EXPONENTIAL CONVERGENCE TO EQUILIBRIUM VIA LYAPOUNOV FUNCTIONALS FOR REACTION-DIFFUSION EQUATIONS ARISING FROM NON REVERSIBLE CHEMICAL KINETICS

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Abstract. We show that the entropy method, that has been used successfully in order to prove exponential convergence towards equilibrium with explicit constants in many contexts, among which reaction-diffusion systems coming out of reversible chemistry, can also be used when one considers a reaction-diffusion system corresponding to an irreversible mechanism of dissociation/recombination, for which no natural entropy is available.

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1. Introduction

1.1. Entropy methods

Entropy methods have recently been used in order to prove exponential convergence towards the equilibrium with explicit constants in many situations (e.g. integral equations, cf. [20], fourth order equations, cf. [5], nonlinear diffusion equations, cf. [6,7,12]). A nice survey of these methods may be found in the review paper [1]. In particular, reaction-diffusion equations in the context of reversible chemistry have been systematically studied in [9–11].

We recall that the principle of this method is to find a Lyapounov functional $E(f)$ and its dissipation $D(f)$ such that

$$\partial_t E(f) = -D(f) \leq 0,$$

when $f$ is a solution of the equation, and such that the following (sometimes called entropy/entropy dissipation) functional inequality holds:

$$D(f) \geq C(E(f) - E(f_{eq})),$$

where $f_{eq}$ is the unique minimum of $E$ (once conservations have been taken into account). In this situation, it is usually possible to prove that

$$\| f - f_{eq} \| \leq C_1 e^{-C_2 t},$$

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where $C_1$ and $C_2$ are explicit (note that a linearization usually leads to exponential decay, but with a constant $C_1$ which is not explicit).

For reaction-diffusion equations appearing in reversible chemistry, it is in general possible to take for $E$ the natural physical entropy of the problem (such an entropy exists because the equations can be obtained as the limit of kinetic equations of Boltzmann type describing the microscopic processes, cf. [2]).

This paper is devoted to showing that in a typical example of irreversible chemistry in which only one equilibrium appears, it is also possible to find a Lyapounov functional $E$ which satisfies the requirements of the entropy method. The model that we intend to study is related to a set of dissociation/recombination chemical reactions.

1.2. A model of dissociation/recombination

We consider a diatomic gas with dissociation/recombination reactions, made up by atoms $A$ with mass $m_1$ and molecules $A_2$ with mass $m_2 = 2m_1$. The two species in the binary mixture are labeled by an index $i = 1, 2$. Generally speaking, the reaction-diffusion system is expected in the form

$$ \partial_t n_i - d_i \Delta_x n_i = Q_i(n_1, n_2), \quad i = 1, 2, $$

(1.1)

where $n_i$ denotes number density, $d_i$ diffusion coefficient, and $Q_i$ the chemical source term. Since all chemical encounters preserve the global mass of participating species (or, equivalently, preserve the total number of atoms), any reasonable dissociation/recombination model must fulfil “a priori” the consistency constraint

$$ Q_1(n_1, n_2) + 2Q_2(n_1, n_2) = 0. $$

(1.2)

If the mixture is embedded in a fixed background, to be labeled by an additional index $i = 0$, dissociation reactions may occur by binary encounter of a molecule $A_2$ with any of the possible collision partners (field particle $A_0$, single atom $A_1$, or other molecule $A_2$), whereas a recombination reaction is due solely to a binary interaction of two atoms between themselves. In the first process, one molecule is lost and two atoms are gained in the collision balance. In the second process, two atoms coalesce in one molecule, and the balance is just reversed. Therefore, the simplest heuristic model one could think of in order to describe the reaction effects on the whole evolution problem is represented by

$$ Q_2^i(n_1, n_2) = \alpha_{11}^i(n_1)^2 - (\alpha_{20}^i n_0 + \alpha_{21}^i n_1 + \alpha_{22}^i n_2)n_2, $$

(1.3)

where $\alpha_{11}^j$ and $\alpha_{2j}^i$ ($j = 0, 1, 2$) are suitable averaged rate constants quantifying the probability of a recombination collision $A_1-A_1$, or of a dissociation collision $A_2-A_j$, respectively. Of course, $Q_2^i$ follows from (1.2), and superscripts are used to denote the type of reaction. The constant background density $n_0$ is given.

The same problem can be tackled at the kinetic level [18], in the frame of a recently developed literature (see for instance [13]). According to a common kinetic model, the gas is described as a mixture of three species, with an additional component, labeled by $i = 3$, representing unstable molecules $A_3 \equiv A_2^*$ (with mass $m_3 = m_2$) and playing the role of a transition state [21]. The mixture is then taken to diffuse in a much denser medium [3], whose evolution is not affected by the collisions going on, assumed in local thermodynamical equilibrium, namely with distribution function $f_0 = n_0 M_0$, where $M_0$ stands for the normalized Maxwellian

$$ M_0 = \left( \frac{m_0}{2\pi T_0} \right)^{\frac{3}{2}} \exp \left( - \frac{m_0}{2T_0} |\psi|^2 \right) $$

(1.4)

and $T_0$ is also constant. According to the model, both atoms $A_1$ and stable molecules $A_2$ may undergo elastic collisions with other atoms, stable molecules and background particles. Moreover, atoms $A_1$ may form a stable molecule $A_2$ passing through the transition state $A_2^*$, while, on the other hand, both stable and unstable diatomic
molecules may dissociate into two atoms. More precisely, the recombination process occurs in two steps:

\[(R) \quad A_1 + A_1 \rightarrow A_2^2 \]
\[(I) \quad A_2^2 + P \rightarrow A_2 + P, \]

where \(P = A, A_2, B\), while dissociation occurs via two possible reactions:

\[(D1) \quad A_2 + P \rightarrow 2 A_1 + P \]
\[(D2) \quad A_2^2 + P \rightarrow 2 A_1 + P. \]

It must be stressed that all above interactions, modelling the chemical reactions at the kinetic level, have to be understood as irreversible processes. Chemical operators are given in terms of total microscopic collision frequencies \(\nu_{ij}^s\) (constant for Maxwellian molecules), where the superscript \(k\) may take the values \(s, r, i, d\), corresponding to elastic scattering, recombination \(R\), inelastic scattering \(I\), dissipations \(D1, D2\), respectively, and of suitable transition probabilities, accounting for the correct exchange rates for mass, momentum, and various forms of energy \([14]\). Then, kinetic integrodifferential equations have been scaled in terms of the typical relaxation times, a small parameter defining the dominant process(es) has been introduced, and the formal various forms of energy \([14]\). Then, kinetic integrodifferential equations have been scaled in terms of the typical relaxation times, a small parameter defining the dominant process(es) has been introduced, and the formal asymptotic limit when this parameter vanishes has been consistently investigated \([3]\). This leads to the derivation of hydrodynamic limiting equations, whose nature varies considerably according to the relative importance of the various processes and to the corresponding pertinent scaling, but which are typically of reaction-diffusion type as long as the scattering with the background plays an important role. Some non-exhaustive examples were given in \([3]\) itself, and also in \([4]\). However, we shall deal here with one of the asymptotic limits which seems more realistic in practice \([3]\), and leads to (1.1), (1.2) with specialization

\[
Q_2(n_1, n_2) = \frac{\nu_{30}^i n_0 + \nu_{31}^i n_1 + \nu_{32}^j n_2}{\nu_{30}^i n_0 + \nu_{31}^i n_1 + \nu_{32}^j n_2} \nu_1^{11} (n_1)^2 - \left( \nu_{20}^d n_0 + \nu_{21}^d n_1 + \nu_{22}^d n_2 \right) n_2 \tag{1.5}
\]

with \(\nu_{3j}^i = \nu_{3j}^i + \nu_{3j}^d, \ j = 0, 1, 2\). This rather simple expression was derived under the simplifying assumption of Maxwell-type interactions, in which reactive collision frequencies are constant. However, with respect to (1.3), this expression shows a more complicated and realistic dependence on the participating species densities (a rational function rather than a quadratic polynomial), with rates well defined in terms of microscopic parameters. The same is true for the (positive) diffusion coefficients, which take the form

\[
d_1 = \frac{m_1 + m_0}{2 m_1 m_0} \frac{T_0}{\nu_{10}^i n_0}, \quad d_2 = \frac{2 m_1 + m_0}{4 m_1 m_0} \frac{T_0}{\nu_{20}^d n_0}, \tag{1.6}
\]

where the constants \(\nu_{10}^i\) are suitable angle averaged scattering collision frequencies \([3]\). The fraction in (1.5) accounts for the important physical fact that recombination occurs via a transition state, and it is remarkable that the microscopic parameters of this metastable species, \(\nu_{3j}^i\) and \(\nu_{3j}^d\), do influence the evolution of the two stable species, though the third species is not present at the hydrodynamic level (its density has collapsed to zero in the asymptotic limit). This is a so-called ghost-effect, not unfrequent in fluid dynamics \([19]\), namely a trace in the evolution of something that does not exist. This is due in our case to the clearance of an indeterminate form, coming from the simultaneous vanishing of the species density and of its relaxation times. It can be noticed that the simple heuristic model (1.3) may be considered as a special case of the more physical model (1.5) under the simplifying assumption \(\nu_{3j}^i = 0, \ j = 0, 1, 2\), with all rate constants \(\alpha_{ij}^k\) provided exactly by the corresponding microscopic collision frequencies \(\nu_{ij}^k\). Another special case of the same type as (1.3) is achieved by assuming instead \(\nu_{3j}^i = \eta \nu_{3j}^d, \ j = 0, 1, 2\), with \(0 < \eta < 1\); in this case the rate constant for recombination \(\alpha_{11}^r\) would be given by the microscopic recombination collision frequency \(\nu_{11}^r\) reduced by the factor \(\eta\).
1.3. Exponential convergence towards equilibrium

The main goal of this paper is to show that the solution of system (1.1)–(1.2)–(1.5), together with the Neumann boundary conditions
\[ \hat{\mathbf{n}}(\mathbf{x}) \cdot \nabla \mathbf{x} n_i = 0, \quad \mathbf{x} \in \partial \Omega, \] (1.7)
(where \( \hat{\mathbf{n}}(\mathbf{x}) \) is the outward normal unit vector to the spatial domain \( \Omega \) at point \( \mathbf{x} \)) and the nonnegative initial conditions
\[ n_i(0, \mathbf{x}) = n_i^0(\mathbf{x}) \geq 0, \] (1.8)
converges exponentially fast with explicit constants towards the unique equilibrium of the system. Since total number of atoms is preserved, we have that
\[ \forall t \geq 0, \quad \int_{\Omega} \left( n_1(t, \mathbf{x}) + 2 n_2(t, \mathbf{x}) \right) d\mathbf{x} = \int_{\Omega} \left( n_1^0(\mathbf{x}) + 2 n_2^0(\mathbf{x}) \right) d\mathbf{x} = \hat{n}^0, \]
or, equivalently, setting \( \bar{n}_i(t) = \int_{\Omega} n_i(t, \mathbf{x}) d\mathbf{x} \),
\[ \forall t \geq 0, \quad \bar{n}_1(t) + 2 \bar{n}_2(t) = \bar{n}^0. \] (1.9)

We are able to prove the following theorem:

**Theorem 1.1.** Let \( \Omega \) be a bounded regular \((C^2)\) open set of \( \mathbb{R}^n \), let \( n_i^0 > 0 \) \((i = 1, 2)\) be initial data in \( C^2(\bar{\Omega}) \) compatible with Neumann boundary conditions. Finally, let \( \nu_{1j}, \nu_{2j}, \nu_{3j} \) \((j = 0, 1, 2)\), \( \nu_{11}, d_1, d_2 \) be strictly positive constants.

Then, there exists a unique strong \((C^2(\mathbb{R}^+ \times \Omega))\) solution \( n_1, n_2 \) to system (1.1)–(1.2)–(1.5) with boundary conditions (1.7) and initial data \( n_i^0 \) \((i = 1, 2)\). This solution is bounded from above and from below:
\[ k_1 \leq n_i(t, \mathbf{x}) \leq K_1, \quad k_2 \leq n_2(t, \mathbf{x}) \leq K_2, \]
where \( k_1, k_2, K_1 \) and \( K_2 \) are strictly positive constants depending only on corresponding bounds for initial data, and moreover it satisfies
\[ \sum_{i=1}^{2} \| n_i(t, \cdot) - n_i^* \|_{L^2(\Omega)}^2 \leq C_1 e^{-\frac{C_2 t}{2}} , \]
where \( C_1 \) and \( C_2 \) are explicitly computable. Here \( n_i^* \) is the unique positive \( \text{(independent of } \mathbf{x} \text{)} \) solution of
\[ Q_i(n_i^*, n_2^*) = 0, \quad i = 1, 2, \]

satisfying the conservation (1.9).

As announced in Section 1.1, the method of proof will consist in finding a suitable Lyapunov functional \( E(n_1, n_2) \) and its dissipation \( D(n_1, n_2) \) such that
\[ \partial_t E(n_1, n_2) = -D(n_1, n_2) \leq 0, \]
when \( n_1, n_2 \) is a solution of system (1.1)–(1.2)–(1.5), and such that
\[ D(n_1, n_2) \geq C \left( E(n_1, n_2) - E(n_i^*, n_2^*) \right), \] (1.10)
for any \( n_1, n_2 \) \((\text{functions of } \mathbf{x} \text{ only})\) satisfying the same \textit{a priori} assumptions as those of system (1.1)–(1.2)–(1.5), \textit{i.e.} conservation of mass, minimum and maximum principle. A suitable scaling of the space variable \( \mathbf{x} \) allows us to carry out the proof under the assumption \(|\Omega| = 1\), without loss of generality.
In Section 2, we begin by treating a particular case in which the computations are quite simple and make the method easily understandable. In that section, we do not prove in detail the existence and uniqueness of the solution of the system, for the sake of conciseness.

Then, in Section 3, we treat the general case. Section 3.1 is devoted to a first study of the reaction terms $Q_i$. Then, in Section 3.2 we present the minimum/maximum principle and existence/uniqueness of solutions to system (1.1)–(1.2)–(1.5). Finally, Section 3.3 is devoted to the establishment of estimate (1.10), which enables to recover the result of exponential convergence with explicit rates.

Remark 1.2. The theorem in the present version will be proved in Section 3. The assumption on strict positivity for the collision frequencies $\nu^d_{ij}$ can be easily weakened, and several of them can be allowed to vanish, making proofs easier, as shown indeed by the simpler case dealt with preliminarily in Section 2. From a mathematical point of view, an interesting question consists in asking if it is really necessary that both diffusivities are strictly positive. We shall show in Remark 3.8 that if one of them vanishes, the results remain true (though with different constants).

Remark 1.3. Concerning computability of the constants in the decay estimate, explicit formulas are too involved to be given in the text of the theorem. However, they can be estimated in terms of the initial distributions $\nu^0_i$ (specifically, of their upper and lower bounds), of the domain $\Omega$ (specifically, of its Poincaré constant), in addition to the physical parameters $\nu^d_{ij}$, $\nu^d_{3j}$, $\nu^d_{j3}$ ($j = 0, 1, 2$), $\nu^s_{1j}$, and the diffusivity constants (more precisely, a lower bound of one of them) $d_1$, $d_2$. The explicit formulas can be put together anyhow by following the various steps of the proofs and the relevant estimates, which are all given in full detail.

2. PARTICULAR CASE

For readers’ convenience, we introduce at first a self-contained particular case, in which the main steps of our procedure may be summarized without tedious technical complications. The generalization to the collision contributions (1.5) will be presented in next section.

If we assume $\nu^d_{3j} = 0$, $j = 0, 1, 2$, and $\nu^d_{20} = 0$ in (1.5), system (1.1) takes the form

\[
\begin{align*}
\frac{\partial n_1}{\partial t} - d_1 \Delta x n_1 &= -2 \nu^s_{11} (n_1)^2 + 2 \left( \nu^d_{21} n_1 + \nu^d_{22} n_2 \right) n_2, \\
\frac{\partial n_2}{\partial t} - d_2 \Delta x n_2 &= \nu^s_{11} (n_1)^2 - \left( \nu^d_{21} n_1 + \nu^d_{22} n_2 \right) n_2,
\end{align*}
\]

so that, after a suitable rescaling, we have to deal with the dimensionless equations

\[
\begin{align*}
\frac{\partial n_1}{\partial t} - d_1 \Delta x n_1 &= -2 (n_1)^2 + 4 \alpha n_1 n_2 + 2 \beta (n_2)^2 := Q^*_1(n_1, n_2), \\
\frac{\partial n_2}{\partial t} - d_2 \Delta x n_2 &= (n_1)^2 - 2 \alpha n_1 n_2 - \beta (n_2)^2 := Q^*_2(n_1, n_2),
\end{align*}
\]

where

\[
\alpha = \frac{\nu^d_{21}}{2 \nu^s_{11}} > 0, \quad \beta = \frac{\nu^d_{22}}{\nu^s_{11}} > 0.
\]

2.1. Study of the reaction terms

Collision equilibria for the set (2.2) are given by the nonnegative solutions of the equation $Q^*_1(n_1, n_2) = 0$. It can be trivially checked that

\[
Q^*_2(n_1, n_2) = (n_1 - \gamma n_2) (n_1 + \delta n_2),
\]

where

\[
\gamma = \alpha + \sqrt{\alpha^2 + \beta} > 0, \quad \delta = -\alpha + \sqrt{\alpha^2 + \beta} > 0,
\]
hence the physical equilibrium states are characterized by \( n_1 = \gamma n_2 \). Taking into account the conservation property (1.9), since the initial data are known we get that the unique global (homogeneous in space) equilibrium is determined by the positive constants
\[
n_1^* = \frac{\gamma}{2 + \gamma} \bar{n}^0, \quad n_2^* = \frac{1}{2 + \gamma} \bar{n}^0. \tag{2.5}
\]

Notice that \( Q_2^* \geq 0 \iff n_1 \geq \gamma n_2 \), and conversely for \( Q_1^* \).

This suggests that a suitable entropy could be given by
\[
E(n_1, n_2) = \int_{\Omega} \left( \frac{1}{4} (n_1)^2 + \frac{\gamma}{2} (n_2)^2 \right) \, dx,
\tag{2.6}
\]
since, in space homogeneous conditions,
\[
\partial_t E(n_1, n_2) = - \int_{\Omega} (n_1 - \gamma n_2)^2 (n_1 + \delta n_2) \, dx \leq 0,
\]
with \( \partial_t E = 0 \) only at the local equilibrium \( n_1 = \gamma n_2 \).

2.2. Minimum principle

We now turn to a result of “minimum principle” type for equation (2.2).

**Proposition 2.1.** Let \( d_1, d_2 > 0, \alpha, \beta > 0 \), and \( \Omega \) be a bounded regular \((C^2)\) open set of \( \mathbb{R}^N \). Let \((n_1(t, x), n_2(t, x))\) be a strong solution (that is, in \( C^2(\mathbb{R}^+ \times \Omega) \)) to system (2.2) with Neumann boundary conditions (1.7) and with initial conditions such that
\[
n_1(0, x) = n_1^0(x) > c_1 > 0, \quad n_2(0, x) = n_2^0(x) > c_2 > 0. \tag{2.7}
\]

This solution \((n_1(t, x), n_2(t, x))\) is strictly positive for \((t, x) \in [0, \infty) \times \Omega\), and satisfies the following lower bounds:
\[
n_1(t, x) \geq k_1, \quad n_2(t, x) \geq k_2, \tag{2.8}
\]
where
\[
k_1 = \min \{ c_1, \gamma c_2 \}, \quad k_2 = \gamma^{-1} k_1. \tag{2.9}
\]

**Proof of Proposition 2.1.** The proof shall be carried out following the same lines as in reference [15]. For any \( \varepsilon > 0 \), let us consider the functions
\[
n_1^\varepsilon(t, x) = n_1(t, x) e^{\varepsilon t}, \quad n_2^\varepsilon(t, x) = n_2(t, x) e^{\varepsilon t}, \tag{2.10}
\]
and let us prove that
\[
n_1^\varepsilon(t, x) > k_1, \quad n_2^\varepsilon(t, x) > k_2. \tag{2.11}
\]
From equations (2.2), it follows that the evolution of \( n_1^\varepsilon, n_2^\varepsilon \) is governed by the system
\[
\begin{align*}
\partial_t n_1^\varepsilon - d_1 \Delta_x n_1^\varepsilon &= \left[ - 2 (n_1^\varepsilon)^2 + 4 \alpha n_1^\varepsilon n_2^\varepsilon + 2 \beta (n_2^\varepsilon)^2 \right] e^{-\varepsilon t} + \varepsilon n_1^\varepsilon, \\
\partial_t n_2^\varepsilon - d_2 \Delta_x n_2^\varepsilon &= \left[ (n_1^\varepsilon)^2 - 2 \alpha n_1^\varepsilon n_2^\varepsilon - \beta (n_2^\varepsilon)^2 \right] e^{-\varepsilon t} + \varepsilon n_2^\varepsilon.
\end{align*} \tag{2.12}
\]
Suppose that the inequalities (2.11) do not hold for all \((t, x) \in [0, \infty) \times \Omega\), and define the set \(B^c\) as
\[
B^c = \left\{ \tau > 0 : n_1^c(t, x) > k_1, \quad n_2^c(t, x) > k_2, \quad \forall (t, x) \in [0, \tau) \times \Omega \right\}.
\] (2.13)
If we denote \(\hat{t} = \sup B^c\), there must exist \(\hat{x} \in \Omega\) such that at least one of the following equalities holds:
\[
n_1^c(\hat{t}, \hat{x}) = k_1 \quad \text{or} \quad n_2^c(\hat{t}, \hat{x}) = k_2.
\]

**Case 1.** \(n_1^c(\hat{t}, \hat{x}) = k_1\).

By definitions of \(\hat{t}\) and \(\hat{x}\), we have \(n_1^c(\hat{t}, \hat{x}) \leq n_1^c(\tilde{t}, x) \forall x \in \Omega\), namely the function \(n_1^c(\hat{t}, x)\) takes minimum for \(x = \hat{x}\). So, if \(\hat{x} \in \Omega\), then \(d_1 \Delta_x n_1^c(\hat{t}, \hat{x}) \geq 0\). If \(\hat{x} \in \partial \Omega\), we can claim again that \(d_1 \Delta_x n_1^c(\hat{t}, \hat{x}) \geq 0\); in fact, if it were \(d_1 \Delta_x n_1^c(\hat{t}, \hat{x}) < 0\) with \(n_1^c(\tilde{t}, \hat{x})\) minimum, it would follow (see references [15,17]) \(\nabla_x n_1^c(\hat{t}, \hat{x}) \cdot \hat{n} < 0\), that would contradict Neumann boundary conditions (1.7).

Moreover, by evaluating the chemical contributions in the first line of (2.12) at \((\tilde{t}, \hat{x})\), we get
\[
-2(n_1^c)^2(\tilde{t}, \hat{x}) + 4 \alpha n_1^c(\tilde{t}, \hat{x}) n_2^c(\tilde{t}, \hat{x}) + 2 \beta (n_2^c)^2(\tilde{t}, \hat{x}) = -2 k_1^2 + 4 \alpha k_1 n_2^c(\tilde{t}, \hat{x}) + 2 \beta (n_2^c)^2(\tilde{t}, \hat{x})
\]
\[
\geq -2 k_1^2 + 4 \alpha k_1 k_2 + 2 \beta k_2^2 = 0
\]
(the inequality holds since \(n_2^c(\tilde{t}, \hat{x}) \geq k_2\), and the last line vanishes bearing in mind that \(k_1 = (\alpha + \sqrt{\alpha^2 + \beta}) k_2\)).

Consequently, the equation (2.12) for \(n_1^c\) implies that \(\partial_t n_1^c(\hat{t}, \hat{x}) \geq \varepsilon n_1^c(\hat{t}, \hat{x}) > 0\), hence \(n_1^c(t, \hat{x}) < n_1^c(\tilde{t}, \hat{x}) = k_1\) for some \(t < \hat{t}\), contradicting the definition of \(\hat{t}\).

**Case 2.** \(n_2^c(\hat{t}, \hat{x}) = k_2\).

In this case, we have \(n_2^c(\tilde{t}, \hat{x}) \leq n_2^c(\tilde{t}, x) \forall x \in \Omega\), namely \(n_2^c(\tilde{t}, x)\) takes its minimum for \(x = \hat{x}\). Therefore we get, as above, \(d_2 \Delta_x n_2^c(\tilde{t}, \hat{x}) \geq 0\).

As concerns the first term on the right hand side of the second line of (2.12), we obtain
\[
(n_1^c)^2(\tilde{t}, \hat{x}) - 2 \alpha n_1^c(\tilde{t}, \hat{x}) n_2^c(\tilde{t}, \hat{x}) - \beta (n_2^c)^2(\tilde{t}, \hat{x}) = (n_1^c(\tilde{t}, \hat{x}) - \gamma k_2) (n_1^c(\tilde{t}, \hat{x}) + \delta k_2)
\]
\[
\geq (k_1 - \gamma k_2) (n_1^c(\tilde{t}, \hat{x}) + \delta k_2) = 0.
\]
It follows \(\partial_t n_2^c(\tilde{t}, \hat{x}) \geq \varepsilon n_2^c(\tilde{t}, \hat{x}) > 0\), which leads to a contradiction as in Case 1.

Consequently, the set \(B^c\) is unbounded, hence \(n_1(t, x) > k_1\) and \(n_2(t, x) > k_2\) for all \(x \in \Omega\) and for all \(t \geq 0\). This means that \(n_1(t, x) > k_1 e^{-\varepsilon t}\) and \(n_2(t, x) > k_2 e^{-\varepsilon t}\), thus, passing to the limit \(\varepsilon \to 0\), we have \(n_1(t, x) \geq k_1\) and \(n_2(t, x) \geq k_2\).

### 2.3. Convergence to equilibrium

In this subsection we shall derive an explicit rate of convergence towards the equilibrium state \((n_1^*, n_2^*)\) given in (2.5). Precisely, we shall prove:

**Theorem 2.2.** Let \(d_1, d_2 > 0\), \(\alpha, \beta > 0\) and \(\Omega\) be a bounded regular \((C^2)\) open set of \(\mathbb{R}^N\). Let \((n_1(t, x), n_2(t, x))\) be a strong solution (that is, in \(C^2(\mathbb{R}^+ \times \Omega)\)) to system (2.2) with Neumann boundary conditions (1.7) and with initial conditions (2.7). Then, this solution satisfies the following property of exponential decay towards equilibrium with explicit constants:
\[
\frac{1}{4} \|n_1 - n_1^0\|^2 + \frac{\gamma}{2} \|n_2 - n_2^0\|^2 \leq \left( E(n_1^0, n_2^0) - E(n_1^*, n_2^*) \right) e^{-C t},
\] (2.14)
with
\[
C = \min \left\{ \frac{1}{2} \frac{2 + \gamma}{6} d_1, \frac{1}{2} \frac{2 + \gamma}{6} \frac{d_2}{P(\Omega)} + \frac{2 + \gamma}{6} \right\}.
\] (2.15)
where \( P(\Omega) \) is the Poincaré constant of \( \Omega \), and \( d_3 = k_1 + \delta k_2 = 2 \sqrt{\alpha^2 + \beta k_2} \) is the lower bound for \( n_1 + \delta n_2 \) (remember that \( \gamma \) and \( \delta \) are defined by (2.4), and that \( E \) is defined by (2.6)).

The proof of this theorem is based on the following functional inequality:

**Lemma 2.3 (entropy dissipation).** Let \( n_1 := n_1(x) \) and \( n_2 := n_2(x) \) be two nonnegative functions of \( L^1(\Omega) \) such that

\[
\int_\Omega (n_1(x) + 2n_2(x)) \, dx = n_0, \quad \text{and} \quad \forall x \in \Omega, \quad n_1(x) + \delta n_2(x) \geq d_3.
\]

Then, the entropy dissipation

\[
D(n_1, n_2) = \frac{d_1}{2} \int_\Omega |\nabla_n n_1|^2 \, dx + d_2 \gamma \int_\Omega |\nabla_n n_2|^2 \, dx + \int_\Omega (n_1 - \gamma n_2) ((n_1)^2 - 2\alpha n_1 n_2 - \beta (n_2)^2) \, dx
\]

fulfils the inequality

\[
D(n_1, n_2) \geq C \left[ E(n_1, n_2) - E(n_1^*, n_2^*) \right],
\]

where the constant \( C \) is defined in (2.15).

**Proof of Lemma 2.3.** For convenience, the proof will be divided into five steps.

**Step 1.** A direct computation shows that the relative entropy with respect to the equilibrium state \((n_1^*, n_2^*)\) is related to the \( L^2 \)-distance from the equilibrium itself:

\[
E(n_1, n_2) - E(n_1^*, n_2^*) = \frac{1}{4} \| n_1 - n_1^* \|^2 + \frac{\gamma}{2} \| n_2 - n_2^* \|^2.
\]

This ensures that the entropy \( E(n_1, n_2) \) takes its minimum for \((n_1, n_2) = (n_1^*, n_2^*)\).

**Step 2.** By resorting to Poincaré inequality, we have

\[
\int_\Omega |\nabla_n n_1|^2 \, dx \geq \frac{1}{P(\Omega)} \| n_1 - \bar{n}_1 \|^2 \quad \text{where} \quad \bar{n}_1 = \int_\Omega n_1 \, dx.
\]

In addition, as concerns the last integral in (2.16) we note that

\[
(n_1 - \gamma n_2) ((n_1)^2 - 2\alpha n_1 n_2 - \beta (n_2)^2) = (n_1 - \gamma n_2)^2 (n_1 + \delta n_2) \geq d_3(n_1 - \gamma n_2)^2.
\]

Hence, the following estimate holds for the entropy dissipation:

\[
D(n_1, n_2) \geq \frac{d_1}{2 P(\Omega)} \| n_1 - \bar{n}_1 \|^2 + \frac{d_2 \gamma}{P(\Omega)} \| n_2 - \bar{n}_2 \|^2 + d_3 \| n_1 - \gamma n_2 \|^2.
\]

**Step 3.** Owing to estimate (2.19) and to the first step, in order to prove Lemma 2.3 it suffices to show that

\[
I := \int_\Omega \left[ \frac{d_1}{2 P(\Omega)} \| n_1 - \bar{n}_1 \|^2 + \frac{d_2 \gamma}{P(\Omega)} \| n_2 - \bar{n}_2 \|^2 + d_3 \| n_1 - \gamma n_2 \|^2 \right] \, dx
\]

\[
\geq C \int_\Omega \left[ \frac{1}{4} \| n_1 - n_1^* \|^2 + \frac{\gamma}{2} \| n_2 - n_2^* \|^2 \right] \, dx.
\]
Step 4. Using the inequality
\[ |n_i - n_i^*|^2 \leq 2 \left[ |n_i - \bar{n_i}|^2 + |\bar{n_i} - n_i^*|^2 \right], \]
we see that in order to get estimate (2.20), it is enough to prove that
\[ I \geq C \int_{\Omega} \left[ \frac{1}{2} |n_1 - \bar{n_1}|^2 + \gamma |n_2 - \bar{n_2}|^2 + \frac{1}{2} |\bar{n_1} - n_1^*|^2 + \gamma |\bar{n_2} - n_2^*|^2 \right] \, dx. \] (2.21)

Since it obviously holds
\[ \frac{1}{2} I \geq C_1 \int_{\Omega} \left[ \frac{1}{2} |n_1 - \bar{n_1}|^2 + \gamma |n_2 - \bar{n_2}|^2 \right] \, dx, \]
with
\[ C_1 = \min \left\{ \frac{d_1}{2 \gamma P(\Omega)}, \frac{d_2}{\gamma P(\Omega)} \right\}, \] (2.22)

it remains to prove that
\[ \frac{1}{2} I \geq C_2 \left[ \frac{1}{2} |\bar{n_1} - n_1^*|^2 + \gamma |\bar{n_2} - n_2^*|^2 \right], \] (2.23)
and to take \( C = \min\{C_1, C_2\} \).

Step 5. Since
\[ |\bar{n_1} - \gamma \bar{n_2}|^2 \leq 3 \left[ |\bar{n_1} - n_1|^2 + |n_1 - n_2|^2 + \gamma^2 |n_2 - \bar{n_2}|^2 \right], \]
we have
\[ \frac{1}{3} \min \left\{ \frac{d_1}{2 \gamma P(\Omega)} , \frac{d_2}{\gamma P(\Omega)} , d_3 \right\} |\bar{n_1} - \gamma \bar{n_2}|^2 \leq I, \]

therefore in order to prove (2.23) it suffices to show that
\[ |\bar{n_1} - \gamma \bar{n_2}|^2 \geq \frac{6}{\min \left\{ \frac{d_1}{2 \gamma P(\Omega)} , \frac{d_2}{\gamma P(\Omega)} , d_3 \right\}} C_2 \left[ \frac{1}{2} |\bar{n_1} - n_1^*|^2 + \gamma |\bar{n_2} - n_2^*|^2 \right]. \] (2.24)

At this point, bearing in mind the expressions of \((n_1^*, n_2^*)\) given in (2.5), together with the fact that \(n_1^* + 2 n_2^* = \bar{n_1} + 2 \bar{n_2} = \bar{n}^3\), we get
\[ \bar{n_2} - n_2^* = -\frac{1}{2} (\bar{n_1} - n_1^*) \quad \text{and} \quad \bar{n_1} - \gamma \bar{n_2} = \frac{2 + \gamma}{2} (\bar{n_1} - n_1^*), \]
so that (2.24) becomes
\[ |\bar{n_1} - n_1^*|^2 \geq \frac{6}{2 + \gamma} \min \left\{ \frac{d_1}{2 \gamma P(\Omega)} , \frac{d_2}{\gamma P(\Omega)} , d_3 \right\} C_2 |\bar{n_1} - n_1^*|^2, \]

that is true once we put
\[ C_2 = \frac{2 + \gamma}{6} \min \left\{ \frac{d_1}{2 \gamma P(\Omega)} , \frac{d_2}{\gamma P(\Omega)} , d_3 \right\}. \] (2.25)

Taking \( C = \min\{C_1, C_2\} \) concludes the proof of Lemma 2.3. \( \square \)
We are now in condition to conclude the Proof of Theorem 2.2. Let’s evaluate the entropy dissipation along the solution of system (2.2):

\[- \partial_t E(n_1, n_2) = - \int_\Omega \left( \frac{n_1}{2} \partial_t n_1 + \gamma n_2 \partial_t n_2 \right) \, dx.\]

Taking into account that for Neumann boundary conditions

\[- \int_\Omega n_i \Delta_x n_i \, dx = - \int_\Omega |\nabla_x n_i|^2 \, dx,\]

we have

\[- \partial_t E(n_1, n_2) = D(n_1, n_2).\]

Owing to Lemma 2.3, we end up with

\[\partial_t \left[ E(n_1, n_2) - E(n_1^*, n_2^*) \right] \leq - C \left[ E(n_1, n_2) - E(n_1^*, n_2^*) \right],\]

thus, by Gronwall’s inequality,

\[E(n_1, n_2) - E(n_1^*, n_2^*) \leq \left( E(n_1^0, n_2^0) - E(n_1^*, n_2^*) \right) e^{-Ct}.\]

Finally, the first step of Lemma 2.3 provides the sought estimate (2.14). \(\square\)

**Remark 2.4.** The same procedure could be applied to whatever bi-species reaction-diffusion system in which chemical reaction contributions take the form

\[Q_2 = C (n_1 - \gamma n_2) W(n_1, n_2),\]

with \(C\) and \(\gamma\) positive constants and \(W(n_1, n_2)\) (smooth) function satisfying the estimate

\[W(n_1, n_2) \geq A n_1 + B n_2\]

(where \(A\) and \(B\) are positive constants).

### 3. Mathematical study in the general case

In this section, we prove Theorem 1.1. In next subsection, we begin by giving a few results about the reaction terms \(Q_i\).

#### 3.1. Study of the reaction terms

System (1.1)–(1.2)–(1.5) writes

\[\partial_t n_i - d_i \Delta_x n_i = Q_i(n_1, n_2)\quad i = 1, 2,\]

(3.1)

where the collision contributions may be rearranged as

\[Q_1 = - 2 Q_2, \quad Q_2 = \frac{1}{\nu_{30} n_0 + \nu_{31} n_1 + \nu_{32} n_2} F(n_1, n_2),\]

(3.2)
we get

\[ F(n_1, n_2) = \nu_{11}^2 (n_1) (n_1) (n_2) (n_2) \]

This function is homogeneous (of third order) in the variables \( n_1, n_2, n_0 \), and it can be seen as a cubic function of the variable \( n_1 \):

\[ F(n_1, n_2) = A(n_1)^3 + B(n_1)^2 - C n_1 - D, \]

where

\[ A = \nu_{11}^2 \nu_{31}^2 > 0, \]
\[ B = (\nu_{11}^2 \nu_{32}^2 - \nu_{31}^2 \nu_{21}^2) n_2 + \nu_{11}^2 \nu_{30}^2 n_0, \]
\[ C = (\nu_{22}^2 \nu_{31}^2 + \nu_{32}^2 \nu_{21}^2) (n_2)^2 + (\nu_{21}^2 \nu_{30}^2 + \nu_{31}^2 \nu_{20}^2) n_2 n_0 > 0, \]
\[ D = \nu_{22}^2 \nu_{32}^2 (n_2)^3 + (\nu_{22}^2 \nu_{32}^2 + \nu_{22}^2 \nu_{30}^2) (n_2)^2 n_0 + \nu_{20}^2 \nu_{30}^2 (n_2)(n_0)^2 > 0. \]

So, using the sign of \( A, B, D \), it can be checked that there exists only one admissible (i.e., strictly positive) root \( n_1 = G(n_2, n_0) \) of \( F(n_1, n_2) = 0 \), with \( G \) homogeneous of order 1 in the variables \( n_2, n_0 \), and that

\[ \frac{\partial F}{\partial n_1}(G(n_2, n_0), n_2) > 0. \]

Therefore, the collision contributions may be rewritten as \( Q_1 = -2 Q_2 \) and

\[ Q_2 = \frac{1}{\nu_{30}^2 n_0 + \nu_{31}^2 n_1 + \nu_{32}^2 n_2} \left[ n_1 - G(n_2, n_0) \right] P(n_1, n_2, n_0), \tag{3.4} \]

where \( P(n_1, n_2, n_0) \) is an homogeneous function of order 2 that is strictly positive (for \( n_1 > 0, n_2 > 0, n_0 > 0 \)).

**Lemma 3.1.** With the notations above, \( \frac{\partial G}{\partial n_2}(n_2) > 0 \) for each \( n_2 > 0 \) (we skip here the dependence of \( G \) on the background fixed density \( n_0 \) to keep reasonable notations).

**Proof of Lemma 3.1.** Differentiating the equality

\[ F(G(n_2), n_2) = 0, \tag{3.5} \]

we get

\[ \frac{\partial G}{\partial n_2}(n_2) = -\frac{\partial F}{\partial n_2}(G(n_2), n_2). \tag{3.6} \]

By resorting to (3.5), identity (3.6) is equivalent to

\[ \frac{\partial G}{\partial n_2}(n_2) = -\frac{\partial F}{\partial n_2}(G(n_2), n_2) - \frac{1}{n_2} F(G(n_2), n_2) - 2 \frac{\partial F}{\partial n_1}(G(n_2), n_2) =: -\frac{N}{D}. \tag{3.7} \]

The numerator of this fraction turns out to be

\[ N = -2 \nu_{22}^2 \nu_{32}^2 (n_2)^2 - \nu_{11}^2 \nu_{31}^2 \frac{G^2(n_2)}{n_2} - (\nu_{22}^2 \nu_{31}^2 + \nu_{32}^2 \nu_{21}^2) G(n_2) n_2 \]

\[ - (\nu_{20}^2 \nu_{32}^2 + \nu_{22}^2 \nu_{30}^2) n_0 n_2 - \nu_{11}^2 \nu_{30}^2 n_0 \frac{G^2(n_2)}{n_2} < 0. \tag{3.8} \]
As concerns the denominator of fraction (3.7), we obtain analogously
\[ D = \nu_1^d \nu_3^d G^2(n_2) + 2 \nu_2^d \nu_3^d \frac{(n_2)^3}{G(n_2)} + (\nu_2^d \nu_3^d + \nu_3^d \nu_2^d)(n_2)^2 \]
\[ + (\nu_1^d \nu_3^d + \nu_2^d \nu_3^d)n_0 n_2 + 2(\nu_2^d \nu_3^d + \nu_3^d \nu_2^d)n_0 \frac{(n_2)^2}{G(n_2)} + 2 \nu_2^d \nu_3^d (n_0)^2 \frac{n_2}{G(n_2)} > 0. \]

By inserting results (3.8) and (3.9) into equality (3.7), we get
\[ \frac{\partial G}{\partial n_2}(n_2) > 0. \]  

Lemma 3.2. If \( n_1 \) and \( n_2 \) are bounded from above and from below (that is, \( k_1 \leq n_1 \leq K_1, \quad k_2 \leq n_2 \leq K_2, \) \( k_1, k_2, K_1, K_2 > 0 \), then there exist positive constants \( g, G, \tilde{g}, \tilde{G}, p, P \) such that
\[ g \leq G(n_2, n_0) \leq G, \]
\[ \tilde{g} \leq \frac{\partial G}{\partial n_2}(n_2, n_0) \leq \tilde{G}, \]
\[ p \leq P(n_1, n_2, n_0) \leq P. \]

Proof of (3.12) of Lemma 3.2. Since \( k_2 \leq n_2 \leq K_2 \) and \( G \) is increasing with respect to \( n_2 \) (we have proved that \( \frac{\partial G}{\partial n_2}(n_2, n_0) > 0 \)), we immediately get
\[ g = G(k_2, n_0) \leq G(n_2, n_0) \leq G(K_2, n_0) = G. \]

Proof of (3.13) of Lemma 3.2. Since \( n_2 \) and \( G(n_2, n_0) \) are bounded from above and from below, we can obtain lower and upper bounds for expressions (3.8) and (3.9), thus, coming back to equality (3.7), also for \( \frac{\partial G}{\partial n_2}(n_2, n_0) \). More precisely, \( c_N < -N < C_N \) where
\[ c_N = 2 \nu_2^d \nu_3^d (K_2)^2 + \nu_1^d \nu_3^d \frac{G^2(K_2)}{K_2} + (\nu_2^d \nu_3^d + \nu_3^d \nu_2^d) G(K_2) k_2 \]
\[ + (\nu_1^d \nu_3^d + \nu_2^d \nu_3^d)n_0 k_2 + \nu_1^d \nu_3^d n_0 \frac{G^2(K_2)}{K_2}, \]
\[ C_N = 2 \nu_2^d \nu_3^d (K_2)^2 + \nu_1^d \nu_3^d \frac{G^2(K_2)}{K_2} + (\nu_2^d \nu_3^d + \nu_3^d \nu_2^d) G(K_2) K_2 \]
\[ + (\nu_1^d \nu_3^d + \nu_2^d \nu_3^d)n_0 K_2 + \nu_1^d \nu_3^d n_0 \frac{G^2(K_2)}{K_2}, \]
and $c_D < D < C_D$ where

$$c_D = \nu_{11} \nu_{31} G^2(k_2) + 2 \nu_{22} \nu_{31}^d (k_2)^3 + \left( \nu_{22} \nu_{31} + \nu_{32} \nu_{21}^d \right) (k_2)^2$$

$$+ \left( \nu_{21} \nu_{30} + \nu_{20} \nu_{31}^d \right) n_0 k_2 + 2 \left( \nu_{20} \nu_{32} + \nu_{22} \nu_{30}^d \right) n_0 \frac{(k_2)^2}{G(k_2)} + 2 \nu_{20} \nu_{30} (n_0)^2 \frac{k_2}{G(k_2)}$$

$$C_D = \nu_{11} \nu_{31} G^2(K_2) + 2 \nu_{22} \nu_{32}^d (K_2)^3 + \left( \nu_{22} \nu_{31} + \nu_{32} \nu_{21}^d \right) (K_2)^2$$

$$+ \left( \nu_{21} \nu_{30} + \nu_{20} \nu_{31}^d \right) n_0 K_2 + 2 \left( \nu_{20} \nu_{32} + \nu_{22} \nu_{30}^d \right) n_0 \frac{(K_2)^2}{G(k_2)} + 2 \nu_{20} \nu_{30} (n_0)^2 \frac{K_2}{G(k_2)}$$

so that

$$\tilde{g} = \frac{c_N}{c_D}, \quad \tilde{G} = \frac{C_N}{c_D} \quad \square$$

**Proof of (3.14) of Lemma 3.2.** Let us recall that

$$\left[ n_1 - G(n_2, n_0) \right] \mathcal{P}(n_1, n_2, n_0) = \mathcal{F}(n_1, n_2, n_0)$$

given in (3.5), hence $\mathcal{P}$ may be written in the form

$$\mathcal{P}(n_1, n_2, n_0) = \Lambda(n_1)^2 + \Upsilon(n_2, n_0) n_1 + \Theta(n_2, n_0), \quad (3.15)$$

with $\Upsilon$ homogeneous of order 1 and $\Theta$ homogeneous of order 2. Let us compare the following “polynomial function”

$$\left[ n_1 - G(n_2, n_0) \right] \mathcal{P}(n_1, n_2, n_0) = \Lambda(n_1)^3 + \left[ \Upsilon(n_2, n_0) - \Lambda G(n_2, n_0) \right] (n_1)^2$$

$$+ \left[ \Theta(n_2, n_0) - \Upsilon(n_2, n_0) G(n_2, n_0) \right] n_1 - \Theta(n_2, n_0) G(n_2, n_0)$$

with the function $\mathcal{F}(n_1, n_2, n_0)$ (see (3.5)). Looking at the coefficient of $(n_1)^3$, we immediately get $\Lambda = \nu_{11} \nu_{31}$. Then, by considering the terms of order 0 in $n_1$, we have

$$\Theta(n_2, n_0) G(n_2, n_0) = \nu_{22} \nu_{32}^d (n_2)^3 + \left( \nu_{20} \nu_{32}^d + \nu_{22} \nu_{30} n_2 \right) (n_2)^2 n_0 + \nu_{20} \nu_{30} n_2 (n_0)^2,$$

so there exist positive constants $\theta_1$, $\theta_2$ such that $\theta_1 \leq \Theta(n_2, n_0) \leq \theta_2$; precisely

$$\theta_1 = \frac{1}{G(k_2, n_0)} \left[ \nu_{22} \nu_{32}^d (k_2)^3 + \left( \nu_{20} \nu_{32}^d + \nu_{22} \nu_{30}^d \right) (k_2)^2 n_0 + \nu_{20} \nu_{30}^d k_2 (n_0)^2 \right],$$

$$\theta_2 = \frac{1}{G(k_2, n_0)} \left[ \nu_{22} \nu_{32}^d (K_2)^3 + \left( \nu_{20} \nu_{32}^d + \nu_{22} \nu_{30}^d \right) (K_2)^2 n_0 + \nu_{20} \nu_{30}^d K_2 (n_0)^2 \right].$$

Finally, by comparing the coefficients of $n_1$, we get

$$\Upsilon(n_2, n_0) G(n_2, n_0) = \Theta(n_2, n_0) + \left( \nu_{22} \nu_{32}^d + \nu_{22} \nu_{21} n_2 \right) (n_2)^2 + \left( \nu_{21} \nu_{30} + \nu_{20} \nu_{31} \right) n_2 n_0,$$
thus there exist positive constants $y_1, y_2$ such that $y_1 \leq \Upsilon(n_2, n_0) \leq y_2$:

$$y_1 = \frac{1}{\mathcal{G}(K_2, n_0)} \left[ \theta_1 + (\nu^d_{22} \nu^d_{31} + \nu^d_{32} \nu^d_{21})(k_2)^2 + (\nu^d_{21} \nu^d_{30} + \nu^d_{20} \nu^d_{31})k_2 n_0 \right],$$

$$y_2 = \frac{1}{\mathcal{G}(k_2, n_0)} \left[ \theta_2 + (\nu^d_{22} \nu^d_{31} + \nu^d_{32} \nu^d_{21})(\mathcal{K}_2)^2 + (\nu^d_{21} \nu^d_{30} + \nu^d_{20} \nu^d_{31})\mathcal{K}_2 n_0 \right].$$

By inserting these results, together with the bounds of $n_1$, into (3.15), we get the sought lower and upper bounds for $P(n_1, n_2, n_0)$:

$$p = \Lambda(k_1)^2 + y_1 \theta_1, \quad P = \Lambda(k_1)^2 + y_2 K_1 + \theta_2.$$  \hfill \Box

### 3.2. Existence and uniqueness of a strong solution

**Proposition 3.3.** Let $d_1, d_2 > 0$ and $\Omega$ be a bounded regular $(C^2)$ open set of $\mathbb{R}^N$. We consider initial data in $C^2(\Omega)$, compatible with Neumann boundary conditions, and satisfying the bounds (for some strictly positive constants $c_1, c_2, C_1$ and $C_2$):

\begin{align*}
0 < c_1 < n_1(0, x) < C_1, \quad &0 < c_2 < n_2(0, x) < C_2. 
\end{align*}

Then, for each $T > 0$, there exists a unique (strong) solution $n_1(t, x), n_2(t, x)$ in $C^2([0, T] \times \Omega)$ to system (3.1)–(1.7) such that, for $(t, x) \in [0, T] \times \Omega$,

\begin{align*}
k_1 \leq n_1(t, x) \leq K_1, \quad &k_2 \leq n_2(t, x) \leq K_2,
\end{align*}

where

\begin{align*}
k_1 &= \min \left\{ c_1, \mathcal{G}(c_2, n_0) \right\}, \quad &k_2 = \mathcal{G}^{-1}(k_1, n_0) = \min \left\{ \mathcal{G}^{-1}(c_1, n_0), c_2 \right\}, \\
K_1 &= \max \left\{ C_1, \mathcal{G}(C_2, n_0) \right\}, \quad &K_2 = \mathcal{G}^{-1}(K_2, n_0) = \max \left\{ \mathcal{G}^{-1}(C_1, n_0), C_2 \right\}. 
\end{align*}

**Remark 3.4.** Given a fixed positive value $n_0$, we have proved that the function $n_2 \mapsto \mathcal{G}(n_2, n_0)$ is strictly increasing, and moreover it can be checked that $\mathcal{G}(0, n_0) = 0$ and $\lim_{n_2 \to +\infty} \mathcal{G}(n_2, n_0) = +\infty$; consequently, the function $\mathcal{G}^{-1}$ is well defined.

**Proof of Proposition 3.3.** Initial bounds (3.16) imply of course

\begin{align*}
k_1 < n_1(0, x) < K_1, \quad &k_2 < n_2(0, x) < K_2.
\end{align*}

At first let us prove that the “maximum principle” holds for $(t, x) \in [0, T] \times \Omega$.

**Lemma 3.5.** Let $\varepsilon > 0$ and $\Omega$ be a bounded regular $(C^2)$ open set of $\mathbb{R}^N$. For any $T > 0$, there exists a strong solution $(N_1^\varepsilon(t, x), N_2^\varepsilon(t, x))$ (in $C^2([0, T] \times \Omega)$) to system

\begin{align*}
\partial_t N_1^\varepsilon - d_1 \Delta_x N_1^\varepsilon &= -\frac{2\mathcal{P}(N_1^\varepsilon, N_2^\varepsilon, n_0 \varepsilon^t)}{\nu^d_{30}n_0 \varepsilon^t + \nu^d_{31}N_1^\varepsilon + \nu^d_{32}N_2^\varepsilon} \left[ N_1^\varepsilon - \mathcal{G}(N_2^\varepsilon, n_0 \varepsilon^t) \right], \\
\partial_t N_2^\varepsilon - d_2 \Delta_x N_2^\varepsilon &= \frac{\mathcal{P}(N_1^\varepsilon, N_2^\varepsilon, n_0 \varepsilon^t)}{\nu^d_{30}n_0 \varepsilon^t + \nu^d_{31}N_1^\varepsilon + \nu^d_{32}N_2^\varepsilon} \left[ N_1^\varepsilon - \mathcal{G}(N_2^\varepsilon, n_0 \varepsilon^t) \right]
\end{align*}

with Neumann boundary conditions (1.7) and initial conditions

\begin{align*}
N_1^\varepsilon(0, x) = n_1(0, x), \quad &N_2^\varepsilon(0, x) = n_2(0, x),
\end{align*}
where $n_1(0, x), n_2(0, x)$ satisfy the bounds (3.16). This solution $(N_1^\varepsilon(t,x), N_2^\varepsilon(t,x))$ satisfies the upper bounds:

$$
N_1^\varepsilon(t,x) < K_1 e^{\varepsilon t}, \quad N_2^\varepsilon(t,x) < K_2 e^{\varepsilon t},
$$

where $K_1, K_2$ are the constants defined in (3.19).

Proof of Lemma 3.5. We first suppose to have a strong solution $(N_1^\varepsilon, N_2^\varepsilon)$ (in $C^2([0,T] \times \Omega)$) to system (3.21) with Neumann boundary conditions (1.7) and initial conditions (3.22). Let us consider the functions

$$
\tilde{N}_1^\varepsilon(t,x) = N_1^\varepsilon(t,x) e^{-\varepsilon t}, \quad \tilde{N}_2^\varepsilon(t,x) = N_2^\varepsilon(t,x) e^{-\varepsilon t},
$$

and let us prove that

$$
\tilde{N}_1^\varepsilon(t,x) < K_1, \quad \tilde{N}_2^\varepsilon(t,x) < K_2.
$$

From equations (3.1), it follows that the evolution of $\tilde{N}_1^\varepsilon, \tilde{N}_2^\varepsilon$ is governed by

$$
\begin{align*}
\partial_t \tilde{N}_1^\varepsilon - d_1 \Delta_x \tilde{N}_1^\varepsilon &= -\frac{2\mathcal{P}(\tilde{N}_1^\varepsilon, \tilde{N}_2^\varepsilon, n_0)}{\nu_{30} n_0 + \nu_{31} \tilde{N}_1^\varepsilon + \nu_{32} \tilde{N}_2^\varepsilon} \left[ \tilde{N}_1^\varepsilon - G(\tilde{N}_2^\varepsilon, n_0) \right] - \varepsilon \tilde{N}_1^\varepsilon, \\
\partial_t \tilde{N}_2^\varepsilon - d_2 \Delta_x \tilde{N}_2^\varepsilon &= \frac{\mathcal{P}(\tilde{N}_1^\varepsilon, \tilde{N}_2^\varepsilon, n_0)}{\nu_{30} n_0 + \nu_{31} \tilde{N}_1^\varepsilon + \nu_{32} \tilde{N}_2^\varepsilon} \left[ \tilde{N}_1^\varepsilon - G(\tilde{N}_2^\varepsilon, n_0) \right] - \varepsilon \tilde{N}_2^\varepsilon.
\end{align*}
$$

Suppose that the inequalities (3.25) do not hold for all $(t, x) \in [0,T] \times \Omega$, and define

$$
B^\varepsilon = \left\{ \tau > 0 : \tilde{N}_1^\varepsilon(t,x) < K_1, \quad \tilde{N}_2^\varepsilon(t,x) < K_2 \quad \forall (t,x) \in [0,\tau) \times \Omega \right\}.
$$

If we denote $\bar{t} = \sup B^\varepsilon$, there must exist $\bar{x} \in \Omega$ such that at least one of the following equalities holds:

$$
\tilde{N}_1^\varepsilon(\bar{t}, \bar{x}) = K_1 \quad \text{or} \quad \tilde{N}_2^\varepsilon(\bar{t}, \bar{x}) = K_2.
$$

Case 1. $\tilde{N}_1^\varepsilon(\bar{t}, \bar{x}) = K_1$.

By definitions of $\bar{t}$ and $\bar{x}$, we have $\tilde{N}_1^\varepsilon(\bar{t}, \bar{x}) = N_1^\varepsilon(\bar{t}, \bar{x})$ $\forall x \in \Omega$, namely the function $N_1^\varepsilon(\bar{t}, x)$ takes maximum for $x = \bar{x} \in \Omega$, and consequently $d_1 \Delta_x N_1^\varepsilon(\bar{t}, \bar{x}) \leq 0$ (for more details, see the proof of Prop. 2.1).

Moreover, by evaluating the chemical contribution of equation (3.26) for $N_1^\varepsilon$ at $(\bar{t}, \bar{x})$, we get

$$
- \frac{2\mathcal{P}(\tilde{N}_1^\varepsilon(\bar{t}, \bar{x}), \tilde{N}_2^\varepsilon(\bar{t}, \bar{x}), n_0)}{\nu_{30} n_0 + \nu_{31} N_1^\varepsilon(\bar{t}, \bar{x}) + \nu_{32} N_2^\varepsilon(\bar{t}, \bar{x})} \left[ \tilde{N}_1^\varepsilon(\bar{t}, \bar{x}) - G(\tilde{N}_2^\varepsilon(\bar{t}, \bar{x}), n_0) \right] =
$$

$$
- \frac{2\mathcal{P}(K_1, \tilde{N}_2^\varepsilon(\bar{t}, \bar{x}), n_0)}{\nu_{30} n_0 + \nu_{31} K_1 + \nu_{32} N_2^\varepsilon(\bar{t}, \bar{x})} \left[ K_1 - G(K_2, n_0) \right] \leq
$$

$$
- \frac{2\mathcal{P}(K_1, \tilde{N}_2^\varepsilon(\bar{t}, \bar{x}), n_0)}{\nu_{30} n_0 + \nu_{31} K_1 + \nu_{32} N_2^\varepsilon(\bar{t}, \bar{x})} \left[ K_1 - G(\tilde{N}_2^\varepsilon(\bar{t}, \bar{x}), n_0) \right] = 0,
$$

where the inequality holds since $\tilde{N}_2^\varepsilon(\bar{t}, \bar{x}) \leq K_2$ and $n_2 \rightarrow G(n_2, n_0)$ is increasing. Therefore, from equation (3.21) for $\tilde{N}_1^\varepsilon$, we get that $\partial_t \tilde{N}_1^\varepsilon(\bar{t}, \bar{x}) \leq -\varepsilon \tilde{N}_1^\varepsilon(\bar{t}, \bar{x}) = -\varepsilon K_1 < 0$, hence $\tilde{N}_1^\varepsilon(\bar{t}, \bar{x}) > \tilde{N}_1^\varepsilon(\bar{t}, \bar{x}) = K_1$ for some $t < \bar{t}$, contradicting the definition of $\bar{t}$. 
Case 2. $N_2^x(\tilde{t}, \tilde{x}) = K_2$.

In this case we have $N_2^x(\tilde{t}, \tilde{x}) \geq \tilde{N}_2^x(\tilde{t}, \tilde{x})$ for all $\tilde{x} \in \tilde{\Omega}$, namely $N_2^x(\tilde{t}, \tilde{x})$ takes its maximum for $\tilde{x} = \tilde{x} \in \tilde{\Omega}$, therefore $d_2 \Delta_{x} \tilde{N}_2^x(\tilde{t}, \tilde{x}) \leq 0$. For the chemical contributions at $(\tilde{t}, \tilde{x})$ appearing in the equation for $\tilde{N}_2^x$, we get

$$\frac{\mathcal{P}(\tilde{N}_2^x(\tilde{t}, \tilde{x}), \tilde{N}_2^x(\tilde{t}, \tilde{x}), n_0) e^{-\varepsilon t}}{\nu_{30} n_0 + \nu_{31} N_2^x(\tilde{t}, \tilde{x}) + \nu_{32} \tilde{N}_2^x(\tilde{t}, \tilde{x})} \left[ \tilde{N}_2^x(\tilde{t}, \tilde{x}) - G(\tilde{N}_2^x(\tilde{t}, \tilde{x}), n_0) \right] =$$

$$\frac{\mathcal{P}(\tilde{N}_2^x(\tilde{t}, \tilde{x}), K_2, n_0) e^{-\varepsilon t}}{\nu_{30} n_0 + \nu_{31} N_2^x(\tilde{t}, \tilde{x}) + \nu_{32} K_2} \left[ \tilde{N}_2^x(\tilde{t}, \tilde{x}) - G(K_2, n_0) \right] \leq$$

$$\frac{\mathcal{P}(\tilde{N}_2^x(\tilde{t}, \tilde{x}), K_2, n_0) e^{-\varepsilon t}}{\nu_{30} n_0 + \nu_{31} N_2^x(\tilde{t}, \tilde{x}) + \nu_{32} K_2} \left[ K_1 - G(K_2, n_0) \right] = 0,$$

due to the bound $\tilde{N}_2^x(\tilde{t}, \tilde{x}) \leq -\varepsilon \tilde{N}_2^x(\tilde{t}, \tilde{x}) = -\varepsilon K_2 < 0$, which leads to a contradiction as in Case 1.

Consequently, the set $B^t$ is unbounded, hence $\tilde{N}_2^x(\tilde{t}, \tilde{x}) < K_1$ and $\tilde{N}_2^x(\tilde{t}, \tilde{x}) < K_2$ for all $\tilde{x} \in \tilde{\Omega}$ and for all $t \in [0, T]$. This means that $N_2^x(t, x) < K_1 e^{-\varepsilon t}$ and $N_2^x(t, x) < K_2 e^{-\varepsilon t}$.

In order to prove the existence of a solution (in $C^2([0, T] \times \Omega)$) to system (3.21) with Neumann boundary conditions (1.7) and initial conditions (3.22), let us consider a function $\chi := \chi(N_1^x, N_2^x)$ in $C^2$ such that $\chi(N_1^x, N_2^x) = 1$ when $|N_1^x| \leq e^{\varepsilon t} \max \{K_1, K_2\}$, and $\chi(N_1^x, N_2^x) = 0$ when $|N_1^x| \geq e^{\varepsilon t} \max \{K_1, K_2\}$. Denoting by $R_i$ the right hand side of the $i$-th equation (3.21), the function

$$f(N_1^x, N_2^x) = \left( -2 R_2(N_1^x, N_2^x) \chi(N_1^x, N_2^x), R_2(N_1^x, N_2^x) \chi(N_1^x, N_2^x) \right)$$

lies in $C^2 \cap W^{1,\infty}$. Thus, we can resort to a simple fixed point argument (see for instance [8]) that guarantees that, under these assumptions on the function $f$, there exists a unique strong solution on $[0, T]$ to the problem

$$\begin{cases}
\partial_t U - D \cdot \Delta_x U = f(U) \\
\hat{n} \cdot \nabla_x U = 0 \quad \text{for } x \in \partial \Omega \\
U(0, x) = U_0(x),
\end{cases}$$

(3.27)

where in our case the vector $U \equiv (N_1^x, N_2^x)$, and $D$ stands for the diagonal matrix containing the diffusion coefficients $d_1, d_2$ as diagonal entries.

As long as $\forall x \in \Omega$, $|N_1^x(t, x)|$ and $|N_2^x(t, x)| \leq e^{\varepsilon t} \max \{K_1, K_2\}$, the solution $U(t, x) = (N_1^x(t, x), N_2^x(t, x))$ is also a (strong) solution of system (3.21) (in this range $\chi \equiv 1$, hence $f \equiv (R_1, R_2)$), and satisfies therefore the maximum principle (3.23), so that at the end, $|N_1^x(t, x)|$ and $|N_2^x(t, x)| \leq e^{\varepsilon T} \max \{K_1, K_2\}$ holds up to time $T$.

By using Lemma 3.5, we get a sequence of solutions $\{N_j^x\}_{j>0}$ of system (3.21), which is uniformly bounded in $L^\infty([0, T] \times \tilde{\Omega})$. Therefore, it converges (up to a subsequence) weakly * in $L^\infty$ to some function $n_1$. The same holds for $\{N_j^x\}_{j>0}$.

Boundingness from above of $N_1^x$ and $N_2^x$ implies boundedness (in $[0, T]$) of the polynomial function $[N_1^x - G(N_2^x, n_0 e^{\varepsilon t})]$, appearing in the right hand sides of (3.21). Therefore, using the smoothing properties of the heat equation, we can pass to the limit $\varepsilon \to 0$ in the weak form of equations (3.21), obtaining that the limit functions $(n_1, n_2)$ are weak solutions of the system (3.1). Passing to the limit in (3.23), we get the upper bounds in (3.17).

Since (3.1) is a strict parabolic system, it is clear that $(n_1, n_2)$ lies in $C^2([0, T] \times \tilde{\Omega})$ (cf. [16]), and is therefore the (unique) solution of system (3.1) satisfying the upper bounds in (3.17).
We now turn to the minimum principle, that is the lower bounds in estimate (3.17). We begin by establishing the following result:

**Lemma 3.6.** Let \( \varepsilon > 0 \) and \( \Omega \) be a bounded regular \((C^2)\) open set of \( \mathbb{R}^N \). Let \((N_1^\varepsilon(t,x), N_2^\varepsilon(t,x))\) be a strong solution (that is, in \( C^2([0,T] \times \Omega) \)) to system

\[
\begin{align*}
\partial_t N_1^\varepsilon - d_1 \Delta \varepsilon N_1^\varepsilon &= -\frac{2P(N_1^\varepsilon, N_2^\varepsilon, n_0 e^{-\varepsilon t})}{\nu_{30} n_0 e^{-\varepsilon t} + \nu_{31} N_1^\varepsilon + \nu_{32} N_2^\varepsilon} \left[ N_1^\varepsilon - G(N_2^\varepsilon, n_0 e^{-\varepsilon t}) \right] \\
\partial_t N_2^\varepsilon - d_2 \Delta \varepsilon N_2^\varepsilon &= \frac{P(N_1^\varepsilon, N_2^\varepsilon, n_0 e^{-\varepsilon t})}{\nu_{30} n_0 e^{-\varepsilon t} + \nu_{31} N_1^\varepsilon + \nu_{32} N_2^\varepsilon} \left[ N_1^\varepsilon - G(N_2^\varepsilon, n_0 e^{-\varepsilon t}) \right]
\end{align*}
\]  

with Neumann boundary conditions (1.7) and with initial conditions satisfying the bounds (3.16). This solution \((N_1^\varepsilon(t,x), N_2^\varepsilon(t,x))\) satisfies the lower bounds:

\[
k_1 e^{-\varepsilon t} < N_1^\varepsilon(t,x), \quad k_2 e^{-\varepsilon t} < N_2^\varepsilon(t,x),
\]

where \(k_1, k_2\) are the constants defined in (3.18).

**Proof of Lemma 3.6.** The proof is very similar to the one performed for the maximum principle. We now have to resort to the auxiliary functions

\[
\tilde{N}_1^\varepsilon(t,x) = N_1^\varepsilon(t,x) e^{\varepsilon t}, \quad \tilde{N}_2^\varepsilon(t,x) = N_2^\varepsilon(t,x) e^{\varepsilon t},
\]

and to prove that

\[
k_1 < \tilde{N}_1^\varepsilon(t,x), \quad k_2 < \tilde{N}_2^\varepsilon(t,x).
\]

The evolution of \(\tilde{N}_1^\varepsilon, \tilde{N}_2^\varepsilon\) is now governed by

\[
\begin{align*}
\partial_t \tilde{N}_1^\varepsilon - d_1 \Delta \varepsilon \tilde{N}_1^\varepsilon &= -\frac{2P(\tilde{N}_1^\varepsilon, \tilde{N}_2^\varepsilon, n_0 e^{-\varepsilon t})}{\nu_{30} n_0 e^{-\varepsilon t} + \nu_{31} \tilde{N}_1^\varepsilon + \nu_{32} \tilde{N}_2^\varepsilon} \left[ \tilde{N}_1^\varepsilon - G(\tilde{N}_2^\varepsilon, n_0) \right] + \varepsilon \tilde{N}_1^\varepsilon \\
\partial_t \tilde{N}_2^\varepsilon - d_2 \Delta \varepsilon \tilde{N}_2^\varepsilon &= \frac{P(\tilde{N}_1^\varepsilon, \tilde{N}_2^\varepsilon, n_0 e^{-\varepsilon t})}{\nu_{30} n_0 e^{-\varepsilon t} + \nu_{31} \tilde{N}_1^\varepsilon + \nu_{32} \tilde{N}_2^\varepsilon} \left[ \tilde{N}_1^\varepsilon - G(\tilde{N}_2^\varepsilon, n_0) \right] + \varepsilon \tilde{N}_2^\varepsilon,
\end{align*}
\]

and we conclude like in Lemma 3.5. \(\square\)

Let us now take initial data such that (3.16), and consequently (3.20), hold. Then, there exists a time \(T^* > 0\) depending only on \(K_1, K_2\) (and not on \(\varepsilon\) nor \(n_1^0, n_2^0\)) such that (3.28) (with the initial datum \(N_1^\varepsilon(0,x) = n_1(0,x), N_2^\varepsilon(0,x) = n_2(0,x)\), and the Neumann boundary conditions) admits a strong solution on \([0,T^*]\) satisfying (3.29) and such that

\[
N_1^\varepsilon(t,x) \leq C, \quad N_2^\varepsilon(t,x) \leq C \quad \text{for } t \in [0,T^*], \quad x \in \Omega,
\]

where \(C\) does not depend on \(\varepsilon\) (this can be obtained by a standard fixed point argument).

As a consequence, passing to the limit in (3.29), we obtain that \(n_1(t,x)\) and \(n_2(t,x)\) satisfy the lower bounds in (3.17) on \([0,T]\). Since we can apply this argument at time \(T^*\) (because the data at this time are still bounded from above by \(K_1, K_2\)), we see that the lower bounds in (3.17) hold on \([T^*, 2T^*]\). By induction, they will hold on \([0,T]\), and this concludes the proof of Proposition 3.3. \(\square\)

By sticking together the solutions on \([0,T]\) (for \(T \in \mathbb{R}^+\)), we obtain a (unique) solution in \(C^2(\mathbb{R}^+ \times \Omega)\) to system (3.1)–(1.7). This concludes the proof of existence and uniqueness of a solution stated in Theorem 1.1, since the assumption of strictly positive initial data in \(C^2(\bar{\Omega})\) compatible with the Neumann boundary conditions (introduced in Thm. 1.1) coincides with assumption (3.16).
Entropy functionals and convergence to equilibrium

A crucial feature of the reaction-diffusion system (3.1) is that it admits a unique collision equilibrium \((n^*_1, n^*_2)\), characterized by the relation \(n^*_1 = \mathcal{G}(n^*_2, n_0)\), plus the conservation of number of atoms \(n^*_1 + 2 n^*_2 = n_0\). This allows to build up several different Lyapounov functionals.

Let \(\varphi(\cdot)\) be a strictly increasing function (regular enough). It can be proved that the following functional is suitable for our problem:

\[
E_\varphi = \int_\Omega \left[ \frac{1}{2} \Psi(n_1) + \Phi(n_2) \right] \ dx,
\]

with \(\Psi\) and \(\Phi\) such that

\[
\Psi(n_1) = \varphi(n_1), \quad \frac{\partial \Phi}{\partial n_2}(n_2, n_0) = \varphi(\mathcal{G}(n_2, n_0)).
\]

Let us explain now why \(E_\varphi\) is indeed suitable:

\[\text{Dissipation of the functional}\]

\[
D(n_1, n_2) = - \partial_t E_\varphi(n_1, n_2) = - \int_\Omega \left( \frac{1}{2} \Psi'(n_1) \partial_t n_1 + \frac{\partial \Phi}{\partial n_2}(n_2, n_0) \partial_t n_2 \right) \ dx
\]

\[
= \frac{d_1}{2} \int_\Omega \varphi(n_1) \Delta x n_1 \ dx - d_2 \int_\Omega \varphi(\mathcal{G}(n_2, n_0)) \Delta x n_2 \ dx
\]

\[
- \int_\Omega \left( \frac{1}{2} \varphi(n_1) \mathcal{Q}_1 + \varphi(\mathcal{G}(n_2, n_0)) \mathcal{Q}_2 \right) \ dx
\]

\[
= \frac{d_1}{2} \int_\Omega \varphi'(n_1) \nabla x n_1^2 \ dx + d_2 \int_\Omega \varphi'(\mathcal{G}(n_2, n_0)) \frac{\partial \mathcal{G}}{\partial n_2}(n_2, n_0) |\nabla x n_2|^2 \ dx
\]

\[
+ \int_\Omega \nu_3 n_0 + \nu_3^* n_1 + \nu_3^* n_2 \left[ n_1 - \mathcal{G}(n_2, n_0) \right] \left[ \varphi(n_1) - \varphi(\mathcal{G}(n_2, n_0)) \right] \ dx \geq 0,
\]

since \(\varphi(\cdot)\) is strictly increasing.

\[\text{Strict coercivity of the Lyapounov functional}\]

\[
E_\varphi(n_1, n_2) - E_\varphi(n^*_1, n^*_2) = \int_\Omega \left[ \frac{1}{2} \Psi(n_1) - \frac{1}{2} \Psi(n^*_1) + \Phi(n_2) - \Phi(n^*_2) \right] \ dx
\]

\[
= \int_\Omega \left\{ \frac{1}{2} \left[ \Psi'(n_1)|_{n_1=n_1^*} (n_1 - n_1^*) + \frac{1}{2} \Psi''(\xi)(n_1 - n_1^*)^2 \right]
\]

\[
+ \left[ \frac{\partial \Phi}{\partial n_2}(n_2, n_0) \right]_{n_2=n_2^*} (n_2 - n_2^*) + \frac{1}{2} \frac{\partial^2 \Phi}{\partial n_2^2}(\xi, n_0)(n_2 - n_2^*)^2 \right\} \ dx
\]

\[
= \int_\Omega \left\{ \frac{1}{2} \varphi(n_1^*)(n_1 - n_1^*) + \varphi(\mathcal{G}(n_2^*, n_0))(n_2 - n_2^*)
\]

\[
+ \frac{1}{4} \varphi'(\xi)(n_1 - n_1^*)^2 + \frac{1}{2} \varphi'(\mathcal{G}(\xi, n_0)) \frac{\partial \mathcal{G}}{\partial n_2}(\xi, n_0)(n_2 - n_2^*)^2 \right\} \ dx
\]

\[
= \int_\Omega \left\{ \frac{1}{4} \varphi'(\xi)(n_1 - n_1^*)^2 + \frac{1}{2} \varphi'(\mathcal{G}(\xi, n_0)) \frac{\partial \mathcal{G}}{\partial n_2}(\xi, n_0)(n_2 - n_2^*)^2 \right\} \ dx,
\]
because of conservation of total number of atoms

\[ \int_\Omega (n_1 + 2 n_2) dx = \tilde{n}^0 = \int_\Omega (n_1^* + 2 n_2^*) dx \]

and of the relation \( n_1^* = G(n_2^*, n_0) \). Since \( \varphi(\cdot) \) and \( n_2 \mapsto G(n_2, n_0) \) are strictly increasing functions, the relative entropy is strictly positive for each \((n_1, n_2) \neq (n_1^*, n_2^*)\).

From now on we shall stick to the choice \( \varphi(s) = s \), already adopted in the simplified case dealt with in Section 2, hence to the corresponding “quadratic” entropy functional:

\[ E(n_1, n_2) = \int_\Omega \left( \frac{1}{4} (n_1)^2 + \Phi(n_2, n_0) \right) dx, \]  \hspace{1cm} (3.34)

with

\[ \frac{\partial \Phi}{\partial n_2}(n_2, n_0) = G(n_2, n_0). \]

**Lemma 3.7** (entropy dissipation inequality). Let \( P \) and \( G \) be defined as in Subsection 3.1 from \( Q_1, Q_2 \) defined by (1.5). We suppose that \( n_0, n_1 := n_1(x), n_2 := n_2(x) \geq 0 \) are such that (for some \( d_3 > 0 \))

\[ d_3 \leq \frac{P(n_1, n_2, n_0)}{\nu_{3\Omega} n_0 + \nu_{\beta_1} n_1 + \nu_{\beta_2} n_2}. \]

We also suppose that estimate (3.13) holds.

Then, the entropy dissipation relevant to the quadratic entropy (3.34):

\[ D(n_1, n_2) = \frac{d_1}{2} \int_\Omega |\nabla n_1|^2 dx + d_2 \int_\Omega \frac{\partial G}{\partial n_2}(n_2, n_0) |\nabla n_2|^2 dx \]

\[ + \int_\Omega \frac{P(n_1, n_2, n_0)}{\nu_{3\Omega} n_0 + \nu_{\beta_1} n_1 + \nu_{\beta_2} n_2} [n_1 - G(n_2, n_0)]^2 dx \]  \hspace{1cm} (3.35)

fulfils the inequality:

\[ D(n_1, n_2) \geq C \left[ E(n_1, n_2) - E(n_1^*, n_2^*) \right], \]  \hspace{1cm} (3.36)

with

\[ C = \min \left\{ \min \left\{ 1, \frac{(2 + \tilde{\gamma})^2}{6(2 + G)} \right\}, \frac{d_1}{2 P(\Omega)} \min \left\{ \frac{1}{2}, \frac{(2 + \tilde{\gamma})^2}{6 G(2 + G)} \right\}, \frac{d_2 \tilde{\gamma}}{P(\Omega) G} \frac{(2 + \tilde{\gamma})^2}{6(2 + G)} d_3 \right\}, \]  \hspace{1cm} (3.37)

where \( P(\Omega) \) is the Poincaré constant.

**Proof of Lemma 3.7.** For convenience we divide the proof into four steps.

**Step 1.** Taking into account the bounds (3.13),

\[ D(n_1, n_2) \geq \frac{d_1}{2} \int_\Omega |\nabla n_1|^2 dx + d_2 \tilde{\gamma} \int_\Omega |\nabla n_2|^2 dx + d_3 \int_\Omega [n_1 - G(n_2, n_0)]^2 dx. \]

Then, by resorting to Poincaré inequality,

\[ D(n_1, n_2) \geq \frac{d_1}{2 P(\Omega)} \|n_1 - \bar{n}_1\|_2^2 + \frac{d_2 \tilde{\gamma}}{P(\Omega)} \|n_2 - \bar{n}_2\|_2^2 + d_3 \|n_1 - G(n_2, n_0)\|_2^2, \]  \hspace{1cm} (3.38)
where
\[ \tilde{n}_i(t) = \int_\Omega n_i(t, x) \, dx. \]

**Step 2.** Considering the relative entropy, we have
\[
E(n_1, n_2) - E(n^*_1, n^*_2) = \int_\Omega \left\{ \frac{1}{4} (n_1 - n^*_1)^2 + \frac{1}{2} \frac{\partial G}{\partial n_2} (\xi, n_0) (n_2 - n^*_2)^2 \right\} \, dx
\leq \frac{1}{4} \| n_1 - n^*_1 \|^2 + \frac{\tilde{G}}{2} \| n_2 - n^*_2 \|^2.
\]
Since
\[ |n_i - n^*_i|^2 \leq 2 [ |n_i - \bar{n}_i|^2 + |\bar{n}_i - n^*_i|^2], \]
we see that in order to prove Lemma 3.7, it is enough to show that
\[
J := \int_\Omega \left[ \frac{d_1}{2 P(\Omega)} |n_1 - \tilde{n}_1|^2 + \frac{d_2 \tilde{g}}{P(\Omega) \tilde{G}^2} |n_2 - \tilde{n}_2|^2 + d_3 |n_1 - G(n_2, n_0)|^2 \right] \, dx
\geq C \int_\Omega \left[ \frac{1}{2} |n_1 - \tilde{n}_1|^2 + G |n_2 - \tilde{n}_2|^2 + \frac{1}{2} |\tilde{n}_1 - n^*_1|^2 + \tilde{G} |\tilde{n}_2 - n^*_2|^2 \right] \, dx.
\]
Since it obviously holds
\[ \frac{1}{2} J \geq C_1 \int_\Omega \left[ \frac{1}{2} |n_1 - \tilde{n}_1|^2 + G |n_2 - \tilde{n}_2|^2 \right] \, dx, \]
with
\[
C_1 = \min \left\{ \frac{d_1}{2 P(\Omega)}, \frac{d_2 \tilde{g}}{2 P(\Omega) \tilde{G}^2} \right\}, \tag{3.39}
\]
it remains to prove that
\[ \frac{1}{2} J \geq C_2 \left[ \frac{1}{2} |\tilde{n}_1 - n^*_1|^2 + \tilde{G} |\tilde{n}_2 - n^*_2|^2 \right], \tag{3.40}
\]
and to take \( C = \min \{C_1, C_2\} \).

**Step 3.** Notice that
\[
|\tilde{n}_1 - G(\tilde{n}_2, n_0)|^2 \leq 3 \left[ |\tilde{n}_1 - n_1|^2 + |n_1 - G(n_2, n_0)|^2 + |G(n_2, n_0) - G(\tilde{n}_2, n_0)|^2 \right]
= 3 \left[ |\tilde{n}_1 - n_1|^2 + |n_1 - G(n_2, n_0)|^2 + \left| \frac{\partial G}{\partial n_2} (\xi, n_0) (n_2 - \tilde{n}_2) \right|^2 \right]
\leq 3 \left[ |\tilde{n}_1 - n_1|^2 + |n_1 - G(n_2, n_0)|^2 + \tilde{G}^2 |n_2 - \tilde{n}_2|^2 \right].
\]
Therefore we have
\[
\frac{1}{6} \min \left\{ \frac{d_1}{2 P(\Omega)}, \frac{d_2 \tilde{g}}{P(\Omega) \tilde{G}^2}, d_3 \right\} |\tilde{n}_1 - G(\tilde{n}_2, n_0)|^2 \leq \frac{1}{2} J.
\]
Thus, in order to prove (3.40), it suffices to show that
\[ |\tilde{n}_1 - G(\tilde{n}_2, n_0)|^2 \geq \frac{6}{\min \left\{ \frac{d_1}{2 P(\Omega)}, \frac{d_2 \tilde{g}}{P(\Omega) \tilde{G}^2}, d_3 \right\}} C_2 \left[ \frac{1}{2} |\tilde{n}_1 - n^*_1|^2 + \tilde{G} |\tilde{n}_2 - n^*_2|^2 \right]. \tag{3.41} \]
Step 4. Bearing in mind the preservation of total number of atoms: $\bar{n}_1 + 2\bar{n}_2 = \bar{n}^0 = n_1^* + 2n_2^*$, we have

$$\bar{n}_1 - n_1^* = -2(n_2^* - \bar{n}_2).$$

Moreover,

$$|\bar{n}_1 - G(\bar{n}_2, n_0)|^2 = |\bar{n}_1 - n_1^* + G(n_2^*, n_0) - G(\bar{n}_2, n_0) + \frac{\partial G}{\partial n_2}(\xi, n_0)(n_2^* - \bar{n}_2)|^2$$

$$= |2 + \frac{\partial G}{\partial n_2}(\xi, n_0)|^2 |n_2^* - \bar{n}_2|^2 \geq [2 + \tilde{g}]^2 |n_2^* - \bar{n}_2|^2.$$

Consequently it suffices to prove that

$$[2 + \tilde{g}]^2 |n_2^* - \bar{n}_2|^2 \geq \min\left\{\frac{d_1}{2P(\Omega)}, \frac{d_2 \tilde{g}}{P(\Omega)G^2}, d_3\right\} C_2[2 + \tilde{G}]|\bar{n}_2 - n_2^*|^2.$$  \(\text{(3.42)}\)

This is true once we put

$$C_2 = \frac{1}{6} \min\left\{\frac{d_1}{2P(\Omega)}, \frac{d_2 \tilde{g}}{P(\Omega)G^2}, d_3\right\} \frac{(2 + \tilde{g})^2}{2 + \tilde{G}}.$$

Taking $C = \min\{C_1, C_2\}$ concludes the proof. \(\square\)

Notice that if we had $G(n_2, n_0) = \gamma n_2$, we would obtain $\tilde{g} = \tilde{G} = \gamma$, hence we would correctly reproduce the constant $C$ found in the particular case treated in Section 2.

End of the Proof of Theorem 1.1. Owing to Proposition 3.3, we consider the unique solution $n_1 = n_1(t, x)$, $n_2 = n_2(t, x)$ of equations (1.1)–(1.2)–(1.5) together with (1.7). It satisfies the bound (3.17), so that estimates (3.13) and (3.14) are also satisfied (for a suitable $d_3 > 0$). As a consequence, we can use Lemma 3.7 for $n_1(t, \cdot)$ and $n_2(t, \cdot)$. Bearing in mind that $\partial_t E(n_1, n_2) = -D(n_1, n_2)$, we have

$$\partial_t \left[ E(n_1, n_2) - E(n_1^*, n_2^*) \right] \leq -C \left[ E(n_1, n_2) - E(n_1^*, n_2^*) \right],$$

thus, by Gronwall inequality,

$$E(n_1, n_2) - E(n_1^*, n_2^*) \leq \left( E(n_1^0, n_2^0) - E(n_1^*, n_2^*) \right) e^{-Ct}.$$

Finally, proceeding like in (3.39), we see that

$$E(n_1, n_2) - E(n_1^*, n_2^*) = \int_{\Omega} \left\{ \frac{1}{4} (n_1 - n_1^*)^2 + \frac{1}{2} \frac{\partial G}{\partial n_2}(\xi, n_0)(n_2 - n_2^*)^2 \right\} dx$$

$$\geq \frac{1}{4} \|n_1 - n_1^*\|_2^2 + \frac{\tilde{g}}{2} \|n_2 - n_2^*\|_2^2.$$

This concludes the proof of Theorem 1.1, and defines properly the constants $C_1$ and $C_2$. \(\square\)

Remark 3.8. We end this section by a remark about the case when one diffusivity constant is 0. In this case, the entropy-entropy dissipation estimate (3.36) still holds (and so does the theorem of exponentially fast decay), though with different constants.
This comes out (in the case when $d_1 = 0$) from the following inequality
\[
\int_{\Omega} |n_1 - \bar{n}_1|^2 \, dx \leq 3 \int_{\Omega} |n_1 - \mathcal{G}(n_2, n_0)|^2 \, dx + 3 \int_{\Omega} |\mathcal{G}(n_2, n_0) - \mathcal{G}(n_2, n_0)|^2 \, dx + 3 \int_{\Omega} |\mathcal{G}(n_2, n_0) - \bar{n}_1|^2 \, dx
\]
\[
\leq 6 \int_{\Omega} |n_1 - \mathcal{G}(n_2, n_0)|^2 \, dx + 3 \mathcal{G}^2 P(\Omega) \int_{\Omega} |\nabla n_2|^2 \, dx.
\]

The case $d_2 = 0$ can be treated in the same way.

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