initial point. The set  $\mathcal{O}$  of  $\mathbb{R}^n$  is said to be positively invariant if  $x_0 \in \mathcal{O}$  implies that  $x(t, x_0) \in \mathcal{O}$  for all  $t \geq 0$ . In other words, once a trajectory of the system enters  $\mathcal{O}$ , it will never leave it again.

DEFINITION B.8 System (113) is called cooperative if the Jacobian matrix  $\mathbb{J}\Psi(x)$  is a cooperative matrix for all  $x \in \mathbb{R}^n_+$ .

To prove the global stability of the system cooperative (113), generally, the following result is used:

THEOREM B.9 [16, 36] If system (113) possesses a positive equilibrium point  $x^*$  satisfying

$$\Psi_i(\xi x^*) \begin{cases} > 0 & \text{for} \quad \xi \in ]0, 1[, \\ < 0 & \text{for} \quad \xi > 1, \end{cases}$$
(114)

then  $x^*$  is globally stable.

We have also the following result of Hirsch [25]:

THEOREM B.10 If the cooperative system (113) has the following proprieties:

- $\mathbb{J}\Psi(x)$  is irreducible for any  $x \ge 0$ ,
- $\mathbb{J}\Psi(x) \leq \mathbb{J}\Psi(y)$  for any  $x \geq y \geq 0$ , and
- all solutions are bounded,

then either the origin is globally stable or there exists a unique positive equilibrium point and all the trajectories in  $\mathbb{R}^n_+ \setminus \{0\}$  tend to it.

## References

- R. Arditi, C. Lobry, and T. Sari. Is dispersal always beneficial to carrying capacity? New insights from the multi-patch logistic equation. *Theoretical Population Biology*, 106:45–59, dec 2015. ISSN 00405809. doi: 10.1016/j.tpb.2015.10.001. Cited on pp. 53, 54, and 59.
- [2] R. Arditi, C. Lobry, and T. Sari. Asymmetric dispersal in the multipatch logistic equation. *Theoretical Population Biology*, 120:11–15, mar 2018. ISSN 00405809. doi: 10.1016/j.tpb.2017.12.006. Cited on pp. 53, 59, and 73.
- [3] J. Arino. Diseases in Metapopulations. In Z. Ma, Y. Zhou, and J. Wu, editors, *Modeling and Dynamics of Infectious Diseases*, volume 11 of *Series in Contemporary Applied Mathematics*, pages 64–122. World Scientific Press, apr 2009. doi: 10.1142/9789814261265\_0003. Cited on p. 53.
- [4] J. Arino, N. Bajeux, and S. Kirkland. Number of Source Patches Required for Population Persistence in a Source–Sink Metapopulation with Explicit Movement. *Bulletin of Mathematical Biology*, 81(6):1916–1942, jun 2019. ISSN 0092-8240. doi: 10.1007/s11538-019-00593-1. Cited on pp. 52, 53, 60, 65, 66, 67, and 85.
- [5] C. Cosner, J. Beier, R. Cantrell, D. Impoinvil, L. Kapitanski, M. Potts, A. Troyo, and S. Ruan. The effects of human movement on the persistence of vector-borne diseases. *Journal of Theoretical Biology*, 258 (4):550–560, jun 2009. ISSN 00225193. doi: 10.1016/j.jtbi.2009.02.016. PMID: 19265711, PMCPMC2684576. Cited on p. 53.
- [6] F. Crick. Diffusion in Embryogenesis. Nature, 225(5231):420–422, jan 1970. ISSN 0028-0836. doi: 10.1038/225420a0. Cited on p. 52.
- [7] A. Cvetković. Stabilizing the Metzler matrices with applications to dynamical systems. *Calcolo*, 57(1):1, mar 2020. ISSN 0008-0624. doi: 10.1007/s10092-019-0350-3. Cited on p. 91.

- [8] D. DeAngelis, C. Travis, and W. Post. Persistence and stability of seed-dispersed species in a patchy environment. *Theoretical Population Biology*, 16(2):107–125, oct 1979. ISSN 00405809. doi: 10.1016/0040-5809(79)90008-X. Cited on pp. 53 and 65.
- [9] D. L. DeAngelis, W.-M. Ni, and B. Zhang. Dispersal and spatial heterogeneity: single species. *Journal of Mathematical Biology*, 72(1-2):239-254, jan 2016. ISSN 0303-6812. doi: 10.1007/s00285-015-0879-y. URL http://link.springer.com/10.1007/s00285-015-0879-y. Cited on p. 53.
- [10] D. L. DeAngelis, W.-M. Ni, and B. Zhang. Effects of diffusion on total biomass in heterogeneous continuous and discrete-patch systems. *Theoretical Ecology*, 9(4):443–453, dec 2016. ISSN 1874-1738. doi: 10.1007/s12080-016-0302-3. Cited on p. 53.
- [11] B. Elbetch. Effect of dispersal in two-patch environment with Richards growth on population dynamics. Journal of Innovative Applied Mathematics and Computational Sciences, 2(3):41–68, dec 2022. ISSN 2773-4196. doi: 10.58205/jiamcs.v2i3.47. Cited on p. 53.
- [12] B. Elbetch. Effects of rapid population growth on total biomass in Multipatch environment. Technical report, Centre pour la communication scientifique directe (CCSD), June 2022. URL https://hal.science/ hal-03698445. Working paper or preprint. pdf. Cited on p. 54.
- B. Elbetch and A. Moussaoui. Nonlinear diffusion in the multi-patch logistic model. Technical report, Centre pour la communication scientifique directe (CCSD), July 2022. URL https://hal.science/hal-03710815. Working paper or preprint. pdf. Cited on p. 53.
- [14] B. Elbetch, T. Benzekri, D. Massart, and T. Sari. The multi-patch logistic equation. *Discrete & Continuous Dynamical Systems B*, 26(12): 6405, 2021. ISSN 1531-3492. doi: 10.3934/dcdsb.2021025. Cited on pp. 53, 54, 55, 65, 69, 76, 88, and 90.
- [15] B. Elbetch, T. Benzekri, D. Massart, and T. Sari. The multi-patch logistic equation with asymmetric migration. *Revista Integración*, 40(1), mar 2022. ISSN 0120419X. doi: 10.18273/revint.v40n1-2022002. Cited on pp. 53, 54, 55, 65, 71, 73, 76, 78, and 88.
- [16] H. Freedman and Y. Takeuchi. Global stability and predator dynamics in a model of prey dispersal in a patchy environment. *Nonlinear Analysis: Theory, Methods & Applications*, 13(8):993–1002, 1989. ISSN 0362546X. doi: 10.1016/0362-546X(89)90026-6. Cited on p. 91.

- [17] H. Freedman, B. Rai, and P. Waltman. Mathematical models of population interactions with dispersal II: Differential survival in a change of habitat. *Journal of Mathematical Analysis and Applications*, 115(1): 140–154, apr 1986. ISSN 0022247X. doi: 10.1016/0022-247X(86)90029-6. Cited on p. 53.
- [18] H. I. Freedman and P. Waltman. Mathematical Models of Population Interactions with Dispersal. I: Stability of Two Habitats with and without a Predator. *SIAM Journal on Applied Mathematics*, 32(3):631–648, may 1977. ISSN 0036-1399. doi: 10.1137/0132052. Cited on p. 53.
- [19] F. R. Gantmacher. The theory of matrices. Vol. 2. Transl. from the Russian by K. A. Hirsch. Providence, RI: AMS Chelsea Publishing, reprint of the 1959 translation edition, 1998. ISBN 0-8284-0133-0; 0-8218-1393-5. Zbl 0927.15002. Cited on p. 91.
- [20] D. Gao. How Does Dispersal Affect the Infection Size? SIAM Journal on Applied Mathematics, 80(5):2144-2169, jan 2020. ISSN 0036-1399. doi: 10.1137/19M130652X. URL https://epubs.siam.org/doi/10.1137/ 19M130652X. Cited on pp. 53 and 84.
- [21] D. Gao and C.-P. Dong. Fast diffusion inhibits disease outbreaks. Proceedings of the American Mathematical Society, 148(4):1709-1722, dec 2019. ISSN 0002-9939. doi: 10.1090/proc/14868. URL https://www.ams.org/proc/2020-148-04/S0002-9939-2019-14868-1/. Cited on pp. 53 and 65.
- [22] D. Gao and S. Ruan. A Multipatch Malaria Model with Logistic Growth Populations. SIAM Journal on Applied Mathematics, 72(3):819-841, jan 2012. ISSN 0036-1399. doi: 10.1137/110850761. URL http://epubs. siam.org/doi/10.1137/110850761. Cited on p. 53.
- [23] H. Guo, M. Y. Li, and Z. Shuai. Global stability of the endemic equilibrium of multigroup SIR epidemic models. *Can. Appl. Math. Q.*, 14 (3):259–284, 2006. ISSN 1073-1849. Zbl 1148.34039. Cited on pp. 53 and 65.
- [24] I. Hanski and O. E. Gaggiotti, editors. Ecology, Genetics and Evolution of Metapopulations. Accademic Press, 2004. ISBN 9780123234483. doi: 10.1016/B978-0-12-323448-3.X5000-4. URL https://linkinghub.elsevier.com/retrieve/pii/B9780123234483X50004. Cited on p. 51.
- [25] M. W. Hirsch. The dynamical systems approach to differential equations. Bull. Am. Math. Soc., New Ser., 11:1–64, 1984. ISSN 0273-0979. doi: 10.1090/S0273-0979-1984-15236-4. Zbl 0541.34026. Cited on pp. 85 and 92.

- [26] R. D. Holt. Population dynamics in two-patch environments: Some anomalous consequences of an optimal habitat distribution. *Theoreti*cal Population Biology, 28(2):181–208, oct 1985. ISSN 00405809. doi: 10.1016/0040-5809(85)90027-9. Cited on p. 51.
- [27] R. A. Horn and C. R. Johnson. *Matrix analysis*. Cambridge: Cambridge University Press, 2nd ed. edition, 2013. ISBN 978-0-521-54823-6; 978-0-521-83940-2. Zbl 1267.15001. Cited on p. 67.
- [28] D. L. DeAngelis and B. Zhang. Effects of dispersal in a non-uniform environment on population dynamics and competition: A patch model approach. *Discrete & Continuous Dynamical Systems - B*, 19(10):3087– 3104, 2014. ISSN 1553-524X. doi: 10.3934/dcdsb.2014.19.3087. Cited on p. 53.
- [29] S. A. Levin. Dispersion and Population Interactions. *The Ameri*can Naturalist, 108(960):207–228, mar 1974. ISSN 0003-0147. doi: 10.1086/282900. Cited on p. 51.
- [30] S. A. Levin. Spatial patterning and the structure of ecological communities. In Some mathematical questions in biology, VII (Proc. Ninth Sympos. Math. Biol., New York, 1975), Lectures on Math. in the Life Sciences, Vol. 8, pages 1–35. Amer. Math. Soc., Providence, R.I., 1976. MR 0465284, Zbl 0338.92017. Cited on p. 51.
- [31] S. A. Levin, T. M. Powell, and J. W. Steele, editors. *Patch Dynamics*, volume 96 of *Lecture Notes in Biomathematics*. Springer Berlin Heidelberg, Berlin, Heidelberg, 1993. ISBN 978-3-540-56525-3. doi: 10.1007/978-3-642-50155-5. Cited on p. 51.
- [32] C. Lobry, T. Sari, and S. Touhami. On Tykhonov's theorem for convergence of solutions of slow and fast systems. *Electron. J. Differ. Equ.*, 1998:22, 1998. ISSN 1072-6691. Id/No 19. https://eudml.org/doc/119570, Zbl 0897.34052, EuDML: 119570. Cited on pp. 60, 62, 69, 71, and 83.
- [33] Z. Y. Lu and Y. Takeuchi. Global asymptotic behavior in single-species discrete diffusion systems. J. Math. Biol., 32(1):67–77, 1993. ISSN 0303-6812. doi: 10.1007/BF00160375. Cited on pp. 68 and 83.
- [34] Y. Nesterov and V. Y. Protasov. Computing Closest Stable Nonnegative Matrix. SIAM Journal on Matrix Analysis and Applications, 41(1):1– 28, jan 2020. ISSN 0895-4798. doi: 10.1137/17M1144568. URL https: //epubs.siam.org/doi/10.1137/17M1144568. Cited on pp. 90 and 91.
- [35] J.-C. Poggiale, P. Auger, D. Nérini, C. Manté, and F. Gilbert. Global Production Increased by Spatial Heterogeneity in a Population Dynamics

Model. Acta Biotheoretica, 53(4):359–370, dec 2005. ISSN 0001-5342. doi: 10.1007/s10441-005-4890-3. URL http://link.springer.com/10. 1007/s10441-005-4890-3. Cited on p. 73.

- [36] Y. Takeuchi. Cooperative Systems Theory and Global Stability of Diffusion Models. In *Evolution and Control in Biological Systems*, pages 49–57. Springer Netherlands, Dordrecht, 1989. doi: 10.1007/978-94-009-2358-4\_6. Cited on pp. 53 and 91.
- [37] A. N. Tihonov. Systems of differential equations containing small parameters in the derivatives. *Mat. Sbornik (N.S.)*, 31(73):575-586, 1952. http://refhub.elsevier.com/S0040-5809(15)00102-1/sbref18; MR 0055515. Cited on pp. 60, 62, 69, 71, and 83.
- [38] W. Wasow. Asymptotic expansions for ordinary differential equations. Robert E. Krieger Publishing Co., Huntington, N.Y., 1976. ISBN 0-88275-173-5. Reprint of the 1965 edition. MR 0460820. Cited on pp. 60, 62, 69, 71, and 83.
- [39] H. Wu, Y. Wang, Y. Li, and D. L. DeAngelis. Dispersal asymmetry in a two-patch system with source-sink populations. *Theoretical Population Biology*, 131:54-65, feb 2020. ISSN 00405809. doi: 10.1016/j.tpb.2019.11.004. URL https://linkinghub.elsevier.com/retrieve/pii/S0040580919301935. Cited on pp. 52, 55, 56, 57, 58, 59, 60, 66, 73, 85, and 88.
- [40] F. Yu, W. Song, and Y. Wang. Effect of diffusion on a consumerresource system with source-sink patches. *Discrete and Continuous Dynamical Systems - B*, 28(3):1905–1937, 2023. ISSN 1531-3492. doi: 10.3934/dcdsb.2022152. Cited on p. 55.
- [41] B. Zhang, X. Liu, D. DeAngelis, W.-M. Ni, and G. G. Wang. Effects of dispersal on total biomass in a patchy, heterogeneous system: Analysis and experiment. *Mathematical Biosciences*, 264:54–62, jun 2015. ISSN 00255564. doi: 10.1016/j.mbs.2015.03.005. Cited on p. 53.
- [42] B. Zhang, D. L. DeAngelis, W.-M. Ni, Y. Wang, L. Zhai, A. Kula, S. Xu, and J. D. Van Dyken. Effect of Stressors on the Carrying Capacity of Spatially Distributed Metapopulations. *The American Naturalist*, 196 (2):E46–E60, aug 2020. ISSN 0003-0147. doi: 10.1086/709293. PMID: 32673100. Cited on p. 55.

## Efekt rozproszenia w jednogatunkowych systemach dyfuzji dyskretnej z populacjami wykluczanymi ekologicznie. Bilel Elbetch

Streszczenie Rozważany jest model wielogatunkowy enklaw wykluczanych ekologicznie z i bez konkurencji wewnątrzgatunkowej w enklawach zanikających. Najpierw badamy dynamikę modelu, gdy macierz migracji jest nieredukowalna i redukowalna. Pokazujemy, że istnieje progowa liczba enklaw, która powoduje, że populacja potencjalnie wymiera poniżej progu i jest ustabilizowana powyżej pewnego pozomu. Następnie, korzystając z teorii perturbacji pojedynczych i twierdzenia Tichonowa, w przypadku doskonałego mieszania, tj. gdy współczynnik dyfuzji dąży do nieskończoności, obliczamy równowagę modelu i podajemy dobre przybliżenia rozwiązań w tym przypadku. Po drugie, określamy, w pewnych szczególnych przypadkach, warunki, w których fragmentacja i istnienie enklaw może prowadzić do tego, że całkowita populacja w równowadze jest większa lub mniejsza niż suma populacji enklaw. Wreszcie, badamy wpływ szybkiego wzrostu populacji źródłowej i szybkiego wymierania populacji populacji zlewowej na dynamikę całkowitej populacji równowagi i na koegzystencję gatunków

2010 Klasyfikacja tematyczna AMS (2010): Primary: 37N25; Secondary: 34D23, 34D15, 92D25..

*Stowa kluczowe:* Population Dynamics, Logistic equation, diffusion rate, Slow-fast systems, Tikhonov's theorem, Perfect mixing..



Bilel Elbetch<sup>a</sup> completed his doctorate (mathematics) at University of Dr. Moulay Tahar, Saida, Algeria, under the supervision of professors Tounsia Benzekri, Daniel Massart and Tewfik Sari. His main academic interests are mathematical modeling and dynamical systems, more specifically, the effects of migration and spatial heterogeneity on the dynamics of a population and on the coexistence of species.

BILEL ELBETCH D UNIVERSITY DR. MOULAY TAHAR OF SAIDA DEPARTMENT OF MATHEMATICS BP 138 CITÉ ENNASR 20000, SAÏDA, ALGÉRIE *E-mail:* elbetchbilal@gmail.com

Communicated by: Urszula Foryś

(Received: 23rd of September 2022; revised: 23rd of December 2022)

<sup>&</sup>lt;sup>a</sup>References to his research papers are found in MathSciNet under ID: 1475779 and the European Mathematical Society, FIZ Karlsruhe, and the Heidelberg Academy of Sciences bibliography database known as zbMath under ai:Elbetch.Bilel.