



**Statistical
optimization for
passive scalar
transport**

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Statistical optimization for passive scalar transport: maximum entropy production vs. maximum Kolmogorov–Sinay entropy

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Abstract

We derive rigorous results on the link between the principle of maximum entropy production and the principle of maximum Kolmogorov–Sinai entropy using a Markov model of the passive scalar diffusion called the Zero Range Process. We show analytically that both the entropy production and the Kolmogorov–Sinai entropy seen as functions of f admit a unique maximum denoted $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$. The behavior of these two maxima is explored as a function of the system disequilibrium and the system resolution N . The main result of this article is that $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$ have the same Taylor expansion at first order in the deviation of equilibrium. We find that $f_{\max_{\text{EP}}}$ hardly depends on N whereas $f_{\max_{\text{KS}}}$ depends strongly on N . In particular, for a fixed difference of potential between the reservoirs, $f_{\max_{\text{EP}}}(N)$ tends towards a non-zero value, while $f_{\max_{\text{KS}}}(N)$ tends to 0 when N goes to infinity. For values of N typical of that adopted by Paltridge and climatologists ($N \approx 10 \sim 100$), we show that $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$ coincide even far from equilibrium. Finally, we show that one can find an optimal resolution N^* such that $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$ coincide, at least up to a second order parameter proportional to the non-equilibrium fluxes imposed to the boundaries. We find that the optimal resolution N^* depends on the non equilibrium fluxes, so that deeper convection should be represented on finer grids. This result points to the inadequacy of using a single grid for representing convection in climate and weather models. Moreover, the application of this principle to passive scalar transport parametrization is therefore expected to provide both the value of the optimal flux, and of the optimal number of degrees of freedom (resolution) to describe the system.

1 Introduction

A major difficulty in the modeling of nonlinear geophysical or astrophysical processes is the taking into account of all the relevant degrees of freedom. For example, fluid motions obeying Navier–Stokes equations usually require of the order of $N = Re^{9/4}$

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5 modes to faithfully describe all scales between the injection scale and the dissipative scale (Frisch, 1995). In atmosphere, or ocean, where the Reynolds number exceeds 10^9 , this amounts to $N = 10^{20}$, a number too large to be handled by any existing computers (Wallace and Hobbs, 2006). The problem is even more vivid in complex systems

10 such as planetary climate, where the coupling of lito-bio-cryo-sphere with ocean and atmosphere increases the number of degrees of freedom beyond any practical figure. This justifies the long historical tradition of parametrization and statistical model reduction, to map the exact equations describing the system onto a set of simpler equations involving few degrees of freedom. The price to pay is the introduction of *free parameters*, describing the action of discarded degrees of freedom, that needs to be prescribed.

When the number of free parameters is small, their prescription can be successfully done empirically through calibrating experiments or by a posteriori tuning (Rotstayn, 2000). When the number of parameters is large, such as in climate models where it reaches several hundreds (Yang et al., 2012), such empirical procedure is inapplicable, because it is impossible to explore the whole parameter space. In that respect, it is of great interest to explore the alternate road to parametrization via application of a statistical optimization principle, such as minimizing or maximizing of a suitable cost functional. As discussed by Turkington (2013), this strategy usually leads to closed reduced equations with adjustable parameters in the closure appearing as weights in the cost functional and can be computed explicitly. A famous example in climate is given by a principle of maximum entropy production (MEP) that allowed Paltridge (1975) to derive the distribution of heat and clouds at the Earth surface with reasonable accuracy, without any parameters and with a model of a dozen of degrees of freedom (boxes). Since then, refinements of Paltridge model have been suggested to increase its generality and range of prediction (Herbert et al., 2011). Rigorous justifications of its application have been searched using e.g. information theory (Dewar, 2003) without convincing success. More recently, we have used the analogy of the climate box model of Paltridge with the asymmetric exclusion Markov process (ASEP) to establish

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numerically a link between the MEP and the principle of maximum Kolmogorov–Sinai entropy (MKS) (Mihelich et al., 2014). The MKS principle is a relatively new concept which extends the classical results of equilibrium physics Monthus (2011). This principle applied to Markov Chains provides an approximation of the optimal diffusion coefficient in transport phenomena (Gómez-Gardeñes and Latora, 2008) or simulates random walk on irregular lattices (Burda et al., 2009). It is therefore a good candidate for a physically relevant cost functional in passive scalar modeling.

The goal of the present paper is to derive rigorous results on the link between MEP and MKS using a Markov model of the passive scalar diffusion called the Zero Range Process (Andjel, 1982). We find that there exists an optimal resolution N_* such that both maxima coincide to second order in the distance to equilibrium. The application of this principle to passive scalar transport parametrization is therefore expected to provide both the value of the optimal flux, and of the optimal number of degrees of freedom (resolution) to describe the system. This suggests that the MEP and MKS principle may be unified when the Kolmogorov–Sinai entropy is defined on opportunely coarse grained partitions.

2 From passive scalar equation to ZRP model

The equation describing the transport of a passive scalar like temperature in a given velocity field $u(x, t)$ reads:

$$\partial_t T + u \nabla T = \kappa \Delta T, \quad (1)$$

with appropriate boundary conditions. Here κ is the diffusivity. To solve this equation, one must know both the velocity field and the boundary conditions, and use as many number of modes as necessary to describe all range of scales up to the scales at which molecular diffusivity takes place i.e. roughly $(RePr)^{3/2}$ modes, where Re is the Reynolds number of the convective flow, and Pr is its Prandtl number. In geophysical flows, this number is too large to be handled even numerically (Troen and Mahrt, 1986).

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Moreover, in typical climate studies, the velocity flow is basically unknown as it must obey a complicated equation involving the influence of all the relevant climate components. In order to solve the equation, one must necessarily prescribe the heat flux $f = -u + \kappa \nabla T$. The idea of Paltridge was then to discretize the passive scale equation in boxes and prescribe the heat flux f_{ij} between boxes i and j by maximizing the associated thermodynamic entropy production $\dot{S} = \sum_i f_{ij}/T_i$. Here, we slightly modify the Paltridge discretization approximation to make it amenable to rigorous mathematical results on Markov Chains. For simplicity, we stick to a one dimensional case (corresponding to boxes varying only in latitude) and impose the boundary conditions through two reservoirs located at each end of the chain (mimicking the solar heat flux at pole and equator). We consider a set of N boxes that can contain an arbitrary number $n \in \mathbb{N}$ of particles. We then allow transfer of particles in between two adjacent boxes via decorrelated jumps (to the right or to the left) following a 1-D Markov dynamics governed by a coupling with the two reservoirs imposing a difference of chemical potential at the ends. The resulting process is called the Zero Range Process (Andjel, 1982). The different jumps are described as follow. At each time step a particle can jump right with probability $p w_n$ or jump left with probability $q w_n$. w_n is a parameter depending of the number of particles inside the box. Physically it represents the interactions between particles. At the edges of the lattice the probability rules are different: At the left edge a particle can enter with probability α and exit with probability γw_n whereas at the right edge a particle can exit with probability βw_n and enter with probability δ . Choices of different w_n give radically different behaviors. For example $w_n = 1 + b/n$ where $b \geq 0$ described condensation phenomena (Großkinsky et al., 2003) whereas $w_1 = w$ and $w_n = 1$ if $n \geq 2$ has been used to modeled road traffic. We will consider in this article the particular case where $w = 1$ by convenience of calculation. Moreover without loss of generality we will take $p \geq q$ which corresponds to a particle flow from the left to the right and note $f = p - q$. After a sufficiently long time the system reaches a non-equilibrium steady state. The interest of this toy model is that it is simple enough so that exact computations are analytically tractable.

Taking the continuous limit of this process, it may be checked that the fugacity z , which is related to the average particle density (see definition below), of stationary solutions of a system consisting of boxes of size $\frac{1}{N}$ follows the continuous equation (Levine et al., 2005) :

$$f \frac{\partial z}{\partial x} - \frac{1}{2N} \frac{\partial^2 z}{\partial x^2} = 0, \quad (2)$$

corresponding to stationary solution of a passive scalar equation with velocity f and diffusivity $\frac{1}{2N}$. Therefore, the fugacity of the Zero Range Process is a passive scalar obeying a convective-diffusion equation. We thus see that $f = 0$ corresponds to a purely conductive regime whereas the larger f the more convective the regime. In the sequel, we calculate the entropy production and the Kolmogorov–Sinai entropy function of f . These two quantities reach a maximum noted respectively $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$. The MEP principle (resp. the MKS principle) states that the system will choose $f = f_{\max_{\text{EP}}}$ (resp. $f = f_{\max_{\text{KS}}}$).

We will show first of all in this article that numerically $f_{\max_{\text{EP}}} \approx f_{\max_{\text{KS}}}$ even far from equilibrium for a number of boxes N roughly corresponding to the resolution taken by Paltridge (1975) in his climate model. This result is similar to what we found for the ASEP model (Mihelich et al., 2014) and thus gives another example of a system in which the two principles are equivalent. Moreover we will see analytically that $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$ have the same behavior in first order in the difference of the chemical potentials between the two reservoirs for N large enough. These results provide a better understanding of the relationship between the MEP and the MKS principles.

3 Notations and useful preliminary results

This Markovian Process is a stochastic process with a infinite number of states in bijection with \mathbb{N}^N . In fact, each state can be written $n = (n_1, n_2, \dots, n_N)$ where n_i is the number of particle lying in site i . We call P_n the stationary probability to be in state

n . In order to calculate this probability it is easier to use a quantum formalism than the Markovian formalism as explain in the following articles (Domb, 2000; Levine et al., 2005).

The probability to find m particles in the site k is equal to: $p_k(n_k = m) = \frac{z_k^m}{Z_k}$ where Z_k is the analogue of the grand canonical repartition function and z_k is the fugacity between 0 and 1. Moreover $Z_k = \sum_{i=0}^{\infty} z_k^i = \frac{1}{1-z_k}$. So, finally

$$p_k(n_k = m) = (1 - z_k)z_k^m. \quad (3)$$

We can show that the probability P over the states is the tensorial product of the probability p_k over the boxes:

$$P = p_1 \otimes p_2 \otimes \dots \otimes p_N.$$

Thus events $(n_k = m)$ and $(n_{k'} = m')$ for $k \neq k'$ are independent and so:

$$P(m_1, m_2, \dots, m_N) = p_1(n_1 = m_1) \cdot \dots \cdot p_N(n_N = m_N). \quad (4)$$

So finally

$$P(m_1, m_2, \dots, m_N) = \prod_{k=1}^N (1 - z_k)z_k^{m_k}. \quad (5)$$

Moreover, with the Hamiltonian equation we can find the exact values of z_k function of the system parameters:

$$z_k = \frac{\left(\frac{p}{q}\right)^{k-1} [(\alpha + \delta)(p - q) - \alpha\beta + \gamma\delta] - \gamma\delta + \alpha\beta \left(\frac{p}{q}\right)^{N-1}}{\gamma(p - q - \beta) + \beta(p - q + \gamma) \left(\frac{p}{q}\right)^{N-1}}, \quad (6)$$

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and the flux of particles c :

$$c = (\rho - q) \frac{-\gamma\delta + \alpha\beta\left(\frac{\rho}{q}\right)^{N-1}}{\gamma(\rho - q - \beta) + \beta(\rho - q + \gamma)\left(\frac{\rho}{q}\right)^{N-1}}. \quad (7)$$

Finally, the stationary density is related to the fugacity by the relation:

$$\rho_k = z_k \frac{\partial \log Z_k}{\partial z_k} = \frac{z_k}{1 - z_k}. \quad (8)$$

3.1 Entropy production

For a system subject to internal forces X_i and associated fluxes J_i the macroscopic entropy production is well known (Balian, 1992) and takes the form:

$$\sigma = \sum_i J_i \cdot X_i.$$

The Physical meaning of this quantity is a measure of irreversibility: the larger σ the more irreversible the system.

In the case of the zero range process irreversibility is created by the fact that $\rho \neq q$. We will parametrize this irreversibility by the parameter $f = \rho - q$ and we will take $\rho + q = 1$. In the remaining of the paper, we take, without loss of generality, $\rho \leq q$ which corresponds to a flow from right to left. Moreover, the only flux to be considered is here the flux of particles c and the associated force is due to the gradient of the density of particles ρ : $X = \nabla \log \rho$.

Thus, when the stationary state is reached when c is constant:

$$\sigma = \sum_{i=1}^{N-1} c \cdot (\log(\rho_i) - \log(\rho_{i+1})) = c \cdot (\log(\rho_1) - \log(\rho_N)). \quad (9)$$

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Thus, according to Eqs. (6)–(9) when N tends to $+\infty$ we obtain:

$$\sigma(f) = \frac{\alpha f}{f + \gamma} \left(\log \left(\frac{\alpha}{f + \gamma - \alpha} \right) - \log \left(\frac{(\alpha + \delta)f + \gamma\delta}{f(\beta - \alpha - \delta) + \beta\gamma - \gamma\delta} \right) \right). \quad (10)$$

Because $f \geq 0$ the entropy production is positive if and only if $\rho_1 \geq \rho_N$ if $z_1 \geq z_N$. This is physically coherent because fluxes are in the opposite direction of the gradient. We remark that if $f = 0$ then $\sigma(f) = 0$. Moreover, when f increases $\rho_1(f)$ decreases and $\rho_2(f)$ increases till they take the same value. Thus it exists f , large enough, for which $\sigma(f) = 0$. Between these two values of f the entropy production has at least one maximum.

3.2 Kolmogorov–Sinai entropy

There are several ways to introduce the Kolmogorov–Sinai entropy which is a mathematical quantity introduced by Kolmogorov and picked up by famous mathematician as Sinai and Billingsley (Billingsley, 1965). Nevertheless, for a Markov process we can give it a simple physic interpretation: the Kolmogorov–Sinai entropy is the time derivative of the Jaynes entropy (entropy over the path).

$$S_{\text{Jaynes}}(t) = - \sum_{\Gamma_{[0,t]}} \rho_{\Gamma_{[0,t]}} \cdot \log \left(\rho_{\Gamma_{[0,t]}} \right). \quad (11)$$

For a Markov Chain we have thus:

$$S_{\text{Jaynes}}(t) - S_{\text{Jaynes}}(t - 1) = - \sum_{(i,j)} \mu_{i,\text{stat}} \rho_{ij} \log(\rho_{ij}), \quad (12)$$

where $\mu_{\text{stat}} = \mu_{i,\text{stat}}$ $i = 1 \dots N$ is the stationary measure and where the ρ_{ij} are the transition probabilities.

Thus the Kolmogorov–Sinai entropy takes the following form:

$$h_{\text{KS}} = - \sum_{(i,j)} \mu_{i,\text{stat}} \rho_{ij} \log(\rho_{ij}). \quad (13)$$

For the Zero Range Process, we show in appendix that it can be written as:

$$\begin{aligned}
 h_{\text{KS}} = & -(\alpha \log \alpha + \delta \log \delta + \gamma \log \gamma + \beta \log \beta + (N - 1)(\rho \log(\rho) + q \log(q))) \\
 & + (\rho \log(\rho) + q \log(q)) \sum_{i=1}^N (1 - z_i) + (\gamma \log(\gamma) + \rho \log(\rho))(1 - z_1) \\
 & + (\beta \log(\beta) + q \log(q))(1 - z_N). \tag{14}
 \end{aligned}$$

4 Results

We will start first by pointing to some interesting properties of f_{maxEP} and f_{maxKS} , then by presenting numerical experiments on the ZRP model and finally concluding with some analytical computations.

Let us first note that for $N, \alpha, \beta, \gamma, \delta$ fixed the entropy production as well as the Kolmogorov–Sinai entropy seen as functions of f admit both a unique maximum. When N tends to infinity and $f = 0$, using Eq. (6) (i.e. the symmetric case), we find that $z_1 = \frac{\alpha}{\gamma}$ and $z_N = \frac{\delta}{\beta}$. Thus, the system is coupled with two reservoirs with respective chemical potential $\frac{\alpha}{\gamma}$ (left) and $\frac{\delta}{\beta}$ (right). For $\frac{\alpha}{\gamma} \neq \frac{\delta}{\beta}$ the system is out of equilibrium. We assume, without loss of generality, $z_1 \geq z_N$ which corresponds to a flow from left to right. As a measure of deviation from equilibrium we take $s = z_1 - z_N$: the larger s , the more density fluxes we expect into the system.

First we remark that f_{maxEP} hardly depends on N whereas f_{maxKS} depends strongly on N . This is easily understood because σ depends only on z_1 and z_N whereas h_{KS} depends on all the z_i . Moreover, the profile of the z_i depends strongly on N . In particular, for a fixed difference of potential between the reservoirs, $f_{\text{maxEP}}(N)$ tends towards a non-zero value, while $f_{\text{maxKS}}(N)$ tends to 0 when N goes to infinity.

Moreover, f_{maxEP} and f_{maxKS} coincide even far from equilibrium for N corresponding to the choice of Paltridge (1975) $N \approx 10 \sim 100$. for N fixed, as large as one wants, and for all ϵ , as small as one wants, it exists ν such that for all $s \in [0; \nu]$ $|f_{\text{maxEP}} - f_{\text{maxKS}}| \leq \epsilon$.

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These observations are confirmed by the results presented in Fig. 1 where EP and KS are calculated using Eqs. (6) and (14) for $s = 0.13$ and three different partitions: $N = 20$, $N = 100$ and $N = 1000$. The figure shows that $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$ coincide with good approximation for $N = 20$ and $N = 100$. But then when N increases $f_{\max_{\text{KS}}}(N)$ tends to 0 whereas $f_{\max_{\text{EP}}}(N)$ tends to a non-zero value.

In Fig. 2 we represent the Entropy Production (top) and KS Entropy (bottom) function of f for $N = 1000$ and for three value of s : $s = 0.13$; $s = 0.2$; $s = 0.04$. This supports the claim that for N fixe, we could tried different values of s such that $s \in [0; \nu]$ $|f_{\max_{\text{EP}}} - f_{\max_{\text{KS}}}| \leq \epsilon$.

Such numerical investigations suggest to understand why $f_{\max_{\text{KS}}}(N)$ and $f_{\max_{\text{EP}}}(N)$ have different behaviors function of N , and why for N large enough $f_{\max_{\text{KS}}}$ and $f_{\max_{\text{EP}}}$ have the same behavior of first order in the deviation from equilibrium measured by the parameter s . We will see that we can get a precise answer to such questions by doing calculations and introducing a sort of Hydrodynamics approximation.

Taylor expansion

From Eq. (14) it is apparent that $f_{\max_{\text{KS}}}$ depends on N whereas from Eq. (9) we get that $f_{\max_{\text{EP}}}$ hardly depends on N . Indeed there is a difference between $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$, i.e. a difference between the two principles for the Zero Range Process. Nevertheless, we have seen numerically that there is a range of N , namely $N \approx 10 \sim 100$ for which the maxima fairly coincide.

Using Eqs. (14), (6) and (10) we compute analytically the Taylor expansion of $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$ in s . We will show the main result: $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$ have the same Taylor expansion in first order in s for N large enough. Their Taylor expansions are different up to the second order in s but it exists an N , i.e. a resolution, such that $f_{\max_{\text{EP}}}$ and $f_{\max_{\text{KS}}}$ coincident up to the second order.

Let us start by compute $f_{\max_{\text{KS}}}$. It is not depends of the constant terms of h_{KS} in Eq. (14) and therefore we need only concern ourselves with :

$$-(p \log(p) + q \log(q)) \left(\sum_{i=1}^N (z_i) - 1 \right) + (\gamma \log(\gamma) + p \log(p))(1 - z_1) + (\beta \log(\beta) + q \log(q))(1 - z_N) = N \cdot H(f, N, \alpha, \gamma, \beta, \delta). \quad (15)$$

5 Using Eq. (6), the expression of $H(f, N, \alpha, \gamma, \beta, \delta)$ takes an easy form. To simplify the calculations, we restrict the space of parameter by assuming $\alpha + \gamma = 1$ and $\beta + \delta = 1$ and we parametrize the deviation from equilibrium by the parameter $\bar{s} = \alpha - \delta$. Moreover let's note $a = \frac{1}{N}$. Thus, we have $H(f, N, \alpha, \gamma, \beta, \delta) = H(f, a, \alpha, \bar{s})$. In order to know the Taylor expansion to the first order in \bar{s} of $f_{\max_{\text{KS}}}$ we develop $H(f, a, \alpha, \bar{s})$ up to the second order in f ; i.e. we have $H(f, a, \alpha, \bar{s}) = C + Bf + Af^2 + o(f^2)$ then we find $f_{\max_{\text{KS}}} = -B/2A$ that we will develop in power of \bar{s} . This is consistent if we assume $f \ll a$.

After some tedious but straightforward calculations, we get at the first order in \bar{s}

$$f_{\max_{\text{KS}}}(\bar{s}) = \frac{1}{4} \frac{(1 - \alpha) - a(\alpha + 2)}{\alpha(1 - \alpha) + 2a\alpha(\alpha - 1)} \bar{s} + o(\bar{s}). \quad (16)$$

And so,

$$15 \quad f_{\max_{\text{KS}}}(\bar{s}) = \frac{1}{4\alpha} \bar{s} + \frac{3a}{4(\alpha - 1)} \bar{s} + o(\bar{s}) + o(a\bar{s}). \quad (17)$$

We repeat the same procedure starting from Eq. (10) and we obtain:

$$f_{\max_{\text{EP}}}(\bar{s}) = \frac{\bar{s}}{4\alpha} + o(\bar{s}) + o(a). \quad (18)$$

Thus, since $a = \frac{1}{N} \ll 1$ the behaviour of $f_{\max_{\text{KS}}}(\bar{s})$ and $f_{\max_{\text{EP}}}(\bar{s})$ is the same for \bar{s} small enough.

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We remark that we can strictly find the same result by solving the hydrodynamics continuous approximation given by Eq. (2). This equation is a classical convection-diffusion equation. We remark that, by varying f , we change the convective behavior: $f = 0$ corresponds to a purely diffusive regime whereas by increasing f we enhance the role of convection. If the system is at equilibrium then $f_{\max_{EP}} = f_{\max_{KS}} = 0$ and the system is purely diffusive. When the system is out of equilibrium $f_{\max_{EP}}$ and $f_{\max_{KS}}$ are different than 0 and corresponds to an (optimal) trade-off between purely diffusive and convective behavior.

One can verify this numerically: We first calculate the exact values of the Entropy Production function of f using Eq. (6) and the Kolmogorov–Sinai Entropy function of f using Eqs. (6) and (14). Then we approximate these two curves with a cubic spline approximation in order to find $f_{\max_{EP}}$ and $f_{\max_{KS}}$.

In order to find the optimal resolution N_* we can go one step further by expanding $f_{\max_{EP}}$ and $f_{\max_{KS}}$ up to the second order in \bar{s} :

$$f_{\max_{EP}}(\bar{s}) = \frac{\bar{s}}{4\alpha} + \frac{\bar{s}^2(\alpha + 1)}{8\alpha^2(\alpha - 1)} + o(\bar{s}^2) + o(a). \quad (19)$$

$$f_{\max_{KS}}(\bar{s}) = \frac{1}{4} \frac{(1 - \alpha) - a(\alpha + 2)}{\alpha(1 - \alpha) + 2a\alpha(\alpha - 1)} \bar{s} + \frac{(1 - \alpha)^2 + a(\alpha^2 - 2\alpha + 1)}{8\alpha^2(\alpha - 1)^2(1 - 2a)} \bar{s}^2 + o(\bar{s}^2). \quad (20)$$

Thus, $f_{\max_{EP}}$ and $f_{\max_{KS}}$ coincide in second order in \bar{s} if a satisfies the equation of second degrees:

$$(4\alpha - 6\alpha^2 + 6\alpha^3 - 4\bar{s} + 3\alpha^2\bar{s})a^2 - \frac{1}{2}(8\alpha - 8\bar{s} + 3\alpha^2\bar{s} - 6\alpha^2 + 6\alpha^3)a - (1 - \alpha). \quad (21)$$

This equation has a unique positive solution because the dominant coefficient is positive for s small enough $(4\alpha - 6\alpha^2 + 6\alpha^3 - 4\bar{s} + 3\alpha^2\bar{s}) \geq 0$ and the constant term is negative $-(1 - \alpha) \leq 0$. We remark that the optimal resolution $N_* = \frac{1}{a_*}$ depends on the parameters of the system namely on the degree of non-equilibrium. This fact can be the explanation for two well known issues in climate/weather modeling. First, it explains that,

to be represented, points to new interesting research patterns, e.g. the introduction of convective models with adaptive grids optimized with maximum entropy principles on the basis of the convective phenomena to be represented.

On another hand, the application of this principle to passive scalar transport parametrization is therefore expected to provide both the value of the optimal flux, and of the optimal number of degrees of freedom (resolution) to describe the system. It would be interesting to apply it to more realistic passive scalar transport problem, to see if it yield to model that can be numerically handled (i.e. corresponding to a number of bow that is small enough to be handled by present computers). Moreover, on a theoretical side, it will be interesting to study whether for general dynamical systems, there exists a smart way to coarse grain the Kolmogorov–Sinai entropy such that its properties coincide with the thermodynamic entropy production. This will eventually justify the use of the MEP principle and explain the deviations as well as the different representations of it due to the dependence of the dynamic (Kolmogorov–Smirnov, Tsallis, Jaynes) entropies on the kind of partition adopted.

Appendix A: Computation of the K–S entropy

In this appendix, we compute the Kolmogorov–Sinai entropy for the Zero Range Process, starting from its definition Eq. (13). In the frame of our Zero Range Process, we use Eqs. (13) and (5) to write it as:

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$$\begin{aligned}
 h_{\text{KS}} &= - \sum_i \mu_{i,\text{stat}} \sum_j p_{ij} \log(p_{ij}) \\
 &= - \sum_{m_1=0}^{+\infty} \dots \sum_{m_N=0}^{+\infty} P(m_1, m_2, \dots, m_N) \sum_j p_{(m_1, \dots, m_N) \rightarrow j} \log(p_{(m_1, \dots, m_N) \rightarrow j}) \\
 &= - \sum_{m_1=0}^{+\infty} P(m_1) \dots \sum_{m_N=0}^{+\infty} P(m_N) \sum_j p_{(m_1, \dots, m_N) \rightarrow j} \log(p_{(m_1, \dots, m_N) \rightarrow j}). \tag{A1}
 \end{aligned}$$

We thus have to calculate $\sum_j p_{(m_1, \dots, m_N) \rightarrow j} \log(p_{(m_1, \dots, m_N) \rightarrow j})$ that we will refer to as (*). We will take $p + q = \alpha + \delta = \beta + \gamma = 1$ and $dt = \frac{1}{N}$ in order to neglect the probabilities to stay in the same state compare to the probabilities of changing state. There are five different cases to consider:

1. If $\forall i m_i \geq 1$ so the possible transitions are:

$(m_1, m_2, \dots, m_N) \rightarrow (m_1 \pm 1, m_2, \dots, m_N)$ with respective probabilities α and δ

$(m_1, m_2, \dots, m_N) \rightarrow (m_1, m_2, \dots, m_N \pm 1)$ with respective probabilities γ and β

and $(m_1, \dots, m_k, \dots, m_N) \rightarrow (m_1, \dots, m_k \pm 1, \dots, m_N)$ with respective probabilities p and q .

Thus,

$$(*) = \alpha \log \alpha + \delta \log \delta + \gamma \log \gamma + \beta \log \beta + (N - 1)(p \log(p) + q \log(q)). \tag{A2}$$

2. If $m_1 \geq 1$ and $m_N \geq 1$ and let i be the number of m_i between 2 and $N - 1$ equal to 0. With the same argument as previously we have:

$$(*) = \alpha \log \alpha + \delta \log \delta + \gamma \log \gamma + \beta \log \beta + (N - 1 - i)(p \log(p) + q \log(q)). \tag{A3}$$

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3. If $m_1 = 0$ and $m_N \geq 1$ and let i the number of m_j between 2 and $N - 1$ equal to 0 we have:

$$(*) = \alpha \log \alpha + \delta \log \delta + \beta \log \beta + (N - 2 - i)p \log(p) + (N - 1 - i)q \log(q). \quad (\text{A4})$$

4. The same applies if $m_1 \geq 1$ and $m_N = 0$ and let i the number of m_j between 2 and $N - 1$ equal to 0 we have:

$$(*) = \alpha \log \alpha + \delta \log \delta + \gamma \log \gamma + (N - 1 - i)p \log(p) + (N - 2 - i)q \log(q). \quad (\text{A5})$$

5. Finally, if $m_1 = 0$ and $m_N = 0$ and let i the number of m_j between 2 and $N - 1$ equal to 0 we have:

$$(*) = \alpha \log \alpha + \delta \log \delta + (N - 2 - i)(p \log(p) + q \log(q)). \quad (\text{A6})$$

10 Using Eq. (3) we find that $P(m_k = 0) = 1 - z_k$ and $\sum_{i=1}^{+\infty} P(m_k = i) = z_k$.
We thus obtain than h_{KS} writes:

$$\begin{aligned} h_{\text{KS}} = & -(\alpha \log \alpha + \delta \log \delta + \gamma \log \gamma + \beta \log \beta + (N - 1)(p \log(p) + q \log(q)) \\ & + (p \log(p) + q \log(q)) \left(\sum_{r=0}^N r \sum_{i_1 \dots i_N} \prod_{i=i_1, \dots, i_r} (1 - z_i) \prod_{i \neq i_1, \dots, i_r} z_i \right) \\ & + (\gamma \log(\gamma) + p \log(p)) z_N (1 - z_1) \left(\sum_{i_2 \dots i_{N-1}} \prod_{i=i_2, \dots, i_r} (1 - z_i) \prod_{i \neq i_2, \dots, i_r} z_i \right) \\ & + (\beta \log(\beta) + q \log(q)) z_1 (1 - z_N) \left(\sum_{i_2 \dots i_{N-1}} \prod_{i=i_2, \dots, i_r} (1 - z_i) \prod_{i \neq i_2, \dots, i_r} z_i \right) \\ & + (\beta \log(\beta) + \gamma \log \gamma + p \log p + q \log q) \left(\sum_{i_2 \dots i_{N-1}} \prod_{i=i_2, \dots, i_r} (1 - z_i) \prod_{i \neq i_2, \dots, i_r} z_i \right). \end{aligned} \quad (\text{A7})$$

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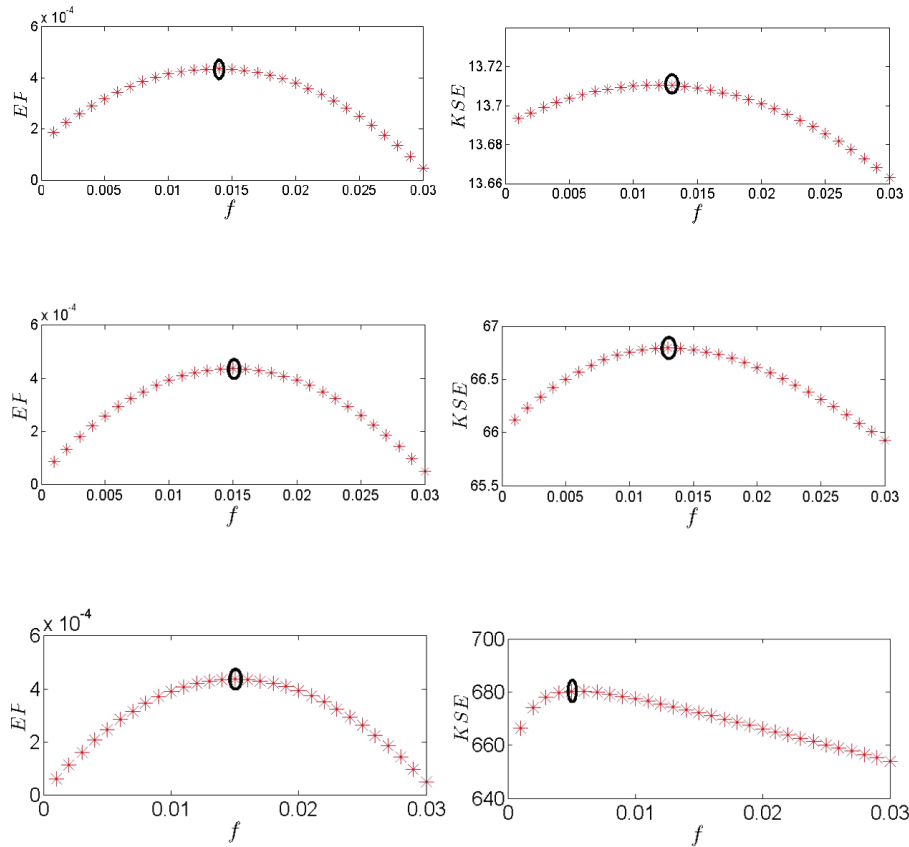


Figure 1. Entropy Production calculate using Eq. (10) (left panels) and KS Entropy calculate using Eqs. (6) and (14) (right panels) function of f for $s = 0.13$ and respectively $N = 20$ $N = 100$ and $N = 1000$.

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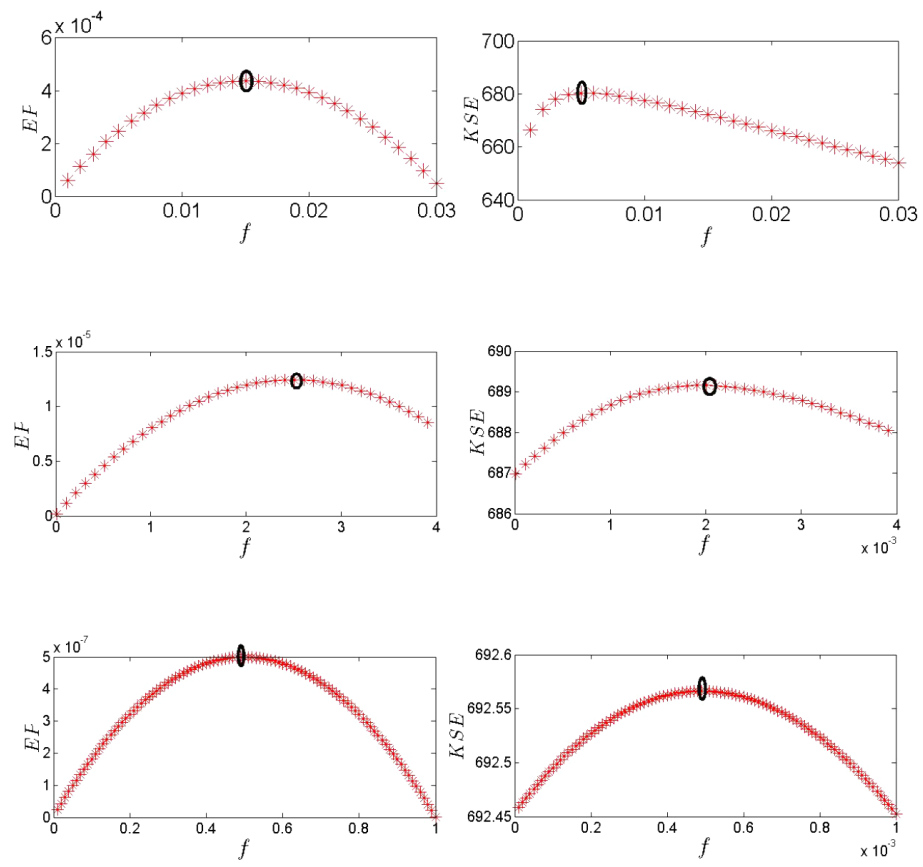


Figure 2. Entropy Production (left panels) and KS Entropy (right panels) function of f for $N = 1000$ and respectively $s = 0.13$; $s = 0.2$; $s = 0.04$.

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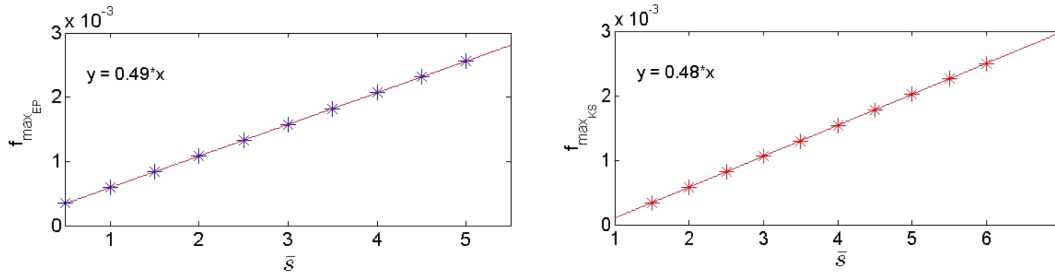


Figure 3. $f_{\max_{EP}}$ (left panel) and $f_{\max_{KS}}$ (right panel) function of \bar{s} for $\alpha = 0.5$ and $N = 100$. We remark that $f_{\max_{KS}}$ and $f_{\max_{EP}}$ have both a linear behaviour with slope respectively 0.48 and 0.49 which is really close to $\frac{1}{4\alpha} = 0.5$.

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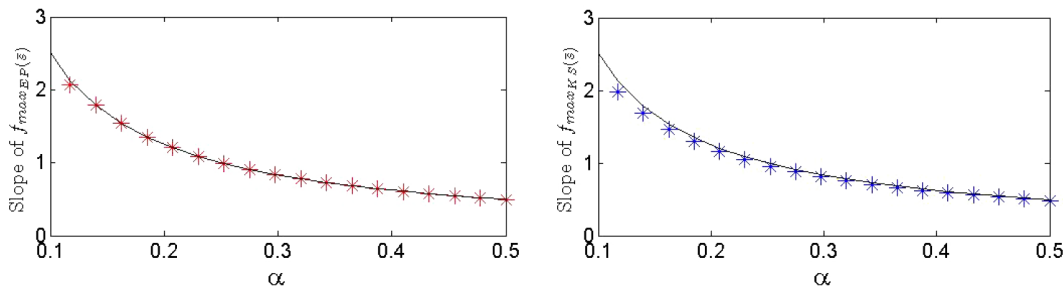


Figure 4. We plot the slope of $f_{\max_{KS}}(\bar{s})$ (left panel) and $f_{\max_{EP}}(\bar{s})$ (right panel) function of α and in black the curve $f(\bar{s}) = \frac{1}{4\alpha}\bar{s}$. We remark that the approximation $f_{\max_{KS}}(\bar{s}) \approx f_{\max_{EP}}(\bar{s}) \approx \frac{1}{4\alpha}\bar{s}$ is good.

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