

A REACTION NETWORK APPROACH TO THE CONVERGENCE TO EQUILIBRIUM OF QUANTUM BOLTZMANN EQUATIONS FOR BOSE GASES

GHEORGHE CRACIUN¹ AND MINH-BINH TRAN^{2,*}

Abstract. When the temperature of a trapped Bose gas is below the Bose-Einstein transition temperature and above absolute zero, the gas is composed of two distinct components: the Bose-Einstein condensate and the cloud of thermal excitations. The dynamics of the excitations can be described by quantum Boltzmann models. We establish a connection between quantum Boltzmann models and chemical reaction networks. We prove that the discrete differential equations for these quantum Boltzmann models converge to an equilibrium point. Moreover, this point is unique for all initial conditions that satisfy the same conservation laws. In the proof, we then employ a toric dynamical system approach, similar to the one used to prove the global attractor conjecture, to study the convergence to equilibrium of quantum kinetic equations.

Mathematics Subject Classification. 35Q20, 45A05, 47G10, 82B40, 82B40, 37N25, 92C42, 37C10, 80A30, 92D25.

Received October 20, 2020. Accepted July 7, 2021.

Dedicated to the 60th birthday of Professor Enrique Zuazua

1. INTRODUCTION

Several years after the invention of the Boltzmann–Nordheim equation, which is the quantum version of the classical Boltzmann one, to describe the evolution of dilute quantum gases (*cf.* [40, 51]), a renewal in the kinetic theory of bosons has started by the pioneering work of Kirkpatrick and Dorfman [34, 35]. This work of Kirkpatrick and Dorfman was later extended by Zaremba, Nikuni and Griffin [54], in which the full coupling system of a quantum Boltzmann equation for the density function of the normal fluid/thermal cloud and a Gross–Pitaevskii equation for the wavefunction of the BEC has been introduced. In an independent work, the same model was derived by Pomeau *et al.* [52]. We prefer to [26, 41] for further discussions on the topic. In the models by Zaremba, Nikuni and Griffin and Pomeau, Brachet, Métens and Rica, there are two type of collisional processes.

Keywords and phrases: Quantum Boltzmann equation, dynamical systems, bosons, Bose-Einstein condensate, rate of convergence to equilibrium, global attractor conjecture, mass-action kinetics, power law systems, biochemical networks, Petri net.

¹ Department of Mathematics and Department of Biomolecular Chemistry, University of Wisconsin-Madison, Madison, WI 53706-1388, USA.

² Department of Mathematics, Southern Methodist University, Dallas, Texas 75275, USA.

* Corresponding author: minhbinht@mail.smu.edu

- The $1 \leftrightarrow 2$ interactions between the condensate and the excited atoms, described by the C_{12} collision operator.
- The C_{22} collision operator describes The $2 \leftrightarrow 2$ interactions between the excited atoms themselves, described by the C_{22} collision operator.

A third collisional process, previously missing, was proposed by Reichl and Gust [28, 44]. This process takes into account $1 \leftrightarrow 3$ type collisions between the excitations and is described by the collision operator C_{31} . However, the derivation of the new collision operator C_{31} was very complicated, since it involves the computations of around 40000 individual terms. As a result, a concise mathematical justification for the existence of the missing collision operator C_{31} had been open for many years, and has been solved only until recently in [49].

The spatial homogeneous kinetic equation for the evolution of the density function $f(t, p)$ of the thermal cloud, derived in Section I of [49], takes the form

$$\partial_t f(p) = C_{12}[f](p) + C_{22}[f](p) + C_{31}[f](p), \quad (1.1)$$

in which the forms of C_{12} , C_{22} , C_{31} are given explicitly below

$$\begin{aligned} C_{12}[f](t, p) &= 4\pi \frac{g^2 n}{V} \sum_{p_1, p_2, p_3 \neq 0} (\delta(p - p_1) - \delta(p - p_2) \\ &\quad - \delta(p - p_3)) \\ &\quad \times \delta(\omega(p_1) - \omega(p_2) - \omega(p_3)) (K_{123}^{12})^2 \delta(p_1 - p_2 - p_3) \\ &\quad \times \left[f(p_2) f(p_3) (f(p_1) + 1) - f(p_1) (f(p_2) + 1) (f(p_3) + 1) \right], \end{aligned} \quad (1.2)$$

$$\begin{aligned} C_{22}[f](t, p) &= \frac{g^2 \pi}{V^2} \sum_{p_1, p_2, p_3, p_4 \neq 0} (\delta(p - p_1) + \delta(p - p_2) \\ &\quad - \delta(p - p_3) - \delta(p - p_4)) (K_{1234}^{22})^2 \\ &\quad \times \delta(p_1 + p_2 - p_3 - p_4) \delta(\omega(p_1) + \omega(p_2) - \omega(p_3) - \omega(p_4)) \\ &\quad \times \left[f(p_3) f(p_4) (f(p_2) + 1) (f(p_1) + 1) \right. \\ &\quad \left. - f(p_1) f(p_2) (f(p_3) + 1) (f(p_4) + 1) \right], \end{aligned} \quad (1.3)$$

and

$$\begin{aligned} C_{31}[f](t, p) &= \frac{3g^2 \pi}{V} \sum_{p_1, p_2, p_3, p_4 \neq 0} (\delta(p - p_1) - \delta(p - p_2) \\ &\quad - \delta(p - p_3) - \delta(p - p_4)) \\ &\quad \times (K_{1234}^{31})^2 \delta(p_1 - p_2 - p_3 - p_4) \\ &\quad \times \delta(\omega(p_1) - \omega(p_2) - \omega(p_3) - \omega(p_4)) \\ &\quad \times \left[f(p_3) f(p_4) f(p_2) (f(p_1) + 1) \right. \\ &\quad \left. - f(p_1) (f(p_2) + 1) (f(p_3) + 1) (f(p_4) + 1) \right], \end{aligned} \quad (1.4)$$

in which n is the density of the condensate, $t \in \mathbb{R}_+$ is the time variable, $p \in (\mathbb{Z}/L)^d \setminus \{O\}$ is the d -dimensional non-zero momentum variable, V is proportional to the volume of the periodic box $[-\frac{L}{2}, \frac{L}{2}]^d$, m is the particle

mass, ω is the Bogoliubov dispersion relation defined as

$$\omega_p = \left[\frac{gn}{m} p^2 + \left(\frac{p^2}{2m} \right)^2 \right]^{\frac{1}{2}} \quad (1.5)$$

and g is the interacting constant. We have normalized the Plank constant to be 1. In the above collision operators, the kernels are defined as follows

$$\begin{aligned} K_{1,2,3}^{1,2} &= u_{p_1} u_{p_2} u_{p_3} - v_{p_1} v_{p_2} v_{p_3} - u_{p_1} u_{p_2} v_{p_3} \\ &\quad + v_{p_1} v_{p_2} u_{p_3} - u_{p_1} v_{p_2} u_{p_3} + v_{p_1} u_{p_2} v_{p_3}, \end{aligned} \quad (1.6)$$

$$\begin{aligned} K_{1,2,3,4}^{2,2} &= u_{p_1} u_{p_2} u_{p_3} u_{p_4} + u_{p_1} v_{p_2} u_{p_3} v_{p_4} + u_{p_1} v_{p_2} v_{p_3} u_{p_4} \\ &\quad + v_{p_1} u_{p_2} v_{p_3} u_{p_4} + v_{p_1} u_{p_2} u_{p_3} v_{p_4} + v_{p_1} v_{p_2} v_{p_3} v_{p_4}, \end{aligned} \quad (1.7)$$

and

$$K_{1,2,3,4}^{3,1} = 2 \left[u_{p_1} u_{p_2} v_{p_3} u_{p_4} + v_{p_1} v_{p_2} u_{p_3} v_{p_4} \right], \quad (1.8)$$

with u_p and v_p being defined as

$$u_p, v_p = \left(\frac{\epsilon_p + gn}{2\omega_p} \pm \frac{1}{2} \right)^{\frac{1}{2}}. \quad (1.9)$$

In the setting of [49], we could fix n as a constant, under the assumption that the thermal could fraction is quite small, in comparison to the condensate. Moreover, in the sum on the momenta $\sum_{p \neq 0}$, the origin is removed due to the fact that the condensate has been factored out in the Bogoliubov diagonalization (*cf.* [49, 50]).

Remark 1.1. As it has been discussed in [49], the BEC is in a cubic box with periodic boundary conditions, the quantum Boltzmann equation is then in the discrete form. In order for the conservations of momentum and energy to be satisfied, the following system needs to have solutions on the lattice

$$\begin{aligned} p_1 &= p_2 + p_3 + p_4, & \omega(p_1) &= \omega(p_2) + \omega(p_3) + \omega(p_4), \\ p'_1 + p'_2 &= p'_3 + p'_4, & \omega(p'_1) + \omega(p'_2) &= \omega(p'_3) + \omega(p'_4), \\ p''_1 &= p''_2 + p''_3, & \omega(p''_1) &= \omega(p''_2) + \omega(p''_3). \end{aligned} \quad (1.10)$$

At the first sign, the system does have solutions due to the complicated form of the Bogoliubov dispersion relation (1.5). However, it has been pointed out in [49, 50] that when the temperature of the system is lower but closed to the Bose-Einstein condensation transition temperature, the Bogoliubov dispersion relation can be replaced by the Hattree-Fock energy ($\omega(p) \approx c|p|^2$). In this regime, the two collision operators C_{12} and C_{22} dominate the collisional processes. The contribution of third collision operator C_{31} becomes non-trivial when both u_p and v_p are large, corresponding to significantly low temperatures. In this low temperature regime, the excitations are phonon-like and the Bogoliubov dispersion relation (1.5) can be replaced by the phonon dispersion relation (1.16). The replacement of (1.5) by (1.16) guarantees the existence of solutions to (1.10), and thus, the conservation laws are satisfied.

Simplified Quantum Boltzmann model of the thermal cloud. In our work, we try to provide a deeper understanding of the property of the system derived in [49] by studying a simplified version of it. If we denote

$$f_1 = f(t, p_1), f_2 = f(t, p_2), f_3 = f(t, p_3), f_4 = f(t, p_4),$$

then our simplified system for f_1 writes

$$\frac{\partial f_1}{\partial t} = C_{12}[f_1] + C_{22}[f_1] + C_{13}[f_1], \quad (1.11)$$

where

$$C_{22}[f_1] := \int_{\mathbb{R}^9} K_{p_1, p_2, p_3, p_4}^{22} \delta(p_1 + p_2 - p_3 - p_4) \delta(\mathcal{E}_{p_1} + \mathcal{E}_{p_2} - \mathcal{E}_{p_3} - \mathcal{E}_{p_4}) \quad (1.12)$$

$$\times [(1 + f_1)(1 + f_2)f_3f_4 - f_1f_2(1 + f_3)(1 + f_4)] dp_2 dp_3 dp_4, \quad (1.13)$$

$$C_{12}[f_1] := \int_{\mathbb{R}^6} K_{p_1, p_2, p_3}^{12} \delta(p_1 - p_2 - p_3) \delta(\mathcal{E}_{p_1} - \mathcal{E}_{p_2} - \mathcal{E}_{p_3}) \quad (1.14)$$

$$\times [(1 + f_1)f_2f_3 - f_1(1 + f_2)(1 + f_3)] dp_2 dp_3$$

$$- 2 \int_{\mathbb{R}^6} K_{p_1, p_2, p_3}^{12} \delta(p_2 - p_1 - p_3) \delta(\mathcal{E}_{p_2} - \mathcal{E}_{p_1} - \mathcal{E}_{p_3})$$

$$\times [(1 + f_2)f_1f_3 - f_2(1 + f_1)(1 + f_3)] dp_2 dp_3,$$

and

$$C_{13}[f_1] = \int_{\mathbb{R}^{3 \times 3}} K_{p_1, p_2, p_3, p_4}^{13} \delta(p_1 - p_2 - p_3 - p_4) \delta(\mathcal{E}_{p_1} - \mathcal{E}_{p_2} - \mathcal{E}_{p_3} - \mathcal{E}_{p_4}) \quad (1.15)$$

$$\times [(1 + f_1)f_2f_3f_4 - f_1(1 + f_2)(1 + f_3)(1 + f_4)] dp_2 dp_3 dp_4$$

$$- 3 \int_{\mathbb{R}^{3 \times 3}} K_{p_1, p_2, p_3, p_4}^{13} \delta(p_2 - p_1 - p_3 - p_4) \delta(\mathcal{E}_{p_2} - \mathcal{E}_{p_1} - \mathcal{E}_{p_3} - \mathcal{E}_{p_4})$$

$$\times [(1 + f_2)f_1f_3f_4 - f_2(1 + f_1)(1 + f_3)(1 + f_4)] dp_2 dp_3 dp_4,$$

The quantities $K_{p_1, p_2, p_3, p_4}^{22}, K_{p_1, p_2, p_3}^{12} \geq 0$ are the collision kernels, which are radially symmetric, and symmetric with respect to the permutation of p_1, p_2, p_3 , and p_4 :

$$\begin{aligned} K_{p_1, p_2, p_3, p_4}^{22} &= K_{|p_1|, |p_2|, |p_3|, |p_4|}^{22} = K_{|p_2|, |p_1|, |p_3|, |p_4|}^{22} = K_{|p_3|, |p_2|, |p_1|, |p_4|}^{22} \\ &= K_{|p_4|, |p_2|, |p_3|, |p_1|}^{22} = K_{|p_1|, |p_3|, |p_2|, |p_4|}^{22} = \mathcal{K}_{|p_1|, |p_4|, |p_3|, |p_2|}^{22} = K_{|p_1|, |p_2|, |p_4|, |p_3|}^{22}, \end{aligned}$$

and

$$K_{p_1, p_2, p_3}^{12} = K_{|p_1|, |p_2|, |p_3|}^{12} = K_{|p_2|, |p_1|, |p_3|}^{12} = K_{|p_3|, |p_2|, |p_1|}^{12} = K_{|p_1|, |p_3|, |p_2|}^{12},$$

where $|p|$ denotes the length of the vector p , and $\mathcal{K}_{p_1, p_2, p_3, p_4}^{13}$ is positive, radially symmetric, and symmetric with respect to the permutation of p_1, p_2, p_3, p_4

$$\begin{aligned} K_{p_1, p_2, p_3, p_4}^{13} &= K_{|p_1|, |p_2|, |p_3|, |p_4|}^{13} = K_{|p_2|, |p_1|, |p_3|, |p_4|}^{13} = K_{|p_3|, |p_2|, |p_1|, |p_4|}^{13} \\ &= K_{|p_4|, |p_2|, |p_3|, |p_1|}^{13} = K_{|p_1|, |p_3|, |p_2|, |p_4|}^{13} = \mathcal{K}_{|p_1|, |p_4|, |p_3|, |p_2|}^{13} = K_{|p_1|, |p_2|, |p_4|, |p_3|}^{13}. \end{aligned}$$

We make a further simplification by supposing that the temperature is very low compared to the Bose-Einstein critical temperature. As a result, the energy $\mathcal{E}_p = \mathcal{E}(p)$ is given by the phonon dispersion law (*cf.* [42]):

$$\mathcal{E}(p) = c|p|, \quad c = \sqrt{\frac{gn_c}{m}}. \quad (1.16)$$

Reaction networks and a toric dynamical system approach for the relaxation to equilibrium problem. The study of the relaxation of BECs to thermodynamic equilibrium has played very important role in the theory of Bose gases [25, 28, 29, 43, 54]. Our main tool is to convert these equations into chemical reaction systems and use an extension of the theory of toric dynamical systems (*cf.* [15]).

In general, there is great interest in understanding the qualitative behavior of deterministically modeled chemical reaction systems, including the existence of positive equilibria, stability properties of equilibria, and the non-extinction, or persistence, of species, which are the constituents of these systems [2–4, 8, 13, 15, 21, 22, 24, 31, 53]. Toric dynamical systems – originally called complex-balanced systems (*cf.* [15, 32]) – are models used to describe an important class of chemical kinetics. The complex-balanced condition was first introduced by Boltzmann [11] for modeling collisions in kinetic gas theory. Based on this condition, it was shown by Horn and Jackson [20, 27, 30, 32] that a complex-balanced system has a unique locally stable equilibrium within each linear invariant subspace. To underline the tight connection to the algebraic study of toric varieties, the name “toric dynamical system” was proposed in [15]. The most important problem in the theory of toric dynamical systems is the Global Attractor Conjecture, which says that the complex balanced equilibrium of a toric dynamical system is a globally attracting point within each linear invariant subspace. This global attractor question is strongly related to the convergence to equilibrium problem in the study of kinetic equations. A proof to the Global Attractor Conjecture for small dimensional systems has been supplied in [17], for strongly connected networks in [2], and a complete proof has been proposed in [14].

Our goal is to use the tools developed in [14, 17] to prove the relaxation to equilibrium of Discrete Velocity Models of a model of (1.11), whose collision operator is C_{12} . Similarly, we will prove the relaxation to equilibrium of another model of (1.11), whose collision operator is $C_{12} + C_{22}$, and modified quantum Boltzmann model of the thermal cloud (1.11), whose collision operator is $C_{12} + C_{22} + C_{13}$. A related approach for the study of acoustic wave turbulence has been used in [48]. Let us also mention that some mathematical results of similar kinetic models have been obtained in [1, 5–7, 9, 10, 12, 18, 19, 23, 33, 36–39, 45–47].

The plan of our paper is the following:

- In Section 2, we show that the discrete version of a simplified version of (1.11), that contains only C_{12} , could be rewritten as a chemical reaction network. By using an approach inspired by the theory of toric dynamical system, we prove in Theorem 2.2 that the solution of the discrete version of a simplified version of (1.11), that contains only C_{12} , converges to the equilibrium exponentially in time.
- In Section 3, we generalize Theorem 2.2 to collision operators of the forms C_{13} and C_{22} . We prove that the solutions of the discrete versions of these equations, associated with the collision operators C_{13} and C_{22} converge to equilibria exponentially in Theorems 3.1 and 3.2. In the case of C_{22} , we consider a one-dimensional version of the model.
- In Theorem 4.1 of Section 4, we extend Theorem 3.2 to a simplified version of (1.11), that contains only $C_{12} + C_{22}$, and the modified quantum Boltzmann model of the thermal cloud (1.11), that contains only $C_{12} + C_{22} + C_{31}$.

2. A REACTION NETWORK APPROACH FOR THE CASE OF C_{12}

2.1. The dynamical system associated to C_{12}

As mentioned in the introduction, the model derived from physics to describe the system that couples BEC-excitations at very low temperature is the discrete version of a simplified version of (1.11), that contains only C_{12} , described below.

Let \mathcal{L}_R denote the lattice of integer points

$$\mathcal{L}_R = \{p \in \mathbb{Z}^3, |p| < R\}.$$

The discrete version of the simplified version of (1.11), that contains only C_{12} , reads

$$\begin{aligned} \dot{f}_{p_1} = & \sum_{\substack{p_2, p_3 \in \mathcal{L}_R, \\ p_1 - p_2 - p_3 = 0, \\ \mathcal{E}(p_1) - \mathcal{E}(p_2) - \mathcal{E}(p_3) = 0}} K_{p_1, p_2, p_3}^{12} \{(f_{p_1} + 1)f_{p_2}f_{p_3} - f_{p_1}(f_{p_2} + 1)(f_{p_3} + 1)\} \\ & - 2 \sum_{\substack{p_2, p_3 \in \mathcal{L}_R, \\ p_1 + p_2 - p_3 = 0, \\ \mathcal{E}(p_1) + \mathcal{E}(p_2) - \mathcal{E}(p_3) = 0}} K_{p_1, p_2, p_3}^{12} \{(f_{p_3} + 1)f_{p_1}f_{p_2} - f_{p_3}(f_{p_1} + 1)(f_{p_2} + 1)\}, \end{aligned} \quad (2.1)$$

for all p_1 in \mathcal{L}_R , where $\mathcal{E}(p)$ is defined in (1.16).

2.2. Decoupling the quantum Boltzmann equation associated to C_{12}

Note that when $p_1 = 0$, K_{p_1, p_2, p_3}^{12} is also 0, and therefore, we get

$$\dot{f}_0 = 0, \quad (2.2)$$

which says that $f_0(t)$ is a constant for all time t . Moreover, f_{p_1} does not depend on f_0 for all $p_1 \neq 0$. Therefore, without loss of generality, we can suppose that $f_0(0) = 0$, which leads to $f_0(t) = 0$ for all t .

Taking into account the fact $\mathcal{E}(p) = c|p|$, note that if $p_1, p_2, p_3 \in \mathcal{L}_R$ are different from 0 and $p_3 = p_1 + p_2$ and $|p_3| = |p_1| + |p_2|$ (like in the second sum of (2.1)), then p_1, p_2, p_3 must be collinear and on the same side of the origin. Therefore, we infer that there exists a vector P and $k_1, k_2, k_3 > 0$, $k_1, k_2, k_3 \in \mathbb{Z}$ such that

$$p_1 = k_1 P; \quad p_2 = k_2 P; \quad p_3 = k_3 P, \quad k_1 + k_2 = k_3.$$

Since \mathcal{L}_R is bounded, it follows that k_1, k_2, k_3 belong to a finite set of integer indices $\mathbb{I} = \{1, \dots, I\}$. Arguing similarly for the first sum in (2.1), we deduce that (2.1) is equivalent with the following system for $k_1 \in \mathbb{I}$

$$\begin{aligned} \dot{f}_{P k_1} = & \sum_{\substack{k_2, k_3 \in \mathbb{I}, \\ k_1 - k_2 - k_3 = 0}} K_{P k_1, P k_2, P k_3}^{12} \{(f_{P k_1} + 1)f_{P k_2}f_{P k_3} - f_{P k_1}(f_{P k_2} + 1)(f_{P k_3} + 1)\} \\ & - 2 \sum_{\substack{k_2, k_3 \in \mathbb{I}, \\ k_1 + k_2 - k_3 = 0}} K_{P k_1, P k_2, P k_3}^{12} \{(f_{P k_3} + 1)f_{P k_1}f_{P k_2} - f_{P k_3}(f_{P k_1} + 1)(f_{P k_2} + 1)\}. \end{aligned} \quad (2.3)$$

Note that the system of equations (2.3) shows a *decoupling* of the system of equations (2.1) along a ray $\{kP_0\}$ with $k > 0$ (see Fig. 1). As a consequence, it is sufficient to study the system of equations (2.3) for a fixed value of P_0 , instead of the system of equations (2.1).

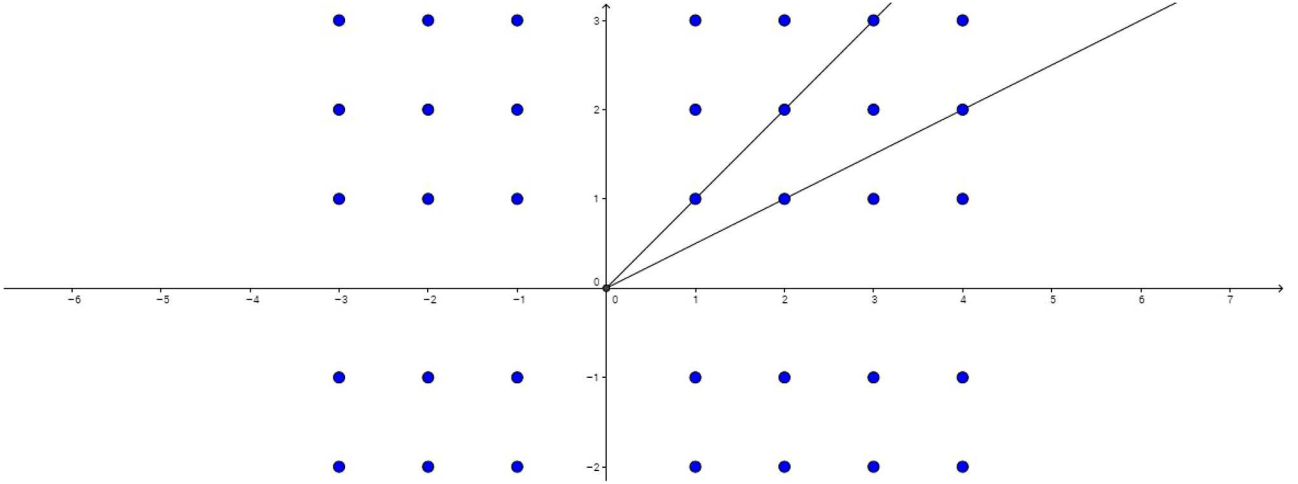


FIGURE 1. We decouple the system (2.1) into rays.

If we denote $f_{k_1 P_0}$ by \bar{f}_{k_1} (with $k_1 \in \mathbb{I}$) and $K_{k_1 P_0, k_2 P_0, k_3 P_0}^{12}$ by $\mathcal{K}_{k_1, k_2, k_3}^{12}$, we obtain the following new system for the ray $\{k_1 P_0 | k_1 > 0\}$:

$$\begin{aligned} \dot{\bar{f}}_{k_1} = & \sum_{\substack{k_2, k_3 \in \mathbb{I}, \\ k_1 = k_2 + k_3}} \mathcal{K}_{k_1, k_2, k_3}^{12} \{(\bar{f}_{k_1} + 1)\bar{f}_{k_2}\bar{f}_{k_3} - \bar{f}_{k_1}(\bar{f}_{k_2} + 1)(\bar{f}_{k_3} + 1)\} \\ & - 2 \sum_{\substack{k_2, k_3 \in \mathbb{I}, \\ k_1 + k_2 = k_3}} \mathcal{K}_{k_1, k_2, k_3}^{12} \{(\bar{f}_{k_3} + 1)\bar{f}_{k_1}\bar{f}_{k_2} - \bar{f}_{k_3}(\bar{f}_{k_1} + 1)(\bar{f}_{k_2} + 1)\}, \quad \forall k_1 \in \mathbb{I}. \end{aligned} \quad (2.4)$$

A simple calculation leads to the following *conservation of energy*

$$\sum_{k=1}^I k \dot{\bar{f}}_k = 0, \quad (2.5)$$

or equivalently

$$\sum_{k=1}^I k \bar{f}_k = \text{const.} \quad (2.6)$$

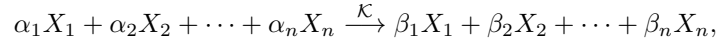
We denote this discrete version of C_{12} by

$$\begin{aligned} \mathcal{C}_{12}[\bar{f}_{k_1}] := & \sum_{k_2 + k_3 = k_1} \mathcal{K}_{k_1, k_2, k_3}^{12} [(\bar{f}_{k_1} + 1)\bar{f}_{k_2}\bar{f}_{k_3} - \bar{f}_{k_1}(\bar{f}_{k_2} + 1)(\bar{f}_{k_3} + 1)] \\ & - 2 \sum_{k_1 + k_3 = k_2} \mathcal{K}_{k_2, k_1, k_3}^{12} [(\bar{f}_{k_2} + 1)\bar{f}_{k_1}\bar{f}_{k_3} - \bar{f}_{k_2}(\bar{f}_{k_1} + 1)(\bar{f}_{k_3} + 1)]. \end{aligned} \quad (2.7)$$

2.3. The chemical reaction network associated to C_{12}

For $x \in \mathbb{R}_{>0}^n$ and $\alpha \in \mathbb{R}_{\geq 0}^n$, we denote by x^α the monomial $\prod_{i=1}^n x_i^{\alpha_i}$.

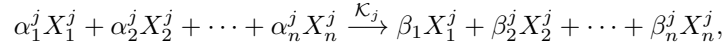
Definition 2.1. Consider a chemical reaction of the form



where \mathcal{K} is a positive parameter, called reaction rate constant. Then the mass-action dynamical system generated by this reaction is

$$\dot{x} = \mathcal{K}x^\alpha(\beta - \alpha), \quad (2.8)$$

where $\alpha = (\alpha_1, \dots, \alpha_n)^T$, $\beta = (\beta_1, \dots, \beta_n)^T$, $\alpha_i, \beta_i \geq 0$ and $x = (x_1, \dots, x_n)^T$, in which x_i is the concentration of the chemical species X_i . For the case of a network that contains several reactions



for $1 \leq j \leq m$, its associated mass-action dynamical system is given by

$$\dot{x} = \sum_{j=1}^m \mathcal{K}_j x^{\alpha^j} (\beta^j - \alpha^j). \quad (2.9)$$

In this section, we will show that the system (2.4) has the form (2.9) for a well-chosen set of reactions.

If $y \rightarrow y'$ and $y' \rightarrow y$ are reactions, we combine them together into a “reversible” reaction $y \leftrightarrow y'$.

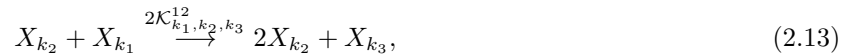
We will derive the system (2.4) from the network of chemical reactions of the form:



for all k_1, k_2, k_3 in \mathbb{I} such that $k_2 + k_3 = k_1$. If we denote by F_k the concentration of the species X_k , we will show that, for appropriate choices of the reaction rate constants in (2.10) and (2.11), the differential equations satisfied by F_k according the mass-action kinetics are exactly the same as (2.4).

In order to describe the connection between the mass-action system given by reactions of the form (2.10)–(2.11) and our system (2.4), we need to consider several cases.

Case 1: For $k_2 + k_3 = k_1$, $k_2 \neq k_3$, $k_1, k_2, k_3 \in \mathbb{I}$, we consider



and for the reversible reaction (2.12) the forward and backward rate constants are the same, *i.e.*, we choose the reaction rate constants of the three reactions $X_{k_2} + X_{k_3} \rightarrow X_{k_1}$, $X_{k_1} \rightarrow X_{k_2} + X_{k_3}$, $X_{k_2} + X_{k_1} \rightarrow 2X_{k_2} + X_{k_3}$ to be $2\mathcal{K}_{k_1, k_2, k_3}^{12}$.

For example, consider the reversible reaction (2.12): in this reaction, X_{k_1} is created from $X_{k_2} + X_{k_3}$ with the rate $2\mathcal{K}_{k_1, k_2, k_3}^{12} F_{k_2} F_{k_3}$ and X_{k_1} is decomposed into $X_{k_2} + X_{k_3}$ with the rate $-2\mathcal{K}_{k_1, k_2, k_3}^{12} F_{k_1}$. Therefore, the rate of change of the species X_{k_1} due to this reaction is $2\mathcal{K}_{k_1, k_2, k_3}^{12} [F_{k_2} F_{k_3} - F_{k_1}]$.

For the irreversible reaction (2.13), X_{k_1} is lost with the rate $-2\mathcal{K}_{k_1, k_2, k_3}^{12} F_{k_2} F_{k_1}$ to create $2X_{k_2} + X_{k_3}$. Therefore the rate of change of the species X_{k_1} due to this reaction is $-2\mathcal{K}_{k_1, k_2, k_3}^{12} F_{k_2} F_{k_1}$. By exchanging the roles of X_{k_2} and X_{k_3} in (2.13), we obtain the rate $-2\mathcal{K}_{k_1, k_2, k_3}^{12} [F_{k_2} F_{k_1} + F_{k_3} F_{k_1}]$.

Therefore, the total rate of change of X_{k_1} due to the reactions in (2.12)–(2.13) is

$$2\mathcal{K}_{k_1,k_2,k_3}^{12}[F_{k_2}F_{k_3} - F_{k_1} - F_{k_2}F_{k_1} - F_{k_3}F_{k_1}]. \quad (2.14)$$

Case 2: For $2k_2 = k_1$, $k_1, k_2 \in \mathbb{I}$, we consider



We choose the reaction rate constant of $2X_{k_2} \rightarrow X_{k_1}$ and the reaction rate constant of $X_{k_1} \rightarrow 2X_{k_2}$ to be $\mathcal{K}_{k_1,k_2,k_3}^{12}$. Also, we choose the reaction rate constant of $X_{k_2} + X_{k_1} \rightarrow 3X_{k_2}$ to be $2\mathcal{K}_{k_1,k_2,k_3}^{12}$.

Consider the first reaction (2.15): In this reaction, X_{k_1} is created from $2X_{k_2}$ with the rate $\mathcal{K}_{k_1,k_2,k_3}^{12}$ and X_{k_1} is decomposed into $2X_{k_2}$ with the rate $-\mathcal{K}_{k_1,k_2,k_3}^{12}F_{k_1}$. The rate of change of the species X_{k_1} is $\mathcal{K}_{k_1,k_2,k_3}^{12}[F_{k_2}^2 - F_{k_1}]$.

For the second reaction (2.16): X_{k_1} is lost with the rate $-2\mathcal{K}_{k_1,k_2,k_3}^{12}F_{k_2}F_{k_1}$ to create $3X_{k_2}$.

As a result, the rate of change of X_{k_1} due to the reactions (2.15)–(2.16) is

$$\mathcal{K}_{k_1,k_2,k_3}^{12}[F_{k_2}^2 - F_{k_1} - 2F_{k_2}F_{k_1}]. \quad (2.17)$$

Case 3: Next, for $k_2 = k_3 + k_1$, $k_1 \neq k_3$, $k_1, k_2, k_3 \in \mathbb{I}$, let us look at the rate of change of X_{k_1} in



For (2.18), the rate of change of X_{k_1} is $2\mathcal{K}_{k_2,k_1,k_3}^{12}[F_{k_2} - F_{k_1}F_{k_3}]$. For (2.19), the rate of change of X_{k_1} is $2\mathcal{K}_{k_2,k_1,k_3}^{12}F_{k_1}F_{k_2}$. By exchanging the roles of X_1 and X_3 , we obtain the rate $2\mathcal{K}_{k_2,k_1,k_3}^{12}[F_{k_1}F_{k_2} + F_{k_2}F_{k_3}]$. Therefore, the rate of change of X_{k_1} due to reactions in (2.18)–(2.20) is

$$-2\mathcal{K}_{k_2,k_1,k_3}^{12}[F_{k_1}F_{k_3} - F_{k_2} - F_{k_2}F_{k_3} - F_{k_1}F_{k_2}]. \quad (2.21)$$

Case 4: Now, for $k_2 = 2k_1$, $k_1, k_2 \in \mathbb{I}$, let us look at the rate of change of X_{k_1} in



For (2.22), the rate of change of X_{k_1} is $2\mathcal{K}_{k_2,k_1,k_3}^{12}[F_{k_2} - F_{k_1}^2]$. For (2.23), the rate of change of X_{k_1} is $4\mathcal{K}_{k_2,k_1,k_3}^{12}F_{k_1}F_{k_2}$. Therefore, the rate of change of X_{k_1} due to the reactions (2.22)–(2.23) is

$$-2\mathcal{K}_{k_2,k_1,k_3}^{12}[F_{k_1}^2 - F_{k_2} - 2F_{k_1}F_{k_2}]. \quad (2.24)$$

From (2.14), (2.17), (2.21), (2.24), the total rate of change of X_{k_1} is

$$\begin{aligned}
& \sum_{k_2+k_3=k_1, k_2 < k_3} 2\mathcal{K}_{k_1, k_2, k_3}^{12} [(F_{k_1} + 1)F_{k_2}F_{k_3} - F_{k_1}(F_{k_2} + 1)(F_{k_3} + 1)] \\
& + \sum_{2k_2=k_1} \mathcal{K}_{k_1, k_2, k_2}^{12} [(F_{k_1} + 1)F_{k_2}F_{k_2} - F_{k_1}(F_{k_2} + 1)(F_{k_2} + 1)] \\
& - \sum_{k_1+k_3=k_2} 2\mathcal{K}_{k_2, k_1, k_3}^{12} [(F_{k_2} + 1)F_{k_1}F_{k_3} - F_{k_2}(F_{k_1} + 1)(F_{k_3} + 1)],
\end{aligned} \tag{2.25}$$

which can be written as

$$\begin{aligned}
\dot{F}_{k_1} &= \sum_{k_2+k_3=k_1} \mathcal{K}_{k_1, k_2, k_3}^{12} [(F_{k_1} + 1)F_{k_2}F_{k_3} - F_{k_1}(F_{k_2} + 1)(F_{k_3} + 1)] \\
& - 2 \sum_{k_1+k_3=k_2} \mathcal{K}_{k_2, k_1, k_3}^{12} [(F_{k_2} + 1)F_{k_1}F_{k_3} - F_{k_2}(F_{k_1} + 1)(F_{k_3} + 1)],
\end{aligned} \tag{2.26}$$

which shows that the system of differential equations satisfied by the concentrations F_k is exactly the same as the system of differential equations (2.4) satisfied by the densities f_k .

2.4. A change of variables

In this section, we introduce a change of variables that will help us to investigate the dynamics of the system (2.26).

Define

$$G_k = \frac{F_k}{F_k + 1},$$

then

$$F_k = \frac{G_k}{1 - G_k},$$

and

$$F_{k_3} + F_{k_1}F_{k_3} + F_{k_2}F_{k_3} - F_{k_1}F_{k_2} = \frac{G_{k_3} - G_{k_1}G_{k_2}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})},$$

$$F_{k_1} + F_{k_1}F_{k_2} + F_{k_1}F_{k_3} - F_{k_3}F_{k_2} = \frac{G_{k_1} - G_{k_2}G_{k_3}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})}.$$

Notice that $0 < F_k < \infty$ and $0 < G_k < 1$.

The system (2.26) is converted into

$$\begin{aligned}
\frac{\dot{G}_{k_1}}{(1 - G_{k_1})^2} &= \tilde{\mathcal{C}}_{12}[G](k_1) := 2 \sum_{k_1+k_2=k_3} \mathcal{K}_{k_1, k_2, k_3}^{12} \frac{G_{k_3} - G_{k_1}G_{k_2}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})} \\
& + \sum_{k_1=k_2+k_3} \mathcal{K}_{k_1, k_2, k_3}^{12} \frac{-G_{k_1} + G_{k_2}G_{k_3}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})}, \forall k_1 \in \mathbb{I}.
\end{aligned} \tag{2.27}$$

Suppose that G represents the column vector $(G_1, \dots, G_I)^T$. Let us also denote by \bar{X}_k , the vector

$$\begin{pmatrix} 0 \\ \dots \\ 1 \\ \dots \\ 0 \end{pmatrix},$$

in which the only element that different from 0 is the k -th one.

Also, for $k_1 \neq k_2$, we denote

$$K_{\bar{X}_{k_1} + \bar{X}_{k_2} \rightarrow \bar{X}_{k_3}}(G) := 2\mathcal{K}_{k_1, k_2, k_3}^{12} \frac{G_{k_1} G_{k_2}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})},$$

$$K_{\bar{X}_{k_3} \rightarrow X_{k_1} + \bar{X}_{k_2}}(G) := 2\mathcal{K}_{k_1, k_2, k_3}^{12} \frac{G_{k_3}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})},$$

$$\mathcal{K}_{\bar{X}_{k_1} + \bar{X}_{k_2} \leftrightarrow \bar{X}_{k_3}} := 2\mathcal{K}_{k_1, k_2, k_3}^{12}.$$

Otherwise, if $k_1 = k_2$, we denote

$$K_{2\bar{X}_{k_1} \rightarrow X_{k_3}}(G) := \mathcal{K}_{k_1, k_1, k_3}^{12} \frac{G_{k_1} G_{k_2}}{(1 - G_{k_1})^2(1 - G_{k_3})},$$

$$K_{\bar{X}_{k_3} \rightarrow 2\bar{X}_{k_1}}(G) := \mathcal{K}_{k_1, k_1, k_3}^{12} \frac{G_{k_3}}{(1 - G_{k_1})^2(1 - G_{k_3})},$$

$$\mathcal{K}_{2\bar{X}_{k_1} \leftrightarrow \bar{X}_{k_3}} := 2\mathcal{K}_{k_1, k_1, k_3}^{12}.$$

Using these notations, the system (2.27) could be rewritten as:

$$\begin{aligned} \dot{G} &= \text{diag} \begin{pmatrix} (1 - G_1)^2 \\ \dots \\ (1 - G_I)^2 \end{pmatrix} \\ &\times \sum_{k_1 + k_2 = k_3} \left[K_{\bar{X}_{k_1} + \bar{X}_{k_2} \rightarrow \bar{X}_{k_3}}(G) - K_{\bar{X}_{k_3} \rightarrow \bar{X}_{k_1} + \bar{X}_{k_2}}(G) \right] (\bar{X}_{k_3} - \bar{X}_{k_1} - \bar{X}_{k_2}). \end{aligned} \quad (2.28)$$

Equivalently, we can also write

$$\dot{G} = \text{diag} \begin{pmatrix} (1 - G_1)^2 \\ \dots \\ (1 - G_I)^2 \end{pmatrix} \sum_{y \leftrightarrow y'} [K_{y \rightarrow y'}(G) - K_{y' \rightarrow y}(G)] (y' - y), \quad (2.29)$$

where $y \leftrightarrow y'$ belongs to the set of reversible reactions

$$\bar{X}_{k_1} + \bar{X}_{k_2} \longleftrightarrow \bar{X}_{k_3}, \quad (2.30)$$

with $k_1 + k_2 = k_3$.

2.5. Convergence to equilibrium

Theorem 2.2. *For any positive initial condition, the solution*

$$f(t) = (f_p(t))_{p \in \mathcal{L}_R}$$

of the discrete quantum Boltzmann equation (2.1) converges to an equilibrium state $f^ = (f_p^*)_{p \in \mathcal{L}_R}$. For each ray $\{kP_0\}_{k \geq 1}$ there exists a positive constant $\rho(P_0)$ such that if $p = kP_0$ then*

$$f_p^* = \frac{1}{e^{k\rho(P_0)} - 1}.$$

Moreover, the solution $f(t)$ of (2.1) converges to f^ exponentially fast in the following sense: there exist positive constants C_1, C_2 such that*

$$\max_{p \in \mathcal{L}_R} |f_p(t) - f_p^*| < C_1 e^{-C_2 t}.$$

Proof. By using the decoupling and the change of variables discussed in the previous sections, for each ray $\{kP_0\}_{k \geq 1}$, we can reduce the study of f to F , which satisfies (2.26). From F , we can switch to study G , which is the solution of (2.29).

Step 1: The Lyapunov function. We recall that (2.27) could be rewritten under the form

$$\dot{G} = \text{diag} \begin{pmatrix} (1 - G_1)^2 \\ \dots \\ (1 - G_I)^2 \end{pmatrix} \sum_{y \leftrightarrow y'} [K_{y \rightarrow y'}(G) - K_{y' \rightarrow y}(G)] (y' - y). \quad (2.31)$$

We define the function

$$L(G) = \sum_{k=1}^I \left(\log(1 - G_k) + \frac{G_k \log G_k}{1 - G_k} - \frac{\log G_k^*}{1 - G_k} \right), \quad (2.32)$$

where $G_k^* = \frac{1}{e^{k\rho}}$, for some $\rho > 0$, and we will show that L is a Lyapunov function for the system (2.27).

We have

$$\nabla L = \begin{pmatrix} \frac{1}{(1-G_1)^2} \log \frac{G_1}{G_1^*} \\ \dots \\ \frac{1}{(1-G_I)^2} \log \frac{G_I}{G_I^*} \end{pmatrix}, \quad (2.33)$$

which implies that

$$\begin{aligned} \text{diag} \begin{pmatrix} (1 - G_1)^2 \\ \dots \\ (1 - G_I)^2 \end{pmatrix} \cdot (y' - y) \cdot \nabla L &= \log \left(\frac{G}{G^*} \right)^{y' - y} \\ &= \log \left(\frac{G}{G^*} \right)^{y'} - \log \left(\frac{G}{G^*} \right)^y. \end{aligned} \quad (2.34)$$

If we define

$$\mathcal{H}_{y,y'}(G) = \frac{K_{y \rightarrow y'}(G)}{\mathcal{K}_{y \leftrightarrow y'} G^y},$$

then $\mathcal{H}_{y,y'} = \mathcal{H}_{y',y}$ for y and y' as in (2.30). Moreover, we have

$$\begin{aligned} & K_{y \rightarrow y'}(G) - K_{y' \rightarrow y}(G) \\ &= \mathcal{K}_{y \leftrightarrow y'} G^y \mathcal{H}_{y,y'}(G) - \mathcal{K}_{y \leftrightarrow y'} G^{y'} \mathcal{H}_{y,y'}(G) \\ &= \mathcal{K}_{y \leftrightarrow y'} \mathcal{H}_{y,y'}(G) [G^y - G^{y'}] \\ &= \mathcal{K}_{y \leftrightarrow y'} (G^*)^y \mathcal{H}_{y,y'}(G) \left[\frac{G^y}{(G^*)^y} - \frac{G^{y'}}{(G^*)^{y'}} \right], \end{aligned} \tag{2.35}$$

since $(G^*)^y = (G^*)^{y'}$.

Combining (2.31), (2.34) and (2.35), we obtain

$$\begin{aligned} & \dot{G} \cdot \nabla L \\ &= \sum_{y \leftrightarrow y'} \left[\log \left(\frac{G}{G^*} \right)^{y'} - \log \left(\frac{G}{G^*} \right)^y \right] \mathcal{K}_{y \leftrightarrow y'} (G^*)^y \mathcal{H}_{y,y'}(G) \left[\frac{G^y}{(G^*)^y} - \frac{G^{y'}}{(G^*)^{y'}} \right] \\ &\leq 0, \end{aligned} \tag{2.36}$$

since log is an increasing function. Also, note that the above inequality is strict unless

$$\frac{G^y}{(G^*)^y} = \frac{G^{y'}}{(G^*)^{y'}}, \tag{2.37}$$

for all reactions $y \leftrightarrow y'$.

Since $(G^*)^y = (G^*)^{y'}$ for all reactions $y \leftrightarrow y'$, this implies $G_{k_1}^* \cdot G_{k_2}^* = G_{k_1+k_2}^*$ for all k_1 and k_2 such that $k_1 + k_2 \leq I$. As a consequence $G_k^* = e^{-\rho k}$, for some positive constant ρ . Moreover, (2.37) implies that at equilibrium $(G)^y = (G)^{y'}$ for all reactions $y \leftrightarrow y'$, which leads to $G_k = e^{-\rho' k}$, for some positive constant ρ' .

By the conservation relation

$$\sum_{k=1}^I k \frac{G_k}{1 - G_k} = \sum_{k=1}^I k \frac{G_k^*}{1 - G_k^*},$$

we deduce that

$$\sum_{k=1}^I k \frac{e^{-\rho k}}{1 - e^{-\rho k}} = \sum_{k=1}^I k \frac{e^{-\rho' k}}{1 - e^{-\rho' k}}.$$

By the monotonicity of the function $\rho \rightarrow \frac{e^{-\rho k}}{1 - e^{-\rho k}}$, we conclude that $\rho = \rho'$, i.e., G^* is the only equilibrium point that satisfies the same conservation relation as the initial condition.

Now, we will prove that there exists exactly one critical point of the Lyapunov function L within each invariant set

$$\mathfrak{S}_c := \left\{ \sum_{k=1}^I k \frac{G_k}{1-G_k} = c \right\}.$$

Since

$$\nabla L = \text{diag} \left(\begin{array}{c} \frac{1}{(1-G_1)^2} \\ \cdots \\ \frac{1}{(1-G_I)^2} \end{array} \right) [\log G - \log G^*],$$

the projection of ∇L on the tangent space to the set \mathfrak{S}_c is 0 if and only if there exists a constant ϱ such that

$$\nabla L = \varrho \cdot \nabla \left(\sum_{k=1}^I k \frac{G_k}{1-G_k} \right),$$

which is equivalent with

$$\text{diag} \left(\begin{array}{c} \frac{1}{(1-G_1)^2} \\ \cdots \\ \frac{1}{(1-G_I)^2} \end{array} \right) [\log G - \log G^*] = \varrho \left(\begin{array}{c} \frac{1}{(1-G_1)^2} \\ \cdots \\ \frac{I}{(1-G_I)^2} \end{array} \right).$$

A direct consequence of the above is the following system of identities

$$\begin{aligned} \log G_1 - \log G_1^* &= \varrho, \\ \log G_2 - \log G_2^* &= 2\varrho, \\ &\cdots \\ \log G_I - \log G_I^* &= I\varrho, \end{aligned}$$

yielding

$$\frac{G_k}{G_k^*} = e^{k\varrho}, \quad \forall k \in \{1, \dots, I\}.$$

Moreover, since G_k and G_k^* satisfy the same conservation law then it follows that $G = G^*$. This implies that G^* is the only critical point of L on the invariant set \mathfrak{S}_c .

Step 2: Differential inclusions and persistence. Now let us observe that (2.4) could be regarded as a \mathcal{K} -variable mass-action system for the reversible network (2.30). For this we write

$$F_{k''} + F_k F_{k''} + F_{k'} F_{k''} = (1 + F_k + F_{k'}) F_{k''},$$

and note that $1 + F_k + F_{k'}$ is bounded below by 1 and above by $1 + 2C$, where

$$C = \sum_{k=1}^I k F_k.$$

Therefore, the results of [14] about persistence of \mathcal{K} -variable reversible mass-action systems can be applied and we conclude that the system is persistent. Alternatively, we can also use the Petri net argument of [4], to prove that the system is persistent, as follows. Note that F_k is the density function of the species X_k . It is straightforward that each *siphon* is $\{X_1, X_2, \dots, X_I\}$, which contains the support of the *P-semiflow* (see [4] for the definition of siphons and P-semiflows) given by

$$\sum_{k=1}^I k F_k = \text{constant}.$$

As a result, the Petri net theory developed in [4] can be applied and it follows that the system is persistent.

Therefore, by using the existence of the globally defined strict Lyapunov function L , and the LaSalle invariance principle, it follows that all trajectories converge to the unique positive equilibrium G^* that we discussed in Step 1.

Step 3: Exponential rate of convergence. Define

$$\begin{aligned} & \mathcal{R}(G) \\ &= \text{diag} \begin{pmatrix} (1 - G_1)^2 \\ \cdots \\ (1 - G_I)^2 \end{pmatrix} \sum_{y \leftrightarrow y'} [K_{y \rightarrow y'}(G) - K_{y' \rightarrow y}(G)] (y' - y) \\ &= \text{diag} \begin{pmatrix} (1 - G_1)^2 \\ \cdots \\ (1 - G_I)^2 \end{pmatrix} \sum_{y \leftrightarrow y'} [\mathcal{K}_{y \leftrightarrow y'} G^y - \mathcal{K}_{y \leftrightarrow y'} G^{y'}] \mathcal{H}_{y, y'}(G) (y' - y), \end{aligned} \quad (2.38)$$

and define

$$\mathcal{S}(G) = \sum_{y \leftrightarrow y'} [\mathcal{K}_{y \leftrightarrow y'} G^y - \mathcal{K}_{y \leftrightarrow y'} G^{y'}] \mathcal{H}_{y, y'}(G) (y' - y).$$

Following [16], we compute the Jacobian of \mathcal{S} at the equilibrium point G^* , applied to an arbitrary vector $\delta \neq 0$ that belongs to the span of the vectors $y' - y$

$$\text{Jac}(\mathcal{S}(G^*))\delta = \sum_{y \leftrightarrow y'} \mathcal{K}_{y \leftrightarrow y'}(G^*)^y ((y - y') * \delta) \mathcal{H}_{y, y'}(G^*) (y - y'), \quad (2.39)$$

in which the inner product $*$ is defined as

$$y * \delta = \sum_1^I \frac{y_k \delta_k}{G_k}.$$

Therefore

$$\begin{aligned} & [\text{Jac}(\mathcal{S}(G^*))\delta] * \delta \\ &= \sum_{y \leftrightarrow y'} \mathcal{K}_{y \leftrightarrow y'}(G^*)^y \mathcal{H}_{y, y'}(G^*) [(y - y') * \delta] [(y' - y) * \delta] < 0. \end{aligned} \quad (2.40)$$

Now, we compute the Jacobian of \mathcal{R} at the equilibrium point G^* ,

$$\begin{aligned} & \text{Jac}(\mathcal{R}(G^*)) \\ &= \text{diag} \begin{bmatrix} \partial_{G_1}(1 - G_1^*)^2 \mathcal{S}(G^*)_1 \\ \cdots \\ \partial_{G_I}(1 - G_I^*)^2 \mathcal{S}(G^*)_I \end{bmatrix} + \text{diag} \begin{bmatrix} (1 - G_1^*)^2 \\ \cdots \\ (1 - G_I^*)^2 \end{bmatrix} \text{Jac}(\mathcal{S}(G^*)) \\ &= \text{diag} \begin{bmatrix} (1 - G_1^*)^2 \\ \cdots \\ (1 - G_I^*)^2 \end{bmatrix} \text{Jac}(\mathcal{S}(G^*)), \end{aligned}$$

where the second equality is due to the fact that since G^* is an equilibrium we have that $\mathcal{S}(G^*) = 0$. Since

$$D := \text{diag} \begin{bmatrix} (1 - G_1^*)^2 \\ \cdots \\ (1 - G_I^*)^2 \end{bmatrix}$$

is a diagonal matrix and $A := \text{Jac}(\mathcal{S}(G^*))$ is negative definite, then $D^{1/2}AD^{1/2}$ is also negative definite with respect to this inner product. Since

$$\det(DA - \lambda Id) = \det(D^{1/2}AD^{1/2} - \lambda Id), \quad \forall \lambda \in \mathbb{R},$$

it follows that $D^{1/2}AD^{1/2}$ and DA have the same eigenvectors, so DA is negative definite. In other words, $\text{Jac}(\mathcal{R}(G^*))$ is negative definite. The exponential rate of convergence

$$\max\{|G_1(t) - G_1^*|, \dots, |G_I(t) - G_I^*|\} \leq C_1 e^{-C_2 t}.$$

then follows from the fact that the Jacobian above is negative definite. This leads to the conclusion of the theorem. \square

Remark 2.3. The Lyapunov function (2.32) in the variable F reads

$$L(F) = \sum_{k=1}^I [F_k \log F_k - (1 + F_k) \log(1 + F_k) + (\log(F_k^* + 1) - \log F_k^*)(F_k + 1)], \quad (2.41)$$

and it is a strictly convex function.

Remark 2.4. If the intersection between the ray $\{kP_0\}_{k \geq 1}$ and \mathcal{L}_R contains a single point, then the solution $f(t)$ of (2.1) has $f_{P_0} \equiv 0$, so $f_{P_0} \equiv \text{constant}$.

3. A REACTION NETWORK APPROACH FOR THE CASE OF C_{13} AND C_{22}

3.1. The dynamical system associated to C_{13}

As we discussed in the Introduction, we are also interested in the dynamics given by the discrete model of the collision operator C_{13} , described in (1.15).

Let \mathcal{L}_R denote the lattice of integer points

$$\mathcal{L}_R = \{p \in \mathbb{Z}^3 \mid |p| < R\}.$$

The discretized quantum Boltzmann equation for C_{13} reads

$$\begin{aligned}
\dot{f}_{p_1} &= C_{13}^D[f_{p_1}] \\
:= & \sum_{\substack{p_2, p_3, p_4 \in \mathcal{L}_R, \\ p_1 = p_2 + p_3 + p_4, \\ \mathcal{E}(p_1) = \mathcal{E}(p_2) + \mathcal{E}(p_3) + \mathcal{E}(p_4)}} K_{p_1, p_2, p_3, p_4}^{13} \{ (f_{p_1} + 1) f_{p_2} f_{p_3} f_{p_4} - (f_{p_2} + 1)(f_{p_3} + 1)(f_{p_4} + 1) f_{p_1} \} \\
& - 3 \sum_{\substack{p_2, p_3, p_4 \in \mathcal{L}_R, \\ p_2 = p_1 + p_3 + p_4, \\ \mathcal{E}(p_2) = \mathcal{E}(p_1) + \mathcal{E}(p_3) + \mathcal{E}(p_4)}} K_{p_2, p_1, p_3, p_4}^{13} \{ (f_{p_2} + 1) f_{p_1} f_{p_3} f_{p_4} - (f_{p_1} + 1)(f_{p_3} + 1)(f_{p_4} + 1) f_{p_2} \},
\end{aligned} \tag{3.1}$$

for all p_1 in \mathcal{L}_R , where $\mathcal{E}(p)$ is defined in (1.16).

Similar to the C_{12} case, when $p = 0$, $K_{p_1, p_2, p_3, p_4}^{13} = 0$, and we obtain

$$\dot{f}_0 = 0,$$

which means $f_0(t)$ is a constant for all time t , and we can assume $f_0(t) = 0$ for all t . Since in the first sum of (3.1), we consider (p_1, p_2, p_3, p_4) satisfying

$$p_1 = p_2 + p_3 + p_4, \quad \mathcal{E}(p_1) = \mathcal{E}(p_2) + \mathcal{E}(p_3) + \mathcal{E}(p_4), \tag{3.2}$$

we infer that there exists a vector P and $k_1, k_2, k_3, k_4 \geq 0$, $k_1, k_2, k_3, k_4 \in \mathbb{Z}$ such that

$$p_1 = k_1 P; \quad p_2 = k_2 P; \quad p_3 = k_3 P; \quad p_4 = k_4 P; \quad k_1 = k_2 + k_3 + k_4.$$

Using the same arguments as the case of C_{12} , we can deduce that equation (3.1) for C_{13} is equivalent with the following family of decoupled systems for $k_1 \in \mathbb{I} = \{1, 2, \dots, I\}$ where P is the closest point to the origin among the lattice points on its ray:

$$\begin{aligned}
& \dot{f}_{k_1 P} \\
= & \sum_{\substack{k_2, k_3, k_4 \in \mathbb{I}, \\ k_1 = k_2 + k_3 + k_4}} K_{k_1 P, k_2 P, k_3 P, k_4 P}^{13} \{ (f_{k_1 P} + 1) f_{k_2 P} f_{k_3 P} f_{k_4 P} \\
& - f_{k_1 P} (f_{k_2 P} + 1)(f_{k_3 P} + 1)(f_{k_4 P} + 1) \} \\
& - 3 \sum_{\substack{k_2, k_3, k_4 \in \mathbb{I}, \\ k_2 = k_1 + k_3 + k_4}} K_{k_2 P, k_1 P, k_3 P, k_4 P}^{13} \{ (f_{k_2 P} + 1) f_{k_1 P} f_{k_3 P} f_{k_4 P} \\
& - f_{k_2 P} (f_{k_1 P} + 1)(f_{k_3 P} + 1)(f_{k_4 P} + 1) \}.
\end{aligned} \tag{3.3}$$

Denoting f_{kP} by F_k (with $k \in \mathbb{I}$) and $K_{k_1 P, k_2 P, k_3 P, k_4 P}^{12}$ by $\mathcal{K}_{k_1, k_2, k_3, k_4}^{12}$, we obtain

$$\begin{aligned}
\dot{F}_{k_1} &= C_{13}[F](k_1) = \sum_{k_1 = k_2 + k_3 + k_4} \mathcal{K}_{k_1, k_2, k_3, k_4}^{13} \{ (F_{k_1} + 1) F_{k_2} F_{k_3} F_{k_4} \\
& - F_{k_1} (F_{k_2} + 1)(F_{k_3} + 1)(F_{k_4} + 1) \} \\
& - 3 \sum_{k_1 + k_2 + k_3 = k_4} \mathcal{K}_{k_1, k_2, k_3, k_4}^{13} \{ (F_{k_4} + 1) F_{k_1} F_{k_2} F_{k_3} \\
& - F_{k_4} (F_{k_1} + 1)(F_{k_2} + 1)(F_{k_3} + 1) \}, \quad \forall k_1 \in \mathbb{I}.
\end{aligned} \tag{3.4}$$

In order to ensure that all the variables F_k are coupled with each other, let us assume that $I \geq 4$. We have the following conservation of energy for C_{13}

$$\sum_{k=1}^I k \dot{F}_k = 0, \quad (3.5)$$

or equivalently

$$\sum_{k=1}^I k F_k = \text{const.} \quad (3.6)$$

Similar to the case of C_{12} , we define

$$G_k = \frac{F_k}{F_k + 1},$$

and then we have

$$F_k = \frac{G_k}{1 - G_k}.$$

Note that, similar to the previous section, $0 < F_k < \infty$ and $0 < G_k < 1$. The system (3.4) can be now written

$$\begin{aligned} & \frac{\dot{G}_{k_1}}{(1 - G_{k_1})^2} = \overline{C_{13}}[G] \\ & := \mathcal{K}_{k_1, k_2, k_3, k_4}^{13} \sum_{\substack{k_1 = k_2 + k_3 + k_4, \\ |k_1| = |k_2| + |k_3| + |k_4|}} \frac{G_{k_2} G_{k_3} G_{k_4} - G_{k_1}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})(1 - G_{k_4})} \\ & \quad - 3\mathcal{K}_{k_2, k_1, k_3, k_4}^{13} \sum_{\substack{k_2 = k_1 + k_3 + k_4, \\ |k_2| = |k_1| + |k_3| + |k_4|}} \frac{G_{k_1} G_{k_3} G_{k_4} - G_{k_2}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})(1 - G_{k_4})}. \end{aligned} \quad (3.7)$$

This system can also be rewritten as

$$\begin{aligned} \dot{G} &= \text{diag} \begin{pmatrix} (1 - G_1)^2 \\ \dots \\ (1 - G_I)^2 \end{pmatrix} \\ & \times \sum_{\substack{k_1 = k_2 + k_3 + k_4, \\ |k_1| = |k_2| + |k_3| + |k_4|}} [K_{\bar{X}_{k_2 + \bar{X}_{k_3} + \bar{X}_{k_4}} \rightarrow \bar{X}_{k_1}}(G) - K_{\bar{X}_{k_1} \rightarrow \bar{X}_{k_2} + \bar{X}_{k_3} + \bar{X}_{k_4}}(G)] (\bar{X}_{k_1} - \bar{X}_{k_2} - \bar{X}_{k_3} - \bar{X}_{k_4}). \end{aligned} \quad (3.8)$$

where \bar{X}_k is, as mentioned earlier, the vector

$$\begin{pmatrix} 0 \\ \dots \\ 1 \\ \dots \\ 0 \end{pmatrix},$$

in which the only element that is 1 is the k -th one, and

$$K_{\bar{X}_{k_2} + \bar{X}_{k_3} + \bar{X}_{k_4} \rightarrow \bar{X}_{k_1}}(G) := \mathcal{K}_{k_1, k_2, k_3, k_4}^{13} \frac{G_{k_1}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})(1 - G_{k_4})},$$

$$K_{\bar{X}_{k_1} \rightarrow \bar{X}_{k_2} + \bar{X}_{k_3} + \bar{X}_{k_4}}(G) := \mathcal{K}_{k_1, k_2, k_3, k_4}^{13} \frac{G_{k_2} G_{k_3} G_{k_4}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})(1 - G_{k_4})}.$$

We can also write

$$\dot{G} = \text{diag} \left(\begin{array}{c} (1 - G_1)^2 \\ \cdots \\ (1 - G_I)^2 \end{array} \right) \sum_{y \leftrightarrow y'} [K_{y \rightarrow y'}(G) - K_{y' \rightarrow y}(G)] (y' - y),$$

where $y \leftrightarrow y'$ rang over the reversible reactions shown above.

Theorem 3.1. *For any initial condition, the solution*

$$f(t) = (f_p(t))_{p \in \mathcal{L}_R}$$

of the quantum Boltzmann equation (3.1) converges to an equilibrium state $f^ = (f_p^*)_{p \in \mathcal{L}_R}$. For each ray $\{kP_0\}_{k \geq 1}$ that intersects \mathcal{L}_R in at least 4 points there exists a constant ρ_{P_0} such that if $p = kP_0$ then*

$$f_p^* = \frac{1}{e^{k\rho_{P_0}} - 1}.$$

Moreover, the solution $f(t)$ of (3.1) converges to f^ exponentially fast in the following sense: there exists positive constants C_1, C_2 such that*

$$\max_{p \in \mathcal{L}_R} |f_p(t) - f_p^*| < C_1 e^{-C_2 t}.$$

Proof. The proof of Theorem 3.2 then follows exactly from the same Lyapunov function (2.32) and arguments as in Theorem 2.2. \square

3.2. The dynamical system associated to C_{22}

Let us consider a discretized version of the quantum Boltzmann model associated to the collision operator given by C_{22} :

Let \mathcal{L}_R denote the lattice of integer points

$$\mathcal{L}_R = \{p \mid |p| \in \mathbb{Z}^3, |p| < R\}.$$

The discretized quantum Boltzmann equation associated to C_{22} reads $\forall p_1 \in \mathcal{L}_R$

$$\begin{aligned} \dot{f}_{p_1} &= C_{22}^D[f_{p_1}] \\ := & \sum_{\substack{p_2, p_3, p_4 \in \mathcal{L}_R, \\ p_1 + p_2 = p_3 + p_4, \\ \mathcal{E}(p_1) + \mathcal{E}(p_2) = \mathcal{E}(p_3) + \mathcal{E}(p_4)}} K_{p_1, p_2, p_3, p_4}^{13} \{ (f_{p_1} + 1)(f_{p_2} + 1)f_{p_3}f_{p_4} - f_{p_1}f_{p_2}(f_{p_3} + 1)(f_{p_4} + 1) \}, \end{aligned} \quad (3.9)$$

where $\mathcal{E}(p)$ is defined in (1.16).

Similar to the C_{12} case, when $p = 0$, $K_{p_1, p_2, p_3, p_4}^{22} = 0$, and we obtain

$$\dot{f}_0 = 0,$$

which means $f_0(t)$ is a constant for all time t . As a consequence, we can suppose that $f_0(0) = 0$, which implies $f_0(t) = 0$ for all t .

In (3.9), the sums for C_{22} are taken over (p_1, p_2, p_3, p_4) satisfying

$$p_1 + p_2 = p_3 + p_4, \text{ and } \mathcal{E}(p_1) + \mathcal{E}(p_2) = \mathcal{E}(p_3) + \mathcal{E}(p_4). \quad (3.10)$$

In this case, unlike in the case of C_{12} and C_{13} , we *cannot* infer from (3.10) that there exists a vector P and $k_1, k_2, k_3, k_4 \geq 0$, $k_1, k_2, k_3, k_4 \in \mathbb{Z}$ such that

$$p_1 = k_1 P; \quad p_2 = k_2 P; \quad p_3 = k_3 P; \quad p_4 = k_4 P, \quad k_1 + k_2 = k_3 + k_4.$$

However, let us consider the following simplified version of (3.9) for C_{22}

$$\begin{aligned} \dot{F}_{k_1} = C_{22}[F](k_1) := & \sum_{\substack{k_1+k_2=k_3+k_4 \\ k_2, k_3, k_4 \in \mathbb{I}}} \mathcal{K}_{k_1, k_2, k_3, k_4}^{13} \{(F_{k_1} + 1)(F_{k_2} + 1)F_{k_3}F_{k_4} \\ & - F_{k_1}F_{k_2}(F_{k_3} + 1)(F_{k_4} + 1)\}, \quad \forall k_1 \in \mathbb{I}. \end{aligned} \quad (3.11)$$

Recall that $\mathbb{I} = \{1, \dots, I\}$. We also suppose that $I \geq 3$. We have the following conservation of energy

$$\sum_{k=1}^I k \dot{F}_k = 0, \quad (3.12)$$

or equivalently

$$\sum_{k=1}^I k F_k = \text{const.} \quad (3.13)$$

For C_{22} , the following ‘‘conservation of mass’’ also holds

$$\sum_{k=1}^I \dot{F}_k = 0, \quad (3.14)$$

or equivalently

$$\sum_{k=1}^I F_k = \text{const.} \quad (3.15)$$

Similar to the case of C_{12} , define

$$G_k = \frac{F_k}{F_k + 1},$$

then

$$F_k = \frac{G_k}{1 - G_k},$$

and the system (3.9) can be now written

$$\begin{aligned} \frac{\dot{G}_{k_1}}{(1 - G_{k_1})^2} &= \overline{\mathcal{C}}_{22}[G] \\ := \mathcal{K}_{k_1, k_2, k_3, k_4}^{13} &\sum_{\substack{k_1 + k_2 = k_3 + k_4, \\ |k_1| + |k_2| = |k_3| + |k_4|}} \frac{G_{k_3} G_{k_4} - G_{k_1} G_{k_2}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})(1 - G_{k_4})}. \end{aligned} \quad (3.16)$$

This system can be rewritten as

$$\begin{aligned} \dot{G} &= \text{diag} \begin{pmatrix} (1 - G_1)^2 \\ \cdots \\ (1 - G_I)^2 \end{pmatrix} \\ &\times \sum_{\substack{k_1 + k_2 = k_3 + k_4, \\ |k_1| + |k_2| = |k_3| + |k_4|}} [K_{\bar{X}_{k_3} + \bar{X}_{k_4} \rightarrow \bar{X}_{k_2} + \bar{X}_{k_1}}(G) \\ &- K_{\bar{X}_{k_2} + \bar{X}_{k_1} \rightarrow \bar{X}_{k_3} + \bar{X}_{k_4}}(G)] (\bar{X}_{k_1} + \bar{X}_{k_2} - \bar{X}_{k_3} - \bar{X}_{k_4}). \end{aligned} \quad (3.17)$$

where \bar{X}_k is, the vector

$$\begin{pmatrix} 0 \\ \cdots \\ 1 \\ \cdots \\ 0 \end{pmatrix},$$

in which the only element that is 1 is the k -th one, and

$$\begin{aligned} K_{\bar{X}_{k_3} + \bar{X}_{k_4} \rightarrow \bar{X}_{k_2} + \bar{X}_{k_1}}(G) &= \mathcal{K}_{k_1, k_2, k_3, k_4}^{13} \frac{G_{k_1} G_{k_2}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})(1 - G_{k_4})}, \\ K_{\bar{X}_{k_2} + \bar{X}_{k_1} \rightarrow \bar{X}_{k_3} + \bar{X}_{k_4}}(G) &= \mathcal{K}_{k_1, k_2, k_3, k_4}^{13} \frac{G_{k_3} G_{k_4}}{(1 - G_{k_1})(1 - G_{k_2})(1 - G_{k_3})(1 - G_{k_4})}. \end{aligned}$$

We can also write

$$\dot{G} = \text{diag} \begin{pmatrix} (1 - G_1)^2 \\ \cdots \\ (1 - G_I)^2 \end{pmatrix} \sum_{y \leftrightarrow y'} [K_{y \rightarrow y'}(G) - K_{y' \rightarrow y}(G)] (y' - y),$$

where $y \leftrightarrow y'$ range over the reversible reactions shown above.

Theorem 3.2. *For any initial condition, the solution*

$$F(t) = (F_k(t))_{k \in \mathbb{I}}$$

of the quantum Boltzmann equation (3.11) converges to an equilibrium state $F^ = (F_k^*)_{k \in \mathbb{I}}$, where*

$$F_k^* = \frac{1}{e^{\rho_2(k-1) - \rho_1(k-2)} - 1}.$$

Moreover, the solution $F(t)$ of (3.11) converges to F^ exponentially fast in the following sense: there exist positive constants C_1, C_2 such that*

$$\max_{k \in \mathbb{I}} |F_k(t) - F_k^*| < C_1 e^{-C_2 t}.$$

Proof. We set

$$G_k^* = \frac{F_k^*}{F_k^* + 1}.$$

By the same argument used to obtain (2.37), we deduce that

$$\frac{G^y}{(G^*)^y} = \frac{G^{y'}}{(G^*)^{y'}} \quad (3.18)$$

holds true for all reactions $y \leftrightarrow y'$, which implies $G^y = G^{y'}$ since $(G^*)^y = (G^*)^{y'}$ for all reactions $y \leftrightarrow y'$. In the case of C_{22} , we obtain the relation $G_{k_1}^* \cdot G_{k_2}^* = G_{k_3}^* \cdot G_{k_4}^*$ for all k_1, k_2, k_3, k_4 such that $k_1 + k_2 = k_3 + k_4 \leq I$. From the relation $G_{k_1} \cdot G_{k_2} = G_{k_3} \cdot G_{k_4}$ and the fact that $k + (k - 2) = 2(k - 1)$, the following identity holds true

$$G_k (G_1)^{k-2} = (G_2)^{k-1}.$$

We then obtain $G_k = (G_2)^{k-1} / (G_1)^{k-2}$, which leads to $G_k = e^{\rho_2'(k-1) - \rho_1'(k-2)}$ for some numbers ρ_1', ρ_2' .

Since

$$\sum_{k=1}^I \frac{k G_k}{1 - G_k} = \sum_{k=1}^I \frac{k G_k^*}{1 - G_k^*},$$

we then have

$$\sum_{k=1}^I \frac{k e^{\rho_2'(k-1) - \rho_1'(k-2)}}{1 - e^{\rho_2'(k-1) - \rho_1'(k-2)}} = \sum_{k=1}^I \frac{k e^{\rho_2(k-1) - \rho_1(k-2)}}{1 - e^{\rho_2(k-1) - \rho_1(k-2)}}.$$

We can proceed like in [32] to obtain $\tilde{\rho}' = \tilde{\rho}$ and $\bar{\rho} = \bar{\rho}'$, yielding $\rho_1 = \rho_1'$ and $\rho_2 = \rho_2'$.

We can still use the Petri net argument of [4] or the result in [14], to prove that the system is persistent. For example, to use the method from [4], we note that we have two *siphons* $\{X_1, X_2, \dots, X_I\}, \{X_2, \dots, X_I\}$. However, we also have the conservations of mass and energy

$$\sum_{k=1}^I F_k = \text{constant},$$

$$\sum_{k=1} k F_k = \text{constant},$$

that leads to the P -semiflow

$$\sum_{k=2} (k-1) F_k = \text{constant}.$$

Therefore, similar to the case of C_{12} , it follows that the system is persistent, and we can use the same Lyapunov function as in the proof of Theorem 2.2 to obtain the desired convergence result. \square

Remark 3.3. If $I < 3$ then $F_k^* \equiv 0$. If $I = 3$ then $F_2^* \equiv 0$ and $F_1^* = \frac{1}{e^\rho - 1}$, $F_3^* = \frac{1}{e^{3\rho} - 1}$ for some $\rho = \rho(P_0)$.

4. A REACTION NETWORK APPROACH FOR THE SUM OF C_{12}, C_{22}, C_{13}

Let us consider the following equations

$$\dot{F}_{k_1} = C_{12}[F](k_1) + C_{22}[F](k_1), \tag{4.1}$$

and

$$\dot{F}_{k_1} = C_{12}[F](k_1) + C_{22}[F](k_1) + C_{13}[F](k_1), \tag{4.2}$$

where C_{12}, C_{22}, C_{13} are the operators defined in (2.7), (3.4), (3.11).

The following theorem then follows by exactly the same argument as in Theorem 3.2

Theorem 4.1. *For any initial condition, the solution*

$$F(t) = (F_k(t))_{k \in \mathbb{I}}$$

of the quantum Boltzmann equation (4.1) or (4.2) converges to an equilibrium state $F^ = (F_k^*)_{k \in \mathbb{I}}$, where $F_k^* = \frac{1}{e^{\rho k} - 1}$ for some constant ρ . Moreover, the solution $F(t)$ of (3.4) converges to F^* exponentially fast in the following sense: there exists positive constants C_1, C_2 such that*

$$\max_{k \in \mathbb{I}} |F_k(t) - F_k^*| < C_1 e^{-C_2 t}.$$

5. CONCLUSION

In this work, we point out a connection between quantum Boltzmann models derived in [49] and chemical reaction network models. We prove that the discrete, simplified versions of some differential equations for these quantum Boltzmann models relax to an equilibrium point, by a toric dynamical system approach, similar to the one used in a recently proposed proof of the global attractor conjecture [14].

CONFLICT OF INTEREST

The authors state that there is no conflict of interest.

Acknowledgements. The authors would like to thank the editors for the invitation to submit the paper for the special issue of the journal. They are also indebted to Professor Roland Glowinski for his encouragement. G. Craciun was supported by NSF grants DMS-1412643 and DMS-1816238, and by a Simons Foundation fellowship. M.-B. Tran is funded in part by the NSF Grant DMS-1854453, NSF RTG Grant DMS-1840260, SMU URC Grant 2020, Humboldt Fellowship, SMU Dedman College Linking Fellowship, NSF CAREER DMS-2044626.

REFERENCES

- [1] R. Alonso, I.M. Gamba and M.-B. Tran, The Cauchy problem and BEC stability for the quantum Boltzmann-condensation system for bosons at very low temperature. Preprint [arXiv:1609.07467](https://arxiv.org/abs/1609.07467) (2016).
- [2] D.F. Anderson, A proof of the global attractor conjecture in the single linkage class case. *SIAM J. Appl. Math.* **71** (2011) 1487–1508.
- [3] D.F. Anderson, G. Craciun and T.G. Kurtz, Product-form stationary distributions for deficiency zero chemical reaction networks. *Bull. Math. Biol.* **72** (2010) 1947–1970.
- [4] D. Angeli, P. De Leenheer and E.D. Sontag, Persistence results for chemical reaction networks with time-dependent kinetics and no global conservation laws. *SIAM J. Appl. Math.* **71** (2011) 128–146.
- [5] L. Arkeryd and A. Nouri, Bose condensates in interaction with excitations: a kinetic model. *Commun. Math. Phys.* **310** (2012) 765–788.
- [6] L. Arkeryd and A. Nouri, A Milne problem from a Bose condensate with excitations. *Kinet. Relat. Models* **6** (2013) 671–686.
- [7] L. Arkeryd and A. Nouri, Bose condensates in interaction with excitations: a two-component space-dependent model close to equilibrium. *J. Stat. Phys.* **160** (2015) 209–238.
- [8] M. Banaji and G. Craciun, Graph-theoretic approaches to injectivity and multiple equilibria in systems of interacting elements. *Commun. Math. Sci.* **7** (2009) 867–900.
- [9] N. Bernhoff, Half-space problems for a linearized discrete quantum kinetic equation. *J. Stat. Phys.* **159** (2015) 358–379.
- [10] N. Bernhoff, Boundary layers for discrete kinetic models: multicomponent mixtures, polyatomic molecules, bimolecular reactions, and quantum kinetic equations. *Kinetic Related Models* **10** (2017) 925.
- [11] L. Boltzmann, Neuer Beweis zweier Satze uber das Warmegleichgewicht unter mehratomigen Gas-molekullen. *Sitzungsberichte der Kaiserlichen Akademie der Wissenschaften in Wien* (1887) 153–164.
- [12] S. Cai and X. Lu, The spatially homogeneous Boltzmann equation for Bose-Einstein particles: rate of strong convergence to equilibrium. Preprint [arXiv:1808.04038](https://arxiv.org/abs/1808.04038) (2018).
- [13] G. Craciun, Polynomial dynamical systems, reaction networks, and toric differential inclusions. *SIAM J. Appl. Algebra Geometry* **3** (2019) 87–106.
- [14] G. Craciun, Toric differential inclusions and a proof of the global attractor conjecture. Submitted.
- [15] G. Craciun, A. Dickenstein, A. Shiu and B. Sturmfels, Toric dynamical systems. *J. Symbolic Comput.* **44** (2009) 1551–1565.
- [16] G. Craciun and M. Feinberg, Multiple equilibria in complex chemical reaction networks. I. The injectivity property. *SIAM J. Appl. Math.* **65** (2005) 1526–1546.
- [17] G. Craciun, F. Nazarov and C. Pantea, Persistence and permanence of mass-action and power-law dynamical systems. *SIAM J. Appl. Math.* **73** (2013) 305–329.
- [18] M. Escobedo and M.-B. Tran, Convergence to equilibrium of a linearized quantum Boltzmann equation for bosons at very low temperature. *Kinetic Related Models* **8** (2015) 493–531.
- [19] M. Escobedo and J.J.L. Velázquez, Finite time blow-up and condensation for the bosonic Nordheim equation. *Invent. Math.* **200** (2015) 761–847.
- [20] M. Feinberg, Lectures on chemical reaction networks. Written version of lectures given at the Mathematical Research Center, University of Wisconsin, Madison WI, 1979. Available at <http://www.crn.tu.edu/LecturesOnReactionNetworks>.
- [21] M. Feinberg, Complex balancing in general kinetic systems. *Arch. Ratl. Mech. Anal.* **49** (1972/73) 187–194.
- [22] M. Feinberg, The existence and uniqueness of steady states for a class of chemical reaction networks. *Arch. Ratl. Mech. Anal.* **132** (1995) 311–370.
- [23] P. Germain, A.D. Ionescu and M.-B. Tran, Optimal local well-posedness theory for the kinetic wave equation. *J. Funct. Anal.* **279** (2020) 108570.
- [24] M. Gopalkrishnan, E. Miller and A. Shiu, A geometric approach to the global attractor conjecture. *SIAM J. Appl. Dyn. Syst.* **13** (2014) 758–797.
- [25] A. Griffin, T. Nikuni and E. Zaremba, Bose-condensed gases at finite temperatures. Cambridge University Press, Cambridge (2009).
- [26] A. Griffin, T. Nikuni and E. Zaremba, Bose-condensed gases at finite temperatures. Cambridge University Press, Cambridge (2009).
- [27] J. Gunawardena, Chemical reaction network theory for in-silico biologists. Lecture notes available online at <http://vcp.med.harvard.edu/papers.html> (2003)
- [28] E.D. Gust and L.E. Reichl, Collision integrals in the kinetic equations of dilute Bose-Einstein condensates. Preprint [arXiv:1202.3418](https://arxiv.org/abs/1202.3418) (2012).
- [29] E.D. Gust and L.E. Reichl, Relaxation rates and collision integrals for Bose-Einstein condensates. *Phys. Rev. A* **170** (2013) 43–59.
- [30] F. Horn, Necessary and sufficient conditions for complex balancing in chemical kinetics. *Arch. Ratl. Mech. Anal.* **49** (1972/73) 172–186.
- [31] F. Horn, The dynamics of open reaction systems. In Mathematical aspects of chemical and biochemical problems and quantum chemistry (*Proc. SIAM-AMS Sympos. Appl. Math., New York, 1974*). SIAM-AMS Proceedings, Vol. VIII. Amer. Math. Soc., Providence, R.I. (1974) 125–137.
- [32] F. Horn and R. Jackson, General mass action kinetics. *Arch. Ratl. Mech. Anal.* **47** (1972) 81–116.
- [33] S. Jin and M.-B. Tran, Quantum hydrodynamic approximations to the finite temperature trapped Bose gases. *Physica D* **380** (2018) 45–57.

- [34] T.R. Kirkpatrick and J.R. Dorfman, Transport theory for a weakly interacting condensed Bose gas. *Phys. Rev. A* **28** (1983) 2576–2579.
- [35] T.R. Kirkpatrick and J.R. Dorfman, Transport in a dilute but condensed nonideal Bose gas: kinetic equations. *J. Low Temp. Phys.* **58** (1985) 301–331.
- [36] X. Lu, On isotropic distributional solutions to the Boltzmann equation for Bose-Einstein particles. *J. Statist. Phys.* **116** (2004) 1597–1649.
- [37] X. Lu, The Boltzmann equation for Bose-Einstein particles: velocity concentration and convergence to equilibrium. *J. Stat. Phys.* **119** (2005) 1027–1067.
- [38] X. Lu, The Boltzmann equation for Bose-Einstein particles: condensation in finite time. *J. Stat. Phys.* **150** (2013) 1138–1176.
- [39] T.T. Nguyen and M.-B. Tran, Uniform in time lower bound for solutions to a quantum Boltzmann equation of bosons. *Arch. Ratl. Mech. Anal.* **231** (2019) 63–89.
- [40] L.W. Nordheim, Transport phenomena in Einstein–Bose and Fermi–Dirac gases. *Proc. Roy. Soc. London A* **119** (1928) 689.
- [41] Y. Pomeau and M.-B. Tran, Vol. 967 of *Statistical Physics of Non Equilibrium Quantum Phenomena*. Springer Nature (2019).
- [42] L.E. Reichl, *A modern course in statistical physics*. A Wiley-Interscience Publication. John Wiley & Sons, Inc., New York (2016) fourth edition.
- [43] L.E. Reichl and E.D. Gust, Transport theory for a dilute Bose-Einstein condensate. *J. Low Temp. Phys.* **88** (2013) 053603.
- [44] L.E. Reichl and M.-B. Tran, A kinetic equation for ultra-low temperature Bose–Einstein condensates. *J. Phys. A: Math. Theor.* **52** (2019) 063001.
- [45] A. Soffer and M.-B. Tran, On coupling kinetic and Schrodinger equations. *J. Differ. Equ.* **265** (2018) 2243–2279.
- [46] A. Soffer and M.-B. Tran, On the dynamics of finite temperature trapped Bose gases. *Adv. Math.* **325** (2018) 533–607.
- [47] H. Spohn, Kinetics of the Bose-Einstein condensation. *Physica D* **239** (2010) 627–634.
- [48] M.-B. Tran, G. Craciun, L.M. Smith and S. Boldyrev, A reaction network approach to the theory of acoustic wave turbulence. *J. Differ. Equ.* **269** (2020) 4332–4352.
- [49] M.-B. Tran and Y. Pomeau, Boltzmann-type collision operators for Bogoliubov excitations of Bose-Einstein condensates: a unified framework. *Phys. Rev. E* **101** (2020) 032119.
- [50] M.-B. Tran and Y. Pomeau, On a thermal cloud - Bose-Einstein condensate coupling system. *Eur. Phys. J. Plus* **136** (2021) 502.
- [51] E.A. Uehling and G.E. Uhlenbeck, On the kinetic methods in the new statistics and its applications in the electron theory of conductivity. *Phys. Rev.* **43** (1933) 552–561.
- [52] S. M’etens Y. Pomeau, M.A. Brachet and S. Rica, Théorie cinétique d’un gaz de Bose dilué avec condensat. *C. R. Acad. Sci. Paris Ser. IIb M’ec. Phys. Astr.* **327** (1999) 791–798.
- [53] P.Y. Yu and G. Craciun, Mathematical analysis of chemical reaction systems. *Israel J. Chem.* **58** (2018) 733–741.
- [54] E. Zaremba, T. Nikuni and A. Griffin, Dynamics of trapped Bose gases at finite temperatures. *J. Low Temp. Phys.* **116** (1999) 277–345.