

Quantum Techniques for Reaction Networks

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March 20, 2014

Abstract

Reaction networks are a general formalism for describing collections of classical entities interacting in a random way. While reaction networks are mainly studied by chemists, they are equivalent to Petri nets, which are used for similar purposes in computer science and biology. As noted by Doi and others, techniques from quantum field theory can be adapted to apply to such systems. Here we use these techniques to study how the ‘master equation’ describing stochastic time evolution for a reaction network is related to the ‘rate equation’ describing the deterministic evolution of the expected number of particles of each species in the large-number limit. We show that the relation is especially strong when a solution of master equation is a ‘coherent state’, meaning that the numbers of entities of each kind are described by independent Poisson distributions.

1 Introduction

A ‘reaction network’ describes how various kinds of classical particles can interact and turn into other kinds. They are commonly used in chemistry. For example, this reaction network gives a very simplified picture of what is happening in a glass of water:



The reactions here are all reversible, but this is not required by the reaction network formalism. The relevant particles here are not atoms, but rather ‘species’ including the water molecule H_2O , the proton H^+ , the hydroxyl ion OH^- and the hydronium ion H_3O^+ .

While the dynamics of the reactions is fundamentally quantum-mechanical, reaction networks give a simplified description of what is going on. First, given a ‘rate constant’ for each reaction, one can write down a differential equation called the *rate equation*, which describes the time evolution of the concentration of each species in a deterministic way. This is especially useful in the limit where there are many particles of each species. At a more fine-grained level, one can describe reactions stochastically, using the *master equation*. In the rate equation the concentration of each species is treated as a continuous variable, but in the master equation, the number of particles of each species is an integer.

In this paper we explain how techniques from quantum field theory can be used to study reaction networks. This work is part of a broader program emphasizing the links between quantum mechanics and a subject we call ‘stochastic mechanics’, where probabilities replace amplitudes [4, 5, 6].

Using ideas from quantum field theory to model classical stochastic systems goes back at least to 1976, with the work of Doi [7]. It has been further developed by Grassberger, Scheunert [9] and many others. The big surprise one meets at the very beginning of this subject is that the canonical commutation relations between annihilation and creation operators, long viewed as a hallmark of quantum theory, are also perfectly suited to systems of identical *classical* particles interacting in a probabilistic way.

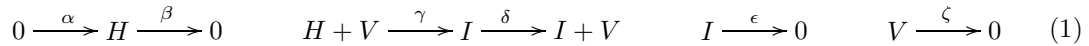
There is an extensive line of research on the rate and master equations in chemistry, initiated by Horn and Jackson [10] along with Feinberg [8] in the late 1970's. This line of work does not use techniques from quantum field theory, but it might profit from them, and also contribute to their development. To illustrate how, here we use these techniques to study how the master equation is related to the 'rate equation', a deterministic equation which applies in the large-number limit. This is formally similar to the 'classical limit' in which a quantum field theory reduces to a classical field theory.

In Theorem 8 we start with the master equation and derive an equation for the rate of change of the expected number of particles of each species. This equation is similar to the rate equation, and reduces to it in a certain approximation. However, in Theorem 9 we show that this equation gives the rate equation *exactly* when the numbers of particles of each species are described by independent Poisson distributions. In this situation, the mathematical technology of coherent states can be applied. Coherent states are eigenvectors of the annihilation operators, and this fact explains their good behavior.

2 Reaction networks

A reaction network, first defined by Aris [3], consists of a set of different kinds of particles—each kind being called a 'species'—together with a set of processes, called 'reactions', that turn collections of particles of various species into collections of particles of other species. As the name suggests, it is convenient to draw a reaction network as a graph.

While reaction networks first arose in chemistry, their uses are not limited to this context. Here is an example that arose in work on HIV, the human immunodeficiency virus [14]:



Here we have three species:

- H : healthy white blood cells,
- I : infected white blood cells,
- V : virions (that is, individual virus particles).

We also have six reactions:

- α : the birth of one healthy cell, which has no input and one H as output.
- β : the death of a healthy cell, which has one H as input and no output.
- γ : the infection of a healthy cell, which has one H and one V as input, and one I as output.
- δ : 'production', or the reproduction of the virus in an infected cell, which has one I as input and one I and one V as output.
- ϵ : the death of an infected cell, which has one I as input and no output.
- ζ : the death of a virion, which has one V as input and no output.

So, we have a finite set of species

$$S = \{H, I, V\}$$

but reactions go between 'complexes', which are finite linear combinations of species with natural number coefficients, such as $H + V$ or $2H + 3V$. In the literature on reaction networks the complex 0, corresponding to nothing at all, is often denoted \emptyset . We can think of complexes as elements of \mathbb{N}^S ,

the set of functions from S to the natural numbers. The reaction network is a graph with certain complexes as vertices and reactions as edges.

We can formalize this as follows. There are various kinds of graphs; the kind we want are sometimes called ‘directed multigraphs’ or ‘quivers’, but to keep our terminology simple, we make the following definition:

Definition 1. A **graph** consists of a set E of **edges**, a set V of **vertices**, and **source** and **target** maps $s, t: E \rightarrow V$ saying where each edge starts and ends.

Then we may say:

Definition 2. A **reaction network** (S, K, T, s, t) consists of:

- a finite set S of **species**,
- a finite set of **complexes** $K \subseteq \mathbb{N}^S$,
- a graph $s, t: T \rightarrow K$ with complexes as vertices and some finite set of **reactions** T as edges.

In the chemistry literature this sort of thing is called a ‘chemical reaction network’, but we want to emphasize that they are useful more generally, so we drop the adjective. Petri nets are an alternative formalism that is entirely equivalent to reaction networks, often used in subjects other than chemistry [12, 13, 15]. In the Petri net literature the species are called ‘places’ and the reactions are called ‘transitions’. For a more complete translation guide between reaction networks and Petri nets, see [6].

For convenience we shall often write $k = |S|$ for the number of species present in a reaction network, and identify the set S with the set $\{1, \dots, k\}$. This lets us write any complex as a k -tuple of natural numbers. In particular, we write the source and target of any reaction τ as

$$s(\tau) = (s_1(\tau), \dots, s_k(\tau)) \in \mathbb{N}^k,$$

$$t(\tau) = (t_1(\tau), \dots, t_k(\tau)) \in \mathbb{N}^k.$$

3 The rate equation

The amount of each species is represented by a ‘state’ of the chemical reaction network. There are various options here. A **pure state** is a vector $x \in \mathbb{N}^k$ with i th entry x_i specifying the number of instances of the i th species. This generalises in two ways. First, we define a **classical state** to be any vector $x \in [0, \infty)^k$ of nonnegative real numbers, and think of such a state as specifying the expected number or concentration of the species present. Second, we define a **mixed state** to be a probability distribution over the pure states. This assigns a probability ψ_ℓ to each $\ell \in \mathbb{N}^k$. The rate equation describes the time evolution of a classical state, while the master equation describes the time evolution of a mixed state.

To write down either of these equations, we must first specify the system’s dynamics. In this paper we consider systems that evolve by the law of mass action [3]. Very roughly, this law states that the rate at which a reactions occur is proportional to the product of the concentrations of all its input species. We call the constant of proportionality for each reaction its **rate constant**:

Definition 3. A **stochastic reaction network** (S, K, T, s, t, r) consists of a reaction network together with a map $r: T \rightarrow (0, \infty)$ assigning a **rate constant** to each reaction.

This concept is equivalent to another concept in the literature, namely that of a stochastic Petri net [6].

Given a stochastic reaction network, the rate equation says that for each reaction τ the time derivative of a classical state x is the product of:

- the vector $t(\tau) - s(\tau)$ describing the change in numbers of each species due to τ ;
- the concentration of each input species i of τ raised to the power given by the number of times it appears as an input, namely $s_i(\tau)$;
- the rate constant $r(\tau)$ of τ .

Definition 4. The **rate equation** for a stochastic reaction network (S, K, T, s, t, r) is

$$\frac{d}{dt}x(t) = \sum_{\tau \in T} r(\tau)(t(\tau) - s(\tau))x(t)^{s(\tau)}$$

where $x: \mathbb{R} \rightarrow [0, \infty)^k$ and we have used multi-index notation to define

$$x^{s(\tau)} = x_1^{s_1(\tau)} \cdots x_k^{s_k(\tau)}.$$

For the HIV reaction network in Equation 1, if we also use the Greek letter names of the reactions as names for their rate constants, we get this rate equation:

$$\begin{aligned} \frac{dH}{dt} &= \alpha - \beta H - \gamma HV \\ \frac{dI}{dt} &= \gamma HV - \epsilon I \\ \frac{dV}{dt} &= -\gamma HV + \delta I - \zeta V \end{aligned}$$

4 The master equation

The master equation describes the time evolution of mixed states. For the rest of this paper we fix a stochastic reaction network (S, K, T, s, t, r) . Recall that for each $\ell \in \mathbb{N}^k$, a ‘mixed state’ gives the probability ψ_ℓ that for each i , exactly $\ell_i \in \mathbb{N}$ instances of the i th species are present. The **master equation** says that

$$\frac{d}{dt}\psi_{\ell'} = \sum_{\ell \in \mathbb{N}^k} H_{\ell'\ell} \psi_\ell$$

for some matrix of numbers H determined by the stochastic reaction network. The matrix element $H_{\ell'\ell}$ is the probability per time for the pure state ℓ to evolve to the pure state ℓ' . This probability is a sum over reactions:

$$H_{\ell'\ell} = \sum_{\tau \in T} H(\tau)_{\ell'\ell}. \quad (2)$$

To describe the matrix $H(\tau)$, we need ‘falling powers’. For any natural numbers n and p , we define the p th **falling power** of n by

$$n^{\underline{p}} = n(n-1) \cdots (n-p+1).$$

This is the number of ways of choosing an ordered p -tuple of distinct elements from a set with n elements. Note that $n^{\underline{p}} = 0$ if $p > n$. More generally, for any multi-indices ℓ and m , we define

$$\ell^{\underline{m}} = \ell_1^{\underline{m_1}} \cdots \ell_k^{\underline{m_k}}$$

This is the number of ways to choose, for each species $i = 1, \dots, k$, an ordered list of m_i distinct things from a collection of ℓ_i things of that species. Thus, we expect a factor of this sort to appear in the master equation.

Using this notation, we define the matrix $H(\tau)$ as follows:

$$H(\tau)_{\ell'\ell} = r(\tau) \frac{\ell^{s(\tau)}}{\ell^{s(\tau)}} (\delta_{\ell', \ell+t(\tau)-s(\tau)} - \delta_{\ell'\ell}). \quad (3)$$

Here δ is the Kronecker delta, which equals 1 if its subscripts are equal and 0 otherwise. The first term in this equation describes the rate for the complex ℓ to become the complex ℓ' via the reaction τ : it equals the rate constant for this reaction times the number of ways this reaction can occur. The second term describes the rate at which ℓ goes away due to this reaction. Thus, the size of the second term is equal to that of the first, but its sign is opposite.

Putting together Equations 2 and 3, we obtain this formula for the Hamiltonian:

$$H_{\ell'\ell} = \sum_{\tau \in T} r(\tau) \frac{\ell^{s(\tau)}}{\ell^{s(\tau)}} (\delta_{\ell', \ell+t(\tau)-s(\tau)} - \delta_{\ell'\ell}). \quad (4)$$

5 The stochastic Fock space

We can understand the Hamiltonian for the master equation at a deeper level using techniques borrowed from quantum field theory. The first step is to introduce a stochastic version of Fock space. To do this, we write any mixed state as a formal power series in some variables z_1, \dots, z_k , with the coefficient of

$$z^\ell = z_1^{\ell_1} \dots z_k^{\ell_k}$$

being the probability ψ_ℓ . That is, we write any mixed state as

$$\Psi = \sum_{\ell \in \mathbb{N}^k} \psi_\ell z^\ell.$$

Because a mixed state is a probability distribution, the coefficients ψ_ℓ must be nonnegative and sum to 1. Indeed, in this formalism **mixed states** are precisely the formal power series Ψ with

$$\sum_{\ell \in \mathbb{N}^k} \psi_\ell = 1, \quad \psi_\ell \geq 0.$$

The simplest mixed states are the monomials z^ℓ ; these are the **pure states**, where there is a definite number of things of each species.

In quantum mechanics we use a similar sort of Fock space, but with power series having complex rather than real coefficients. In quantum mechanics ψ_ℓ represents the amplitude, rather than the probability, of having ℓ_i instances of each species $i = 1, \dots, k$. And in that context, a **quantum state** is a complex formal power series with

$$\sum_{\ell \in \mathbb{N}^k} |\psi_\ell|^2 = 1.$$

To obtain a Hilbert space, we often restrict attention to formal power series with

$$\sum_{\ell \in \mathbb{N}^k} |\psi_\ell|^2 < \infty.$$

However, power series not obeying this condition are also useful, for example to ensure that operators of interest are everywhere defined, instead of merely densely defined.

Thus, for the purposes of studying the master equation, we define the **stochastic Fock space** to be the vector space $\mathbb{R}[[z_1, \dots, z_k]]$ of all real formal power series in the variables z_1, \dots, z_k . We treat the Hamiltonian H for the master equation as the operator on stochastic Fock space with

$$H\Psi = \sum_{\ell, \ell' \in \mathbb{N}^k} H_{\ell'\ell} \psi_\ell z^{\ell'}$$

for all

$$\Psi = \sum_{\ell \in \mathbb{N}^k} \psi_\ell z^\ell,$$

where the matrix entries $H_{\ell'\ell}$ are given by Equation 4.

This allows us to express the Hamiltonian in terms of certain special operators on the stochastic Fock space: the creation and annihilation operators. Our notation here follows that used in quantum field theory, where the creation and annihilation operators are adjoints of each other. Let $1 \leq i \leq k$. The **creation operator** a_i^\dagger is given by

$$a_i^\dagger \Psi = z_i \Psi.$$

This takes any pure state to the pure state with one additional instance of the i th species:

$$a_i^\dagger (z_1^{n_1} \cdots z_i^{n_i} \cdots z_k^{n_k}) = z_1^{n_1} \cdots z_i^{n_i+1} \cdots z_k^{n_k}.$$

The corresponding **annihilation operator** is given by formal differentiation:

$$a_i \Psi = \frac{\partial}{\partial z_i} \Psi.$$

This takes any pure state to the pure state with one fewer instance of the i th species, but multiplied by a coefficient n_i :

$$a_i (z_1^{n_1} \cdots z_i^{n_i} \cdots z_k^{n_k}) = n_i z_1^{n_1} \cdots z_i^{n_i-1} \cdots z_k^{n_k}.$$

This represents the fact that there are n_i ways to annihilate one of the instances of the i th species present.

In what follows we use multi-index notation to define, for any $n \in \mathbb{N}^k$,

$$a^n = a_1^{n_1} \cdots a_k^{n_k}$$

and

$$a^{\dagger n} = a_1^{\dagger n_1} \cdots a_k^{\dagger n_k}.$$

These expressions are well-defined because all the annihilation operators commute with each other, and similarly for the creation operators.

Theorem 5. *For any stochastic reaction network, the Hamiltonian is given by*

$$H = \sum_{\tau \in T} r(\tau) \left(a^{\dagger t(\tau)} - a^{\dagger s(\tau)} \right) a^{s(\tau)}.$$

Proof. We start with a basic result on annihilation and creation operators:

Lemma 6. *For any multi-indices ℓ, m we have*

$$a^{\dagger m} z^\ell = z^{\ell+m}, \quad a^m z^\ell = \ell^{\underline{m}} z^{\ell-m}.$$

Proof. The first equation follows inductively from the definition of the creation operators, which implies

$$a_i^\dagger z^\ell = z_1^{\ell_1} \cdots z_i^{\ell_i+1} \cdots z_k^{\ell_k}.$$

The second follows inductively from the definition of annihilation operators, which implies

$$a_i z^\ell = \ell_i z_1^{\ell_1} \cdots z_i^{\ell_i-1} \cdots z_k^{\ell_k}.$$

Note that $\ell^{\underline{m}} = 0$ if $m_i > \ell_i$ for any i , and in this case we also have $a^m z^\ell = 0$, so the equation $a^m z^\ell = \ell^{\underline{m}} z^{\ell-m}$ holds in this case because both sides vanish. \square

We now prove the theorem. For any multi-index ℓ , Equation 4 gives

$$\begin{aligned} H z^\ell &= \sum_{\ell' \in \mathbb{N}^k} H_{\ell' \ell} z^{\ell'} \\ &= \sum_{\tau \in T} r(\tau) \ell^{s(\tau)} \left(z^{\ell + t(\tau) - s(\tau)} - z^\ell \right) \end{aligned}$$

Using Lemma 6 we obtain

$$H z^\ell = \sum_{\tau \in T} r(\tau) \left(a^{\dagger t(\tau)} - a^{\dagger s(\tau)} \right) a^{s(\tau)} z^\ell$$

and thus

$$H = \sum_{\tau \in T} r(\tau) \left(a^{\dagger t(\tau)} - a^{\dagger s(\tau)} \right) a^{s(\tau)}.$$

The last step requires a little justification. Technically speaking, the monomials z^ℓ are not a basis of the stochastic Fock space $\mathbb{R}[[z_1, \dots, z_k]]$, because not every formal power series is a *finite* linear combination of monomials. However, these monomials form a ‘topological basis’ of $\mathbb{R}[[z_1, \dots, z_k]]$, in the sense that every element of this vector space can be expressed as an *convergent infinite* linear combination of these monomials, with respect to a certain topology. (In this topology, a sequence of formal power series converges if each coefficient converges.) The annihilation and creation operators, and indeed all the operators discussed in this paper, are continuous in this topology. So, to check equations between these operators, it suffices to check them on the monomials z^ℓ . \square

Next we investigate what the master equation says about the time evolution of expected values. For this, we start by defining

$$\langle \Psi \rangle = \sum_{\ell \in \mathbb{N}^k} \psi_\ell$$

for any formal power series

$$\Psi = \sum_{\ell \in \mathbb{N}^k} \psi_\ell z^\ell.$$

In general the sum defining $\langle \Psi \rangle$ may not converge, but it converges and equals 1 when Ψ is a mixed state. Suppose O is a linear operator on the space of formal power series in the variables z_1, \dots, z_k . We define the **expected value** of O in the mixed state Ψ to be $\langle O \Psi \rangle$, assuming this converges. This generalizes the usual definition of the expected value of a random variable, in a way that emphasizes the analogy between stochastic mechanics and quantum mechanics. While in quantum mechanics we compute expected values of observables using expressions of the form $\langle \Psi, O \Psi \rangle$, where the brackets denote an inner product, in stochastic mechanics we use $\langle O \Psi \rangle$. For more details see [4, 5].

In applications of reaction networks, we are especially interested in how the number of items of each species changes with time. We can study this using the **number operators**:

$$N_i = a_i^\dagger a_i.$$

The expected value of N_i in a mixed state Ψ is given by

$$\langle N_i \Psi \rangle = \sum_{\ell \in \mathbb{N}^k} \ell_i \psi_\ell$$

Since ψ_ℓ is the probability of being in a pure state where there are ℓ_i things of the i th species, the above formula indeed gives the expected value of the number of things of this species.

Let us compute the rate of change

$$\frac{d}{dt}\langle N_i \Psi(t) \rangle$$

assuming that $\Psi(t)$ obeys the master equation. For this, we need to define the falling powers of an operator A as follows:

$$A^{\underline{p}} = A(A-1)\cdots(A-p+1)$$

for any $p \in \mathbb{N}$. If m is a multi-index we define

$$N^{\underline{m}} = N_1^{\underline{m_1}} \cdots N_k^{\underline{m_k}}.$$

Since the number operators N_i commute, it does not matter in what order we write this product.

Lemma 7. *For any multi-indices ℓ and m we have*

$$N^{\underline{m}} z^{\ell} = \ell^{\underline{m}} z^{\ell}.$$

Proof. By Lemma 6 and the definition of the number operators we have

$$N_i z^{\ell} = \ell_i z^{\ell}$$

for all $1 \leq i \leq k$. The lemma follows directly from this. \square

There is a way to extract a classical state from a mixed state, which lets us connect the rate equation to the master equation. To do this, we write N for the vector, or list, of number operators (N_1, \dots, N_k) . Then, given a mixed state Ψ , we set

$$\langle N \Psi \rangle = (\langle N_1 \Psi \rangle, \dots, \langle N_k \Psi \rangle).$$

If the expected values here are well-defined, $\langle N \Psi \rangle \in [0, \infty)^k$ is a classical state. It is a simplified description of the mixed state Ψ , which only records the expected number of instances of each species.

The next result says how this sort of classical state evolves in time, assuming that the mixed state it comes from obeys the master equation:

Theorem 8. *For any stochastic reaction network and any mixed state $\Psi(t)$ evolving in time according to the master equation, we have*

$$\frac{d}{dt}\langle N \Psi(t) \rangle = \sum_{\tau \in T} r(\tau) (s(\tau) - t(\tau)) \left\langle N^{\underline{s(\tau)}} \Psi(t) \right\rangle$$

assuming the expected values and their derivatives exist.

Proof. First note using Lemmas 6 and 7 that for any multi-indices ℓ, m, n and any natural number i we have

$$\begin{aligned} \langle N_i a^{\dagger m} a^n z^{\ell} \rangle &= \ell^{\underline{n}} \langle N_i a^{\dagger m} z^{\ell-n} \rangle \\ &= \ell^{\underline{n}} \langle N_i z^{\ell-n+m} \rangle \\ &= \ell^{\underline{n}} (\ell_i - n_i + m_i). \end{aligned}$$

Thus if at any given time

$$\Psi(t) = \sum_{\ell \in \mathbb{N}^k} \psi_{\ell}(t) z^{\ell}$$

and $\Psi(t)$ evolves in time according to the master equation, we have

$$\begin{aligned}
\frac{d}{dt}\langle N_i \Psi(t) \rangle &= \langle N_i H \Psi(t) \rangle \\
&= \sum_{\tau \in T} \sum_{\ell \in \mathbb{N}^k} r(\tau) \left\langle N_i \left(a^{\dagger t(\tau)} - a^{\dagger s(\tau)} \right) a^{s(\tau)} \psi_\ell(t) z^\ell \right\rangle \\
&= \sum_{\tau \in T} \sum_{\ell \in \mathbb{N}^k} r(\tau) \ell^{s(\tau)} (s_i(\tau) - t_i(\tau)) \psi_\ell(t) \\
&= \sum_{\tau \in T} \sum_{\ell \in \mathbb{N}^k} r(\tau) (s_i(\tau) - t_i(\tau)) \left\langle N^{s(\tau)} z^\ell \right\rangle \psi_\ell(t) \\
&= \sum_{\tau \in T} r(\tau) (s_i(\tau) - t_i(\tau)) \left\langle N^{s(\tau)} \Psi(t) \right\rangle
\end{aligned}$$

Recall that N is our notation for the vector of number operators (N_1, \dots, N_k) . Thus, the above equation is equivalent to

$$\frac{d}{dt}\langle N \Psi(t) \rangle = \sum_{\tau \in T} r(\tau) (s(\tau) - t(\tau)) \left\langle N^{s(\tau)} \Psi(t) \right\rangle \quad \square$$

The above theorem makes clear that the rate equation is closely related to the master equation, since the former says

$$\frac{d}{dt}x(t) = \sum_{\tau \in T} r(\tau) (t(\tau) - s(\tau)) x(t)^{s(\tau)}$$

while the latter implies

$$\frac{d}{dt}\langle N \Psi(t) \rangle = \sum_{\tau \in T} r(\tau) (s(\tau) - t(\tau)) \left\langle N^{s(\tau)} \Psi(t) \right\rangle.$$

Suppose we let $x(t)$ be the classical state coming from the mixed state $\Psi(t)$:

$$x(t) = \langle N \Psi(t) \rangle$$

The rate equation for $x(t)$ would then follow from the master equation for $\Psi(t)$ if we had

$$\langle N \Psi(t) \rangle^m = \langle N^m \Psi(t) \rangle$$

for every multi-index m . This equation is not true in general. However, it should hold *approximately* in a suitable limit of large numbers. And surprisingly, it holds *exactly* for coherent states.

6 Coherent states

For any $c \in [0, \infty)^k$, we define the **coherent state** with expected value c to be the mixed state

$$\Psi_c = \frac{e^{c \cdot z}}{e^c}$$

where $c \cdot z$ is the dot product of c and the vector of variables $z = (z_1, \dots, z_k)$, and we set $e^c = e^{c_1 + \dots + c_k}$. Equivalently,

$$\Psi_c = \frac{1}{e^c} \sum_{n \in \mathbb{N}^k} \frac{c^n}{n!} z^n,$$

where c^n and z^n are defined as products in our usual way, and $n! = n_1! \cdots n_k!$. The name ‘coherent state’ comes from quantum mechanics [11], where we think of the coherent state Ψ_c as the quantum state that best approximates the classical state c . In the state Ψ_c , the probability of having n_i things of the i th species is equal to

$$e^{-c_i} \frac{c_i^{n_i}}{n_i!}.$$

This is precisely the definition of a Poisson distribution with mean equal to c_i . The state Ψ_c is thus a product of independent Poisson distributions.

Theorem 9. *Given any stochastic reaction network, let $\Psi(t)$ be a mixed state evolving in time according to the master equation. If $\Psi(t)$ is a coherent state when $t = t_0$, then*

$$x(t) = \langle N\Psi(t) \rangle$$

obeys the rate equation when $t = t_0$.

Proof. We prove this using a series of lemmas:

Lemma 10. *For any multi-index m and any coherent state Ψ_c we have*

$$a^m \Psi_c = c^m \Psi_c$$

and

$$a^{\dagger m} \Psi_c = z^m \Psi_c.$$

Proof. The second equation is immediate from the definition, while the first follows from

$$a_i \Psi_c = \frac{\partial}{\partial z_i} \frac{e^{c \cdot z}}{e^c} = c_i \Psi_c. \quad \square$$

Lemma 11. *For any multi-index m we have*

$$N^{\underline{m}} = a^{\dagger m} a^m.$$

Proof. For any multi-index ℓ , Lemmas 6 and 7 imply

$$a^{\dagger m} a^m z^\ell = \ell^{\underline{m}} a^{\dagger m} z^{\ell-m} = \ell^{\underline{m}} z^\ell = N^{\underline{m}} z^\ell.$$

Since the states z^ℓ form a topological basis of $\mathbb{R}[[z_1, \dots, z_k]]$, and all the operators in question are continuous, the lemma follows. \square

Lemma 12. *For any multi-index m and any $\Psi \in \mathbb{R}[[z_1, \dots, z_k]]$ we have*

$$\langle a^{\dagger m} \Psi \rangle = \langle \Psi \rangle$$

Proof. We have

$$\langle a^{\dagger m} \Psi \rangle = \left\langle \sum_{\ell \in \mathbb{N}^k} \psi_\ell z^{\ell+m} \right\rangle = \sum_{\ell \in \mathbb{N}^k} \psi_\ell = \langle \Psi \rangle. \quad \square$$

Lemma 13. *For any multi-index m and any coherent state Ψ_c we have*

$$\langle N^{\underline{m}} \Psi_c \rangle = \langle N \Psi_c \rangle^m$$

Proof. By Lemma 11 and Lemma 10 we have

$$N^m \Psi_c = a^{\dagger m} a^m \Psi_c = c^m a^{\dagger m} \Psi_c$$

so by Lemma 12

$$\langle N^m \Psi_c \rangle = \langle c^m a^{\dagger m} \Psi_c \rangle = c^m \langle \Psi_c \rangle = c^m.$$

As a special case we have $\langle N_i \Psi_c \rangle = c_i$, so we also have

$$\langle N \Psi_c \rangle^m = c^m$$

and it follows that $\langle N^m \Psi_c \rangle = \langle N \Psi \rangle^m$. □

To prove Theorem 9, we start by setting

$$x(t) = \langle N \Psi(t) \rangle$$

where $\Psi(t)$ is a solution of the master equation. Theorem 8 implies that the time derivative of x equals

$$\dot{x}(t) = \sum_{\tau \in T} r(\tau) (s(\tau) - t(\tau)) \langle N^{s(\tau)} \Psi(t) \rangle.$$

If $\Psi(t)$ is a coherent state at time $t = t_0$, Lemma 13 implies that

$$\langle N^{s(\tau)} \Psi(t_0) \rangle = \langle N \Psi(t_0) \rangle^{s(\tau)} = x(t_0)^{s(\tau)}$$

so we have

$$\dot{x}(t_0) = \sum_{\tau \in T} r(\tau) (s(\tau) - t(\tau)) x(t_0)^{s(\tau)}$$

which is the rate equation at time t_0 , as desired. □

In general, the above result applies at only one moment in time. The reason is that if a solution of the master equation is a coherent state at some time, it need not be a coherent state at later (or earlier) times. Still, Theorem 8 implies that as long as the expected values $\langle N^m \Psi(t) \rangle$ are *close* to the powers $\langle N \Psi(t) \rangle^m$, at least when m is the source of some reaction, the expected values $x(t) = \langle N \Psi(t) \rangle$ will *approximately* obey the rate equation. This could be studied in detail using either the techniques introduced here or those discussed by Anderson and Kurtz [2].

Furthermore, in some cases when $\Psi(t)$ is initially a coherent state it continues to be so for all times. In this case, $x(t)$ continues to exactly obey the rate equation. For example, this is true when all the complexes in the reaction network consist of a single species. It is also true for a large class of equilibrium solutions of the master equation, as shown by Anderson, Craciun and Kurtz [1]. For a discussion of how quantum techniques apply to this situation, see the companion to this paper [6].

Acknowledgements

I thank Jacob Biamonte, Brendan Fong, and the students at U. C. Riverside who helped come up with a preliminary version of the proof of Theorem 8, notably Daniel Estrada, Reeve Garrett, Michael Knap, Tu Pham, Blake Pollard and Franciscus Rebro. I thank John Rowlands and Blake Stacey for catching typos and other mistakes. I also thank the Centre of Quantum Technology and the Mathematics Department of U. C. Riverside for their support.

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