

# Record dynamics in spin glasses, superconductors and biological evolution.

Henrik Jeldtoft Jensen  
Institute of Mathematical Sciences  
and  
Department of Mathematics

Collaborators:

Paolo Sibani, Paul Anderson, Luis P Oliveira and  
Mario Nicodemi

**Imperial College  
London**



# The question:

Is **intermittent**, logarithmically slow, dynamics, driven by **record** events, typical of complex systems?

## List of content:

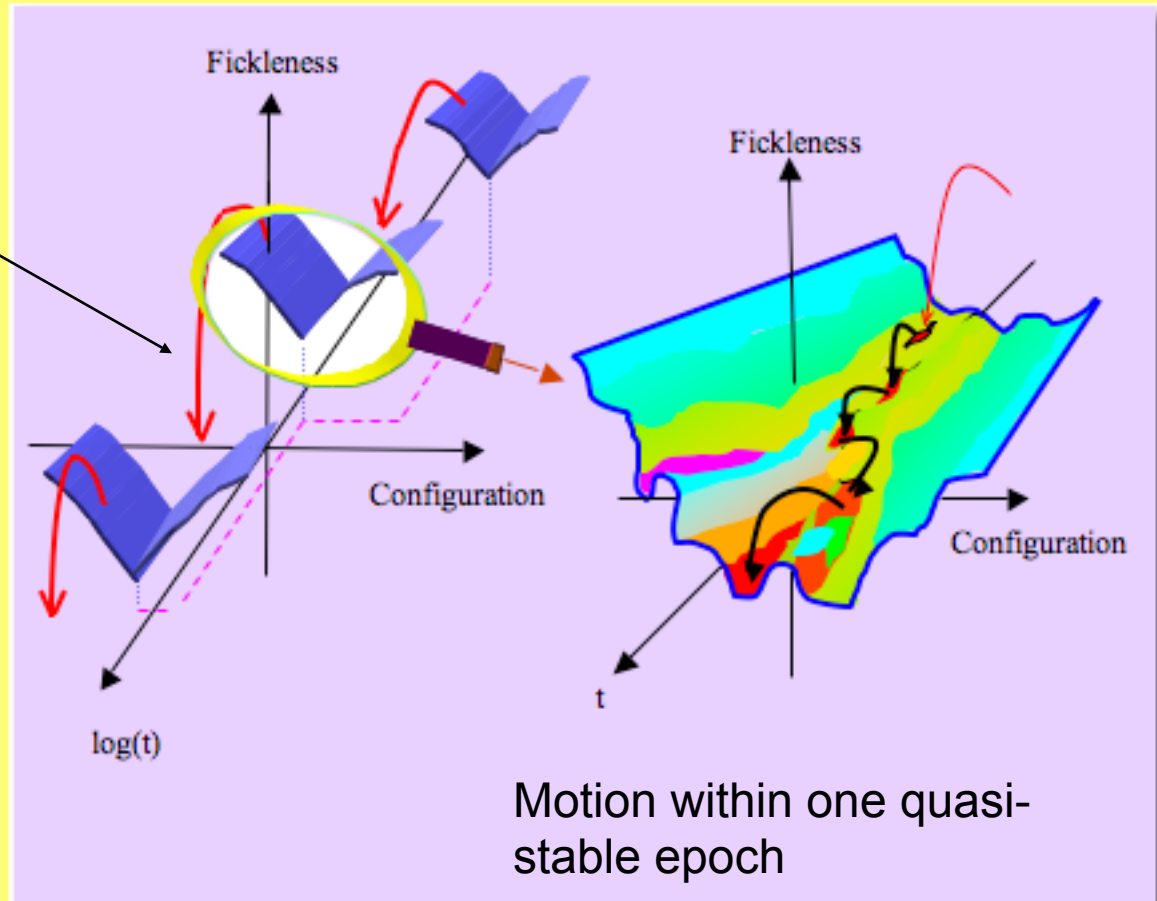
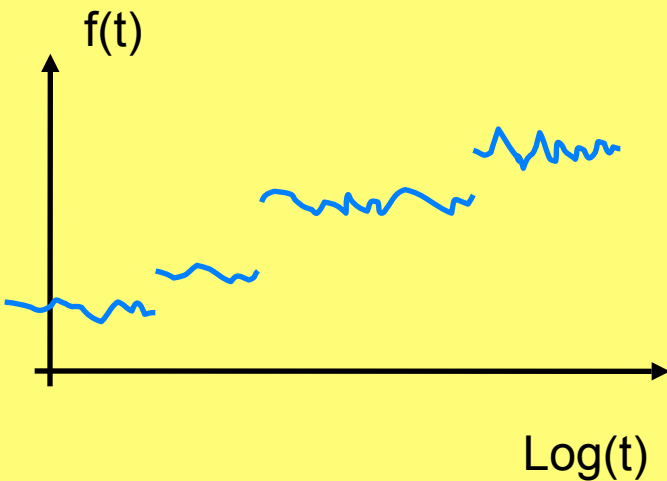
- Dynamics of complex systems
- Three models
  - $\mathbb{Z}^d$  definition and dynamics
- Manifestation of record dynamics
- Consequences
- Conclusion/summary

# Complex dynamics:

Intermittent, non-stationary

Jumping through collective adaptation space: quake driven

Transitions



The models:

**Tangled Nature Model** of co-evolving biological species

**Restricted Occupancy Model** of vortex dynamics in type II superconductors.

**Edward-Anderson Spin Glass** nearest neighbour Gaussian couplings

# The relaxation

## Tangled Nature model

collective adaptation: configurations increasingly coupled together.

## ROM model

magnetic pressure

## Spin Glass

thermal quench

First Model:

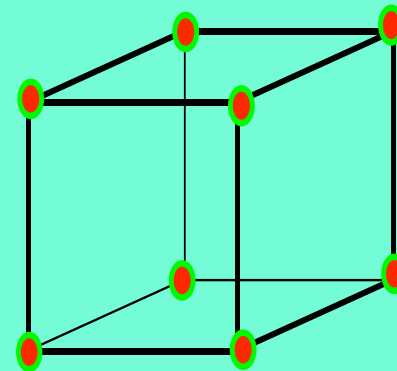
Tangled Nature

# Tangled Nature model of evolution

## Definition:

\* Individuals  $\mathbf{S}^\alpha = (S_1^\alpha, S_2^\alpha, \dots, S_L^\alpha)$ , where  $S_i^\alpha = \pm 1$

and  $\alpha = 1, 2, \dots, N(t)$



$L = 3$

\* Dynamics – a time step:

☹ Annihilation:

Choose indiv. at random, remove with probability

$$p_{kill} = \text{const}$$



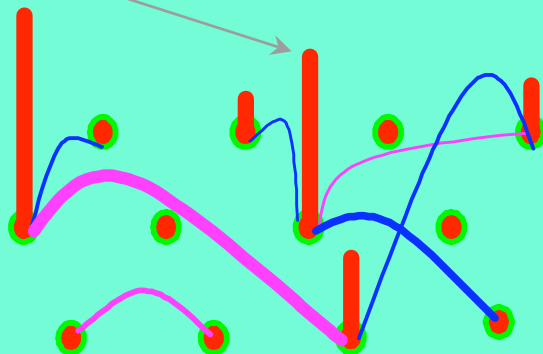


# Reproduction:

- ▶ Choose indiv. at random
- ▶ Determine

$$H(\mathbf{S}^\alpha, t) = \frac{1}{cN(t)} \sum_{\mathbf{S}} J(\mathbf{S}^\alpha, \mathbf{S}) n(\mathbf{S}, t) - \mu N(t)$$

$n(\mathbf{S}, t) =$  occupancy at the location  $\mathbf{S}$

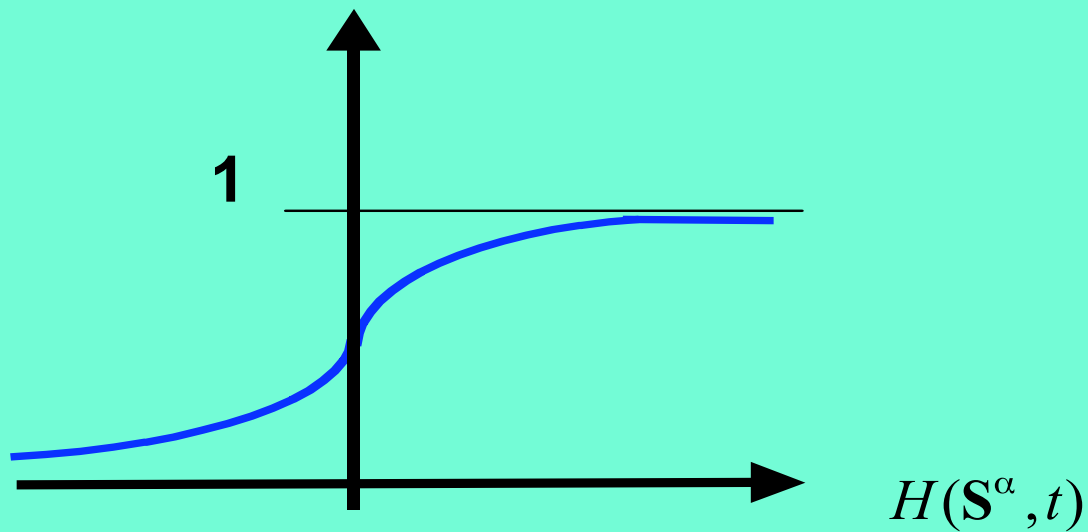


# The coupling matrix $J(S, S')$

- Either consider  $J(S, S')$  to be uncorrelated
- or to vary smoothly through type space.

from  $H(\mathbf{S}^\alpha, t)$  reproduction probability

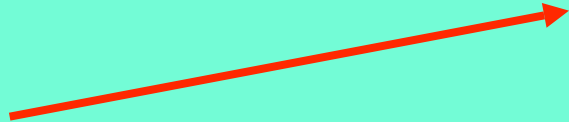
$$p_{off}(\mathbf{S}^\alpha, t) = \frac{\exp[H(\mathbf{S}^\alpha, t)]}{1 + \exp[H(\mathbf{S}^\alpha, t)]} \in [0, 1]$$



☺ Asexual reproduction:

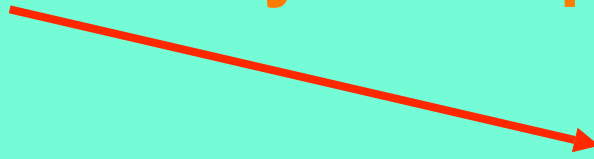
Replace

$S^\alpha$



$S_1^\alpha$

by two copies



$S_2^\alpha$

with probability

$$P_{off}(S^\alpha, t)$$

# Mutations

- ☺ Mutations occur with probability

$P_{mut}$ , i.e.

$$S_i^\gamma \mapsto -S_i^\gamma$$

# Phenomenology

- Long time dynamics
- The evolved networks



# Segregation in genotype space

Non Correlated

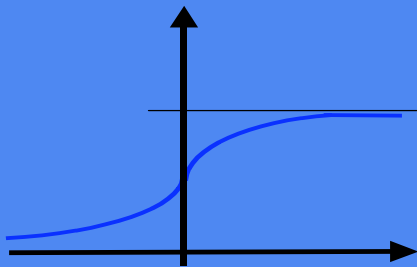
## Initiation

Only one genotype

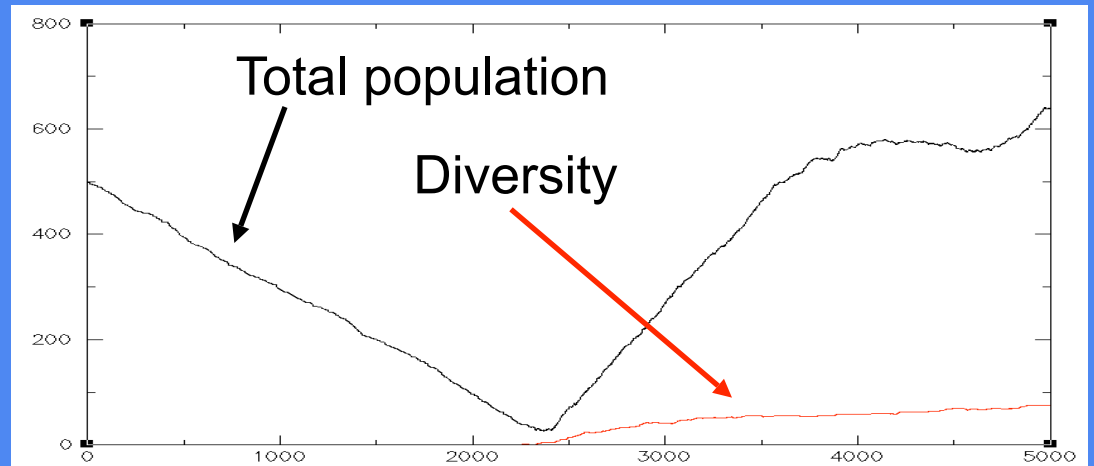
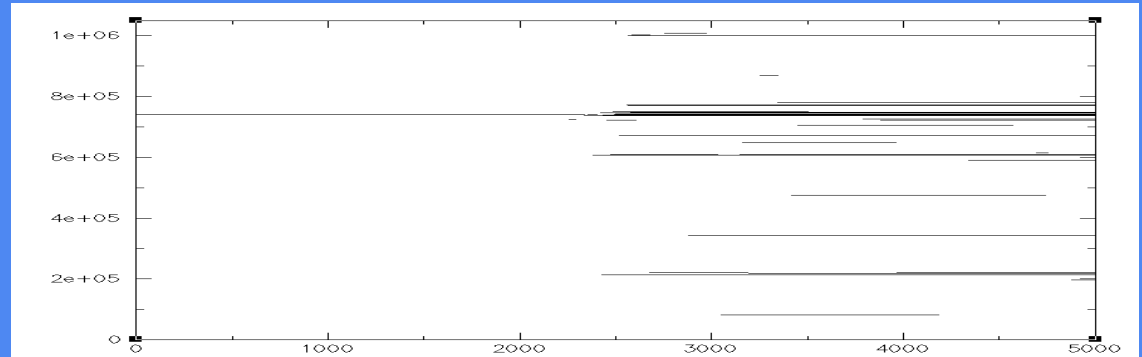
$J_n$  term = 0

$$H = \frac{k}{N(t)} \sum_s J_n - \mu N(t)$$

$N(t)$  adjusts

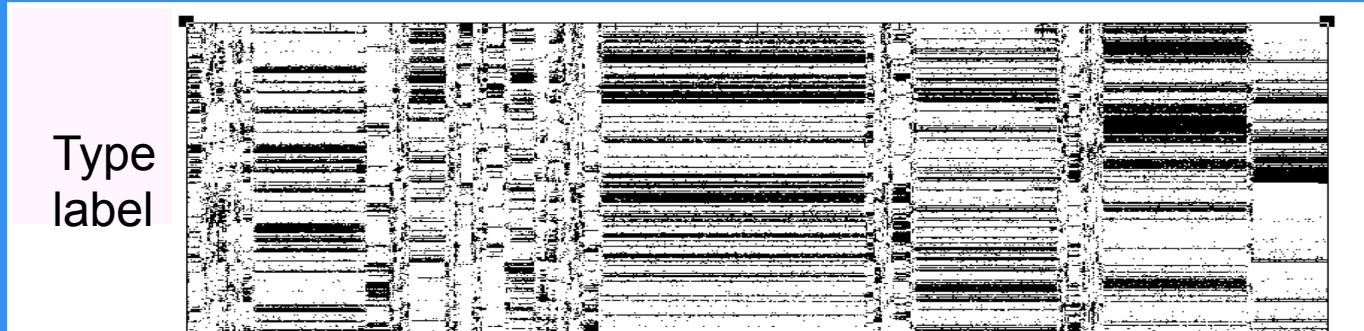


$$H = -\mu N(t)$$



# Intermittency at systems level:

Non Correlated



# generations

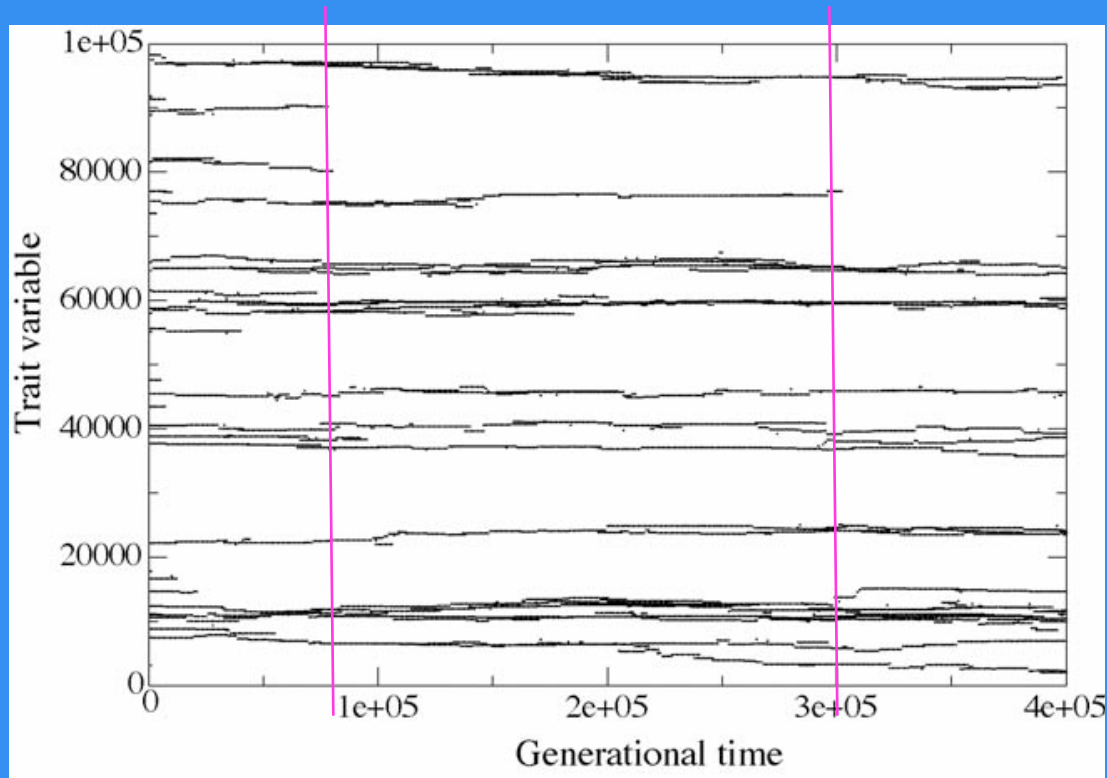
1 generation

$$= N(t) / p_{kill}$$



# Intermittency at systems level:

## Correlated



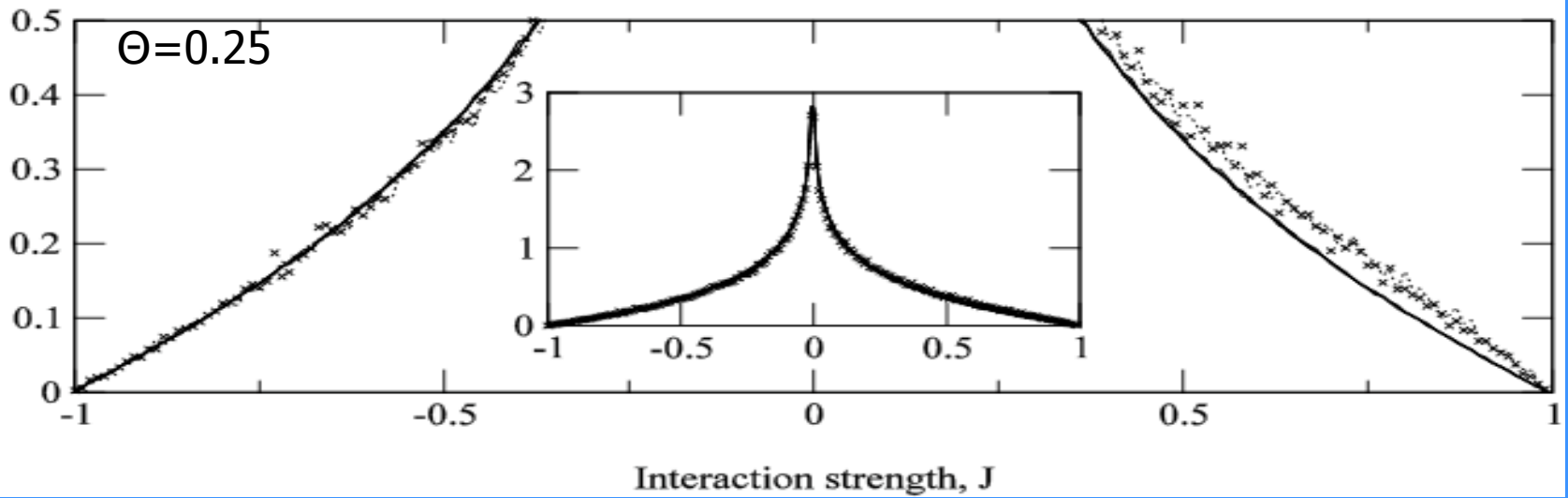
**Fig. 1 – An occupation plot of a single run for a system with  $R = 10,000$ . For each timeslice a point appears where a phenotype is in existence but as the full space is in 16 dimensions a projection onto a single trait is used.**

# Time evolution of

## Distribution of active coupling strengths

Non correlated

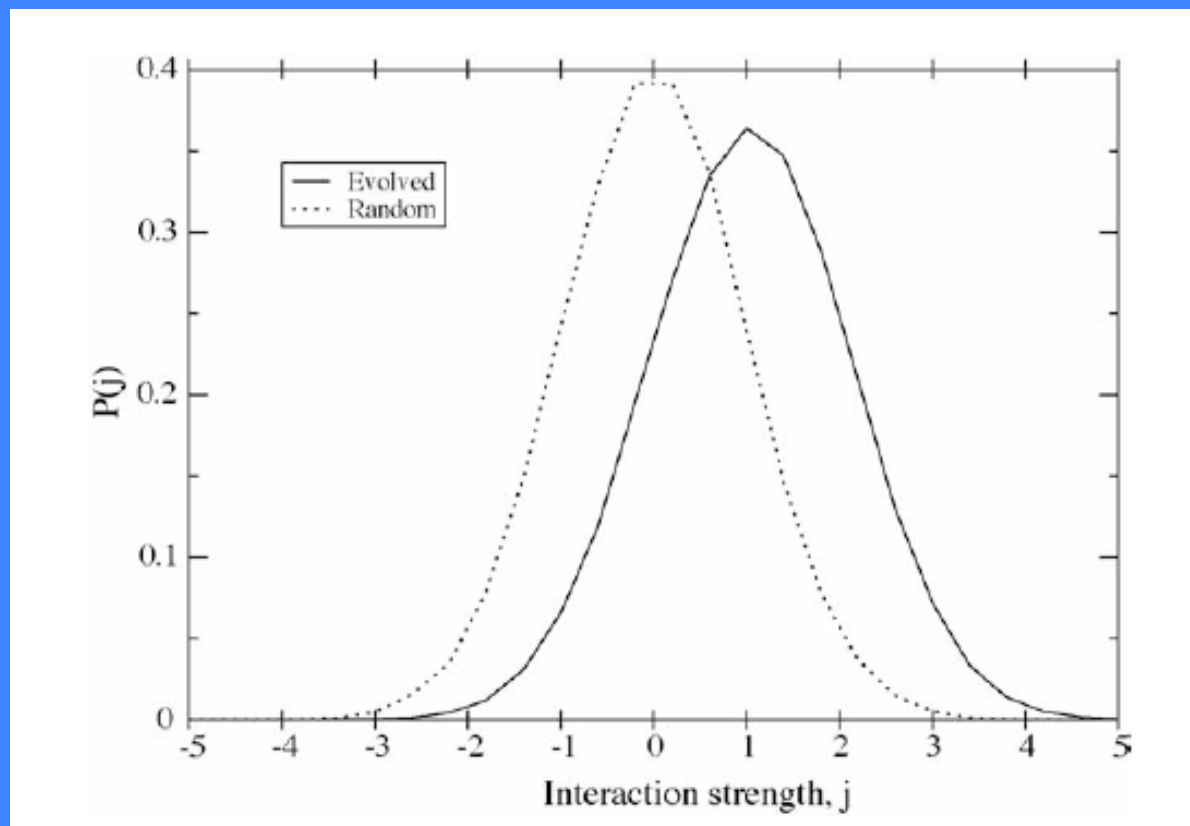
Normalised density of individuals with strength  $J$



# Time evolution of

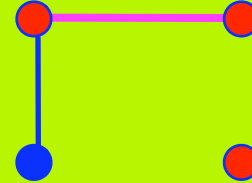
## Distribution of active coupling strengths

Correlated

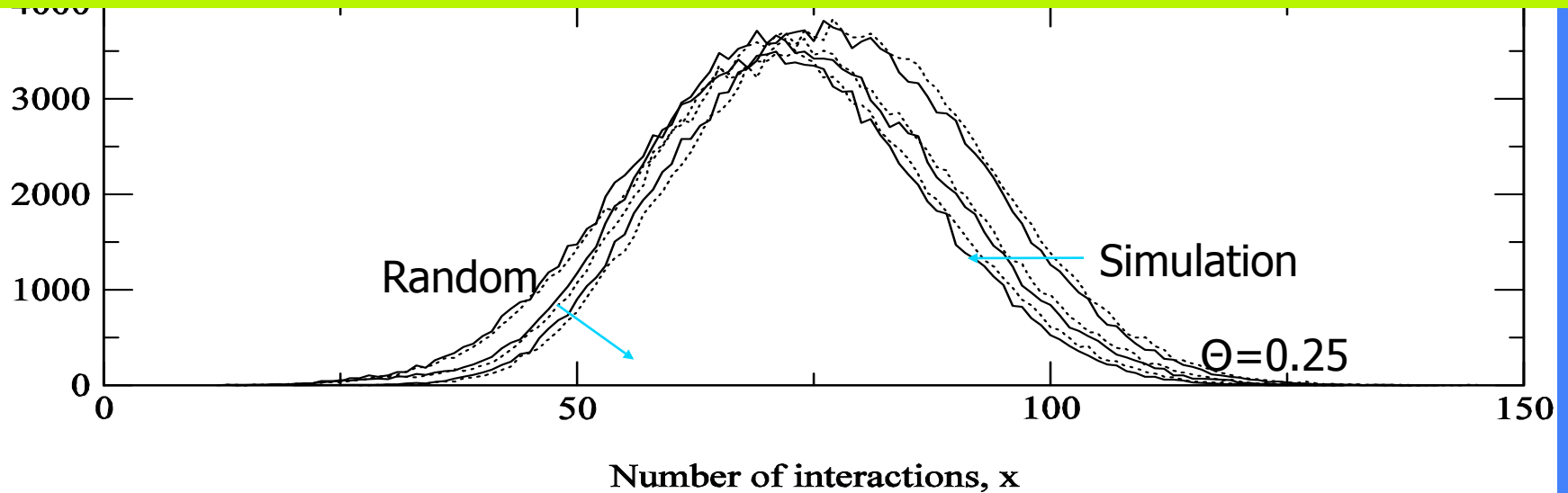


# Increasing complexity ?

$x=1$



Number of sites with  $x$  interactions

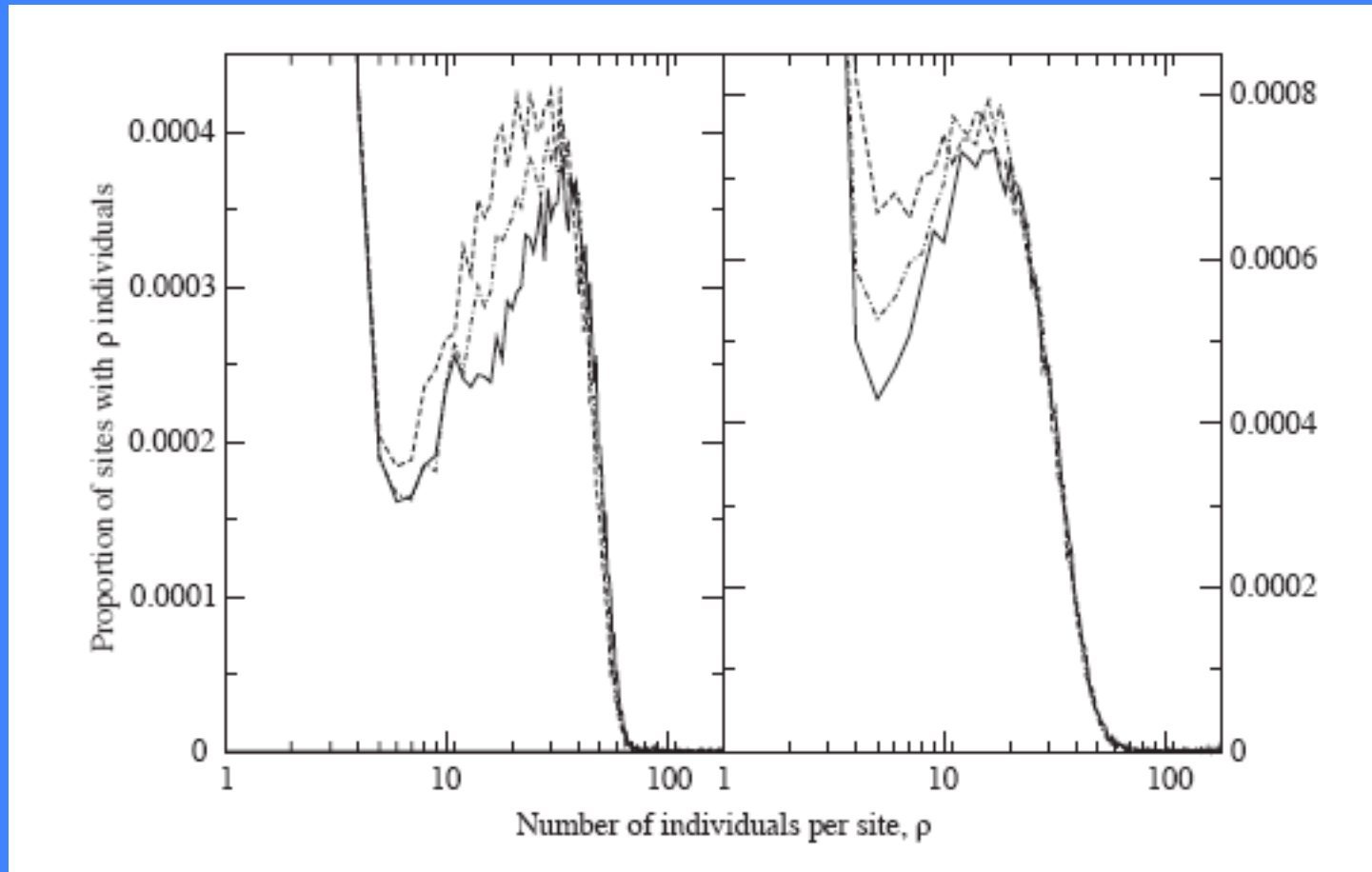


Note: Effect is significant for correlated type space

# Time evolution of

## Species abundance distribution

### Non Correlated



# The evolved degree distribution

## Correlated

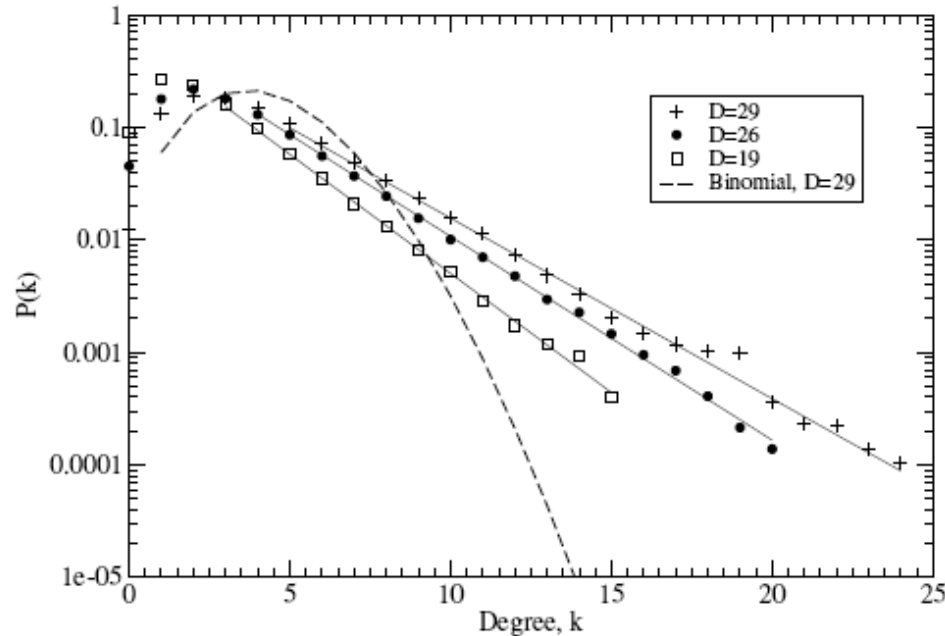


Figure 1: Degree distributions for the Tangled Nature model simulations. Shown are ensemble averaged data taken from all networks with diversity,  $D = \{19, 26, 29\}$  over 50 simulation runs of  $10^6$  generations each. The exponential forms are highlighted by comparison with a binomial distribution of  $D = 29$  and equivalent connectance,  $C \simeq 0.145$  to the simulation data of the same diversity.

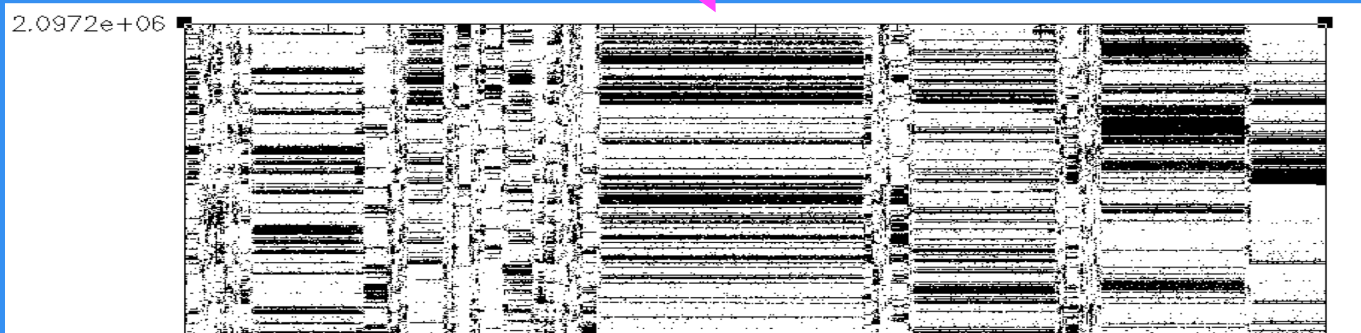
Simon Laird

Exponential becomes  $1/k$  in limit of vanishing mutation rate

# Intermittent dynamics

# Intermittency:

q-ESS = quasi-Evolutionary Stable Strategy

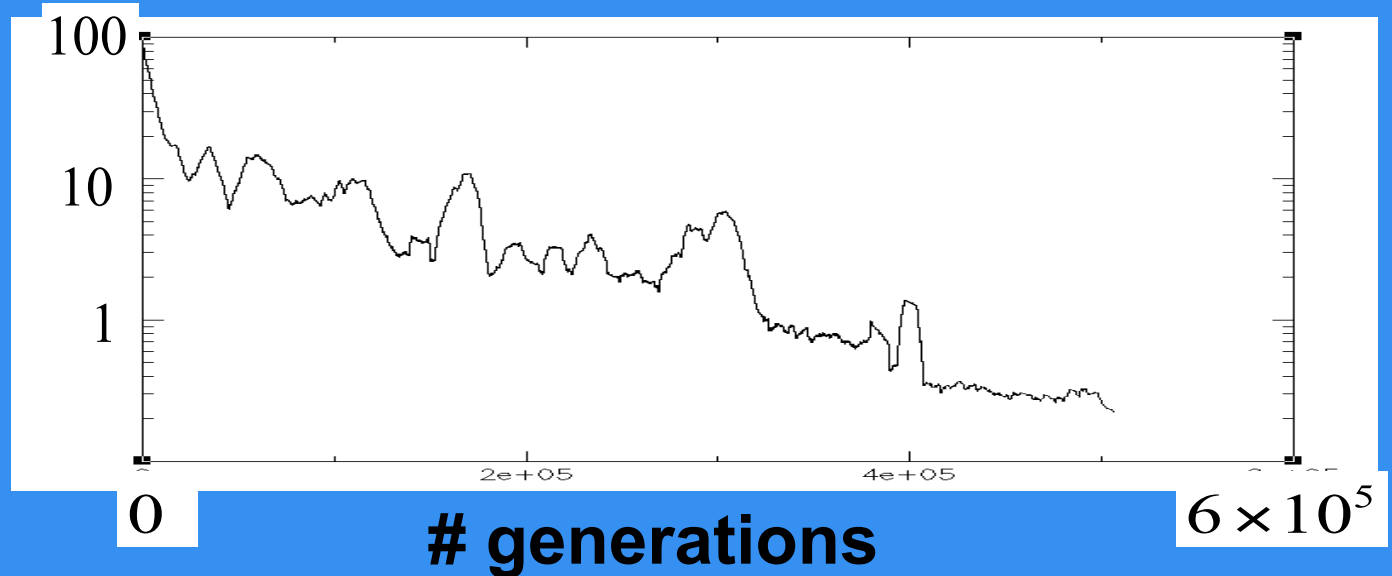


# of transitions in window

Matt Hall

1 generation

$$= N(t) / p_{kill}$$





# Stability of the q-ESS:

Consider simple adiabatic approximation.

Stability of genotype  $S$  assuming:  $n(S', t)$  independent of  $t$  for  $S' \neq S$

Consider 
$$\frac{\partial n(S, t)}{\partial t} = [p_{off}(n(S, t), t) - p_{kill} - p_{mut}] \frac{n(S, t)}{N(t)}$$

Stationary solution  $n_0(S)$  corresponds to  $p_{off}(n_0(S)) - p_{kill} - p_{mut} = 0$

Fluctuation  $\delta = n(S, t) - n_0(S)$

Fulfil 
$$\dot{\delta} = A \frac{n_0}{N_0} \delta$$

with 
$$A = -(1 - p_{mut})(p_{off})^2 e^{-H_0} \left( \frac{J}{N_0^2} + \mu \right) < 0$$

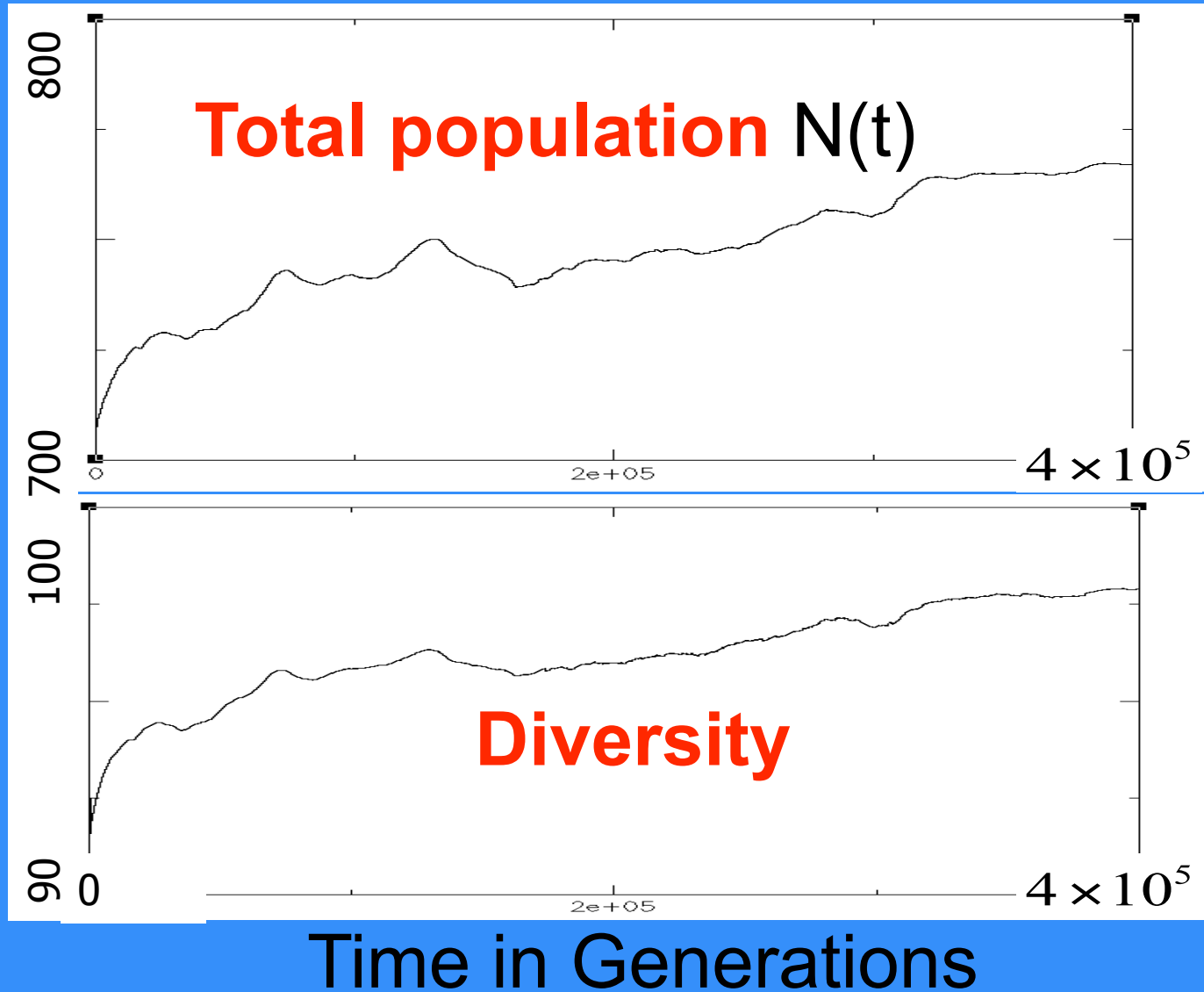
**i.e. stability**

Transitions between q-ESS caused by co-evolutionary collective fluctuations

$n(S', t)$  needs to be considered

dependent of  $t$  for  $S' \neq S$

😊 Time dependence (averaged)





# Origin of drift?

Effect of mutation

Let

$H = \tilde{J} - \mu N$ , then the effect of a mutation is

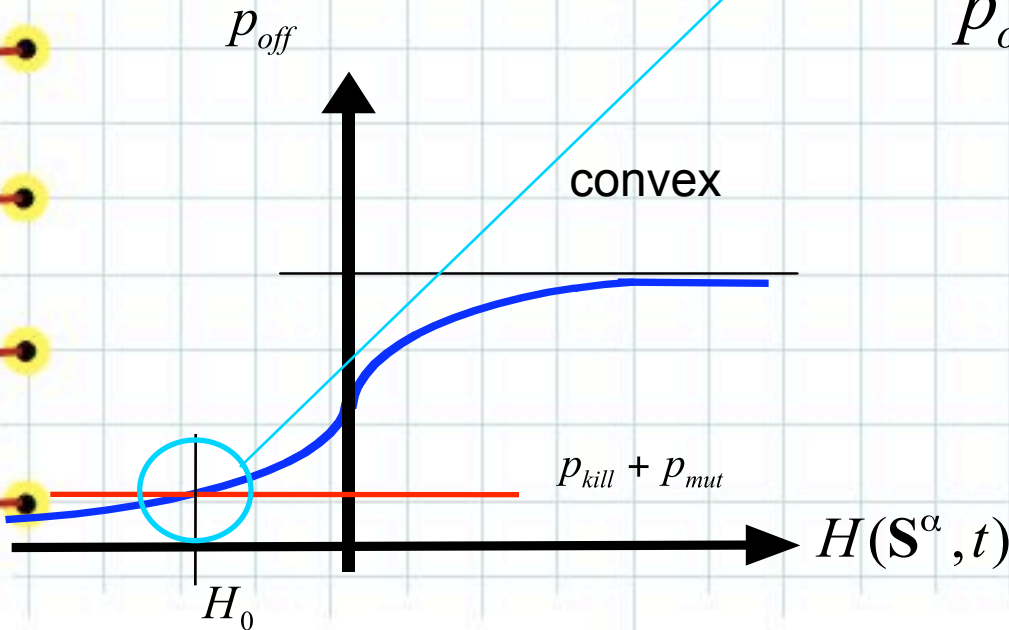
$$H \mapsto H + \delta \tilde{J}.$$

→ Symmetric fluctuations  $prob(\delta \tilde{J}) = prob(-\delta \tilde{J})$

leads to asymmetri

$$p_{off}(H_0 + \delta \tilde{J}) - p_{kill} >$$

$$p_{kill} - p_{off}(H_0 - \delta \tilde{J})$$





$$p_{off}(H_0 + \delta \tilde{J}) - p_{kill} > p_{kill} - p_{off}(H_0 - \delta \tilde{J})$$



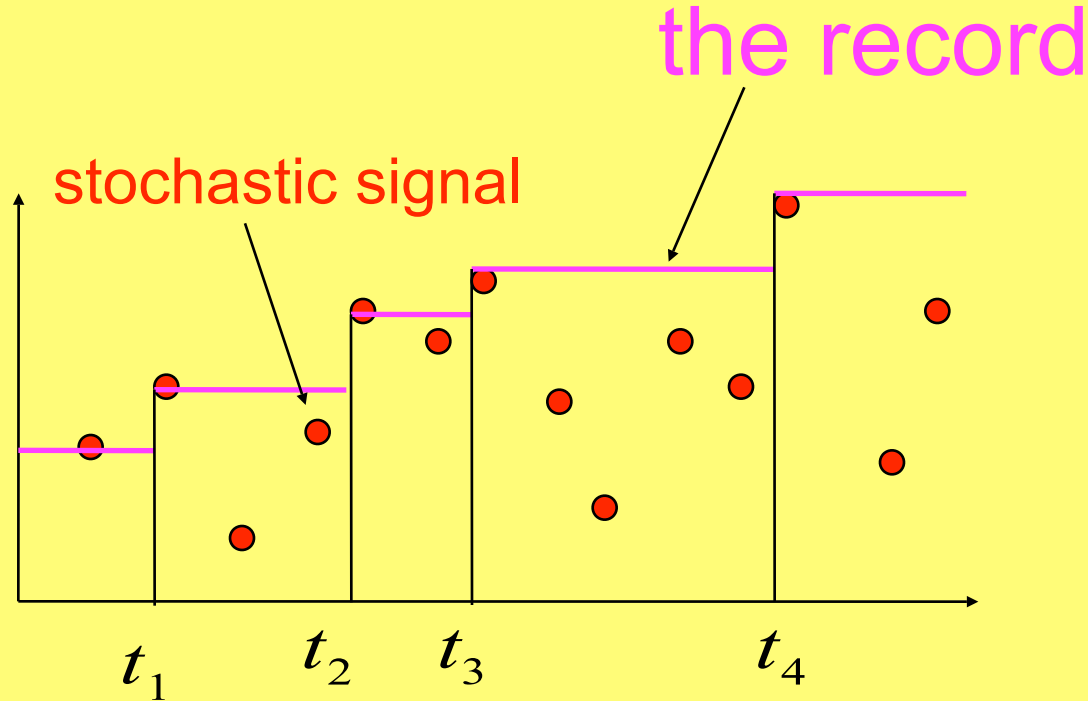
$$\delta N_+ > \delta N_-$$



Not the whole explanation: evolution not smooth.

# Record dynamics

## Record dynamics:



**Paolo Sibani and Peter Littlewood (1992):**

$$\tau = \ln(t_k) - \ln(t_{k-1}) = \ln\left(\frac{t_k}{t_{k-1}}\right) \text{ exponentially distributed}$$

## Record dynamics:

$$\tau = \ln(t_k) - \ln(t_{k-1}) = \ln\left(\frac{t_k}{t_{k-1}}\right) \quad \text{exponentially distributed}$$



- ▶ Poisson process in logarithmic time
- ▶ Mean and variance

$$\langle Q \rangle \propto \ln(t) \quad \text{and} \quad \langle (Q - \langle Q \rangle)^2 \rangle \propto \ln(t)$$

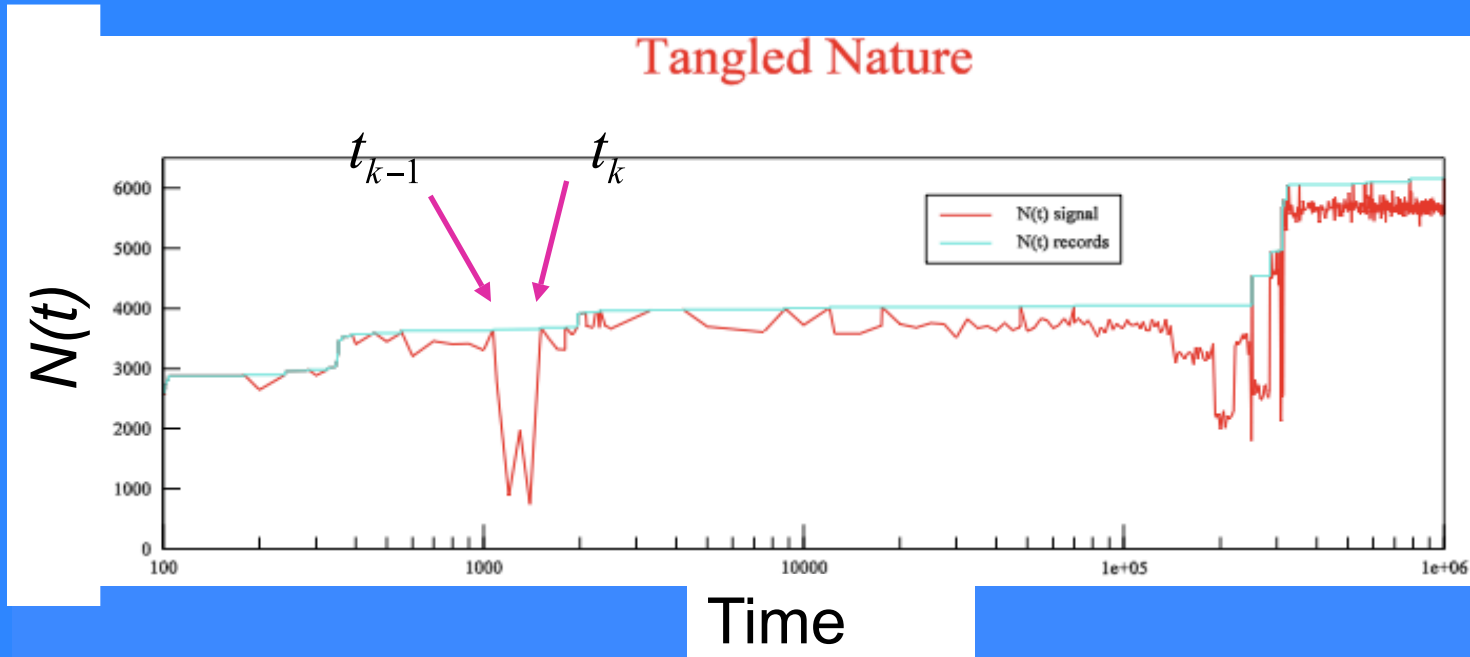
- ▶ Rate of records constant as function of  $\ln(t)$
- ▶ Rate decreases  $\propto \frac{1}{t}$



# Tangled Nature model:

## Single realisation and record dynamics:

Extracting records from the population size

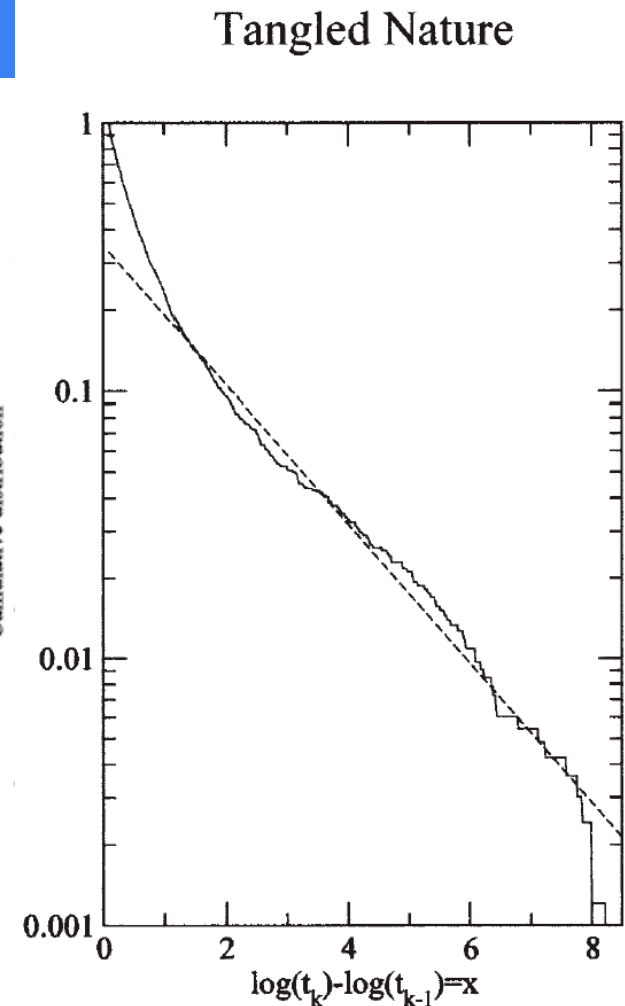
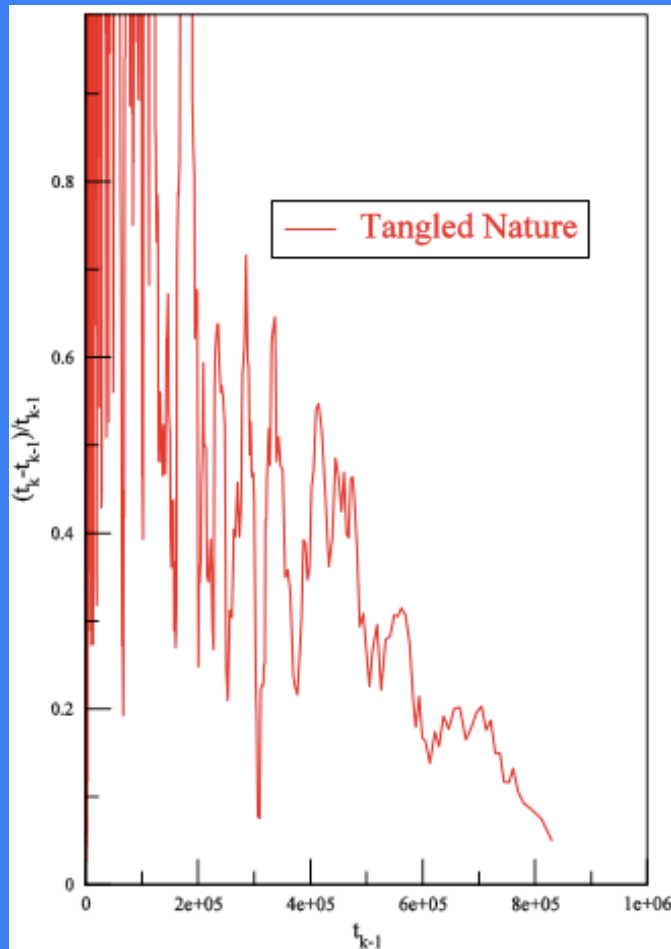


# Record dynamics:

Ratio  $r$  remains  
non-zero

$$r = (t_k - t_{k-1}) / t_{k-1}$$

Cumulative Distribution

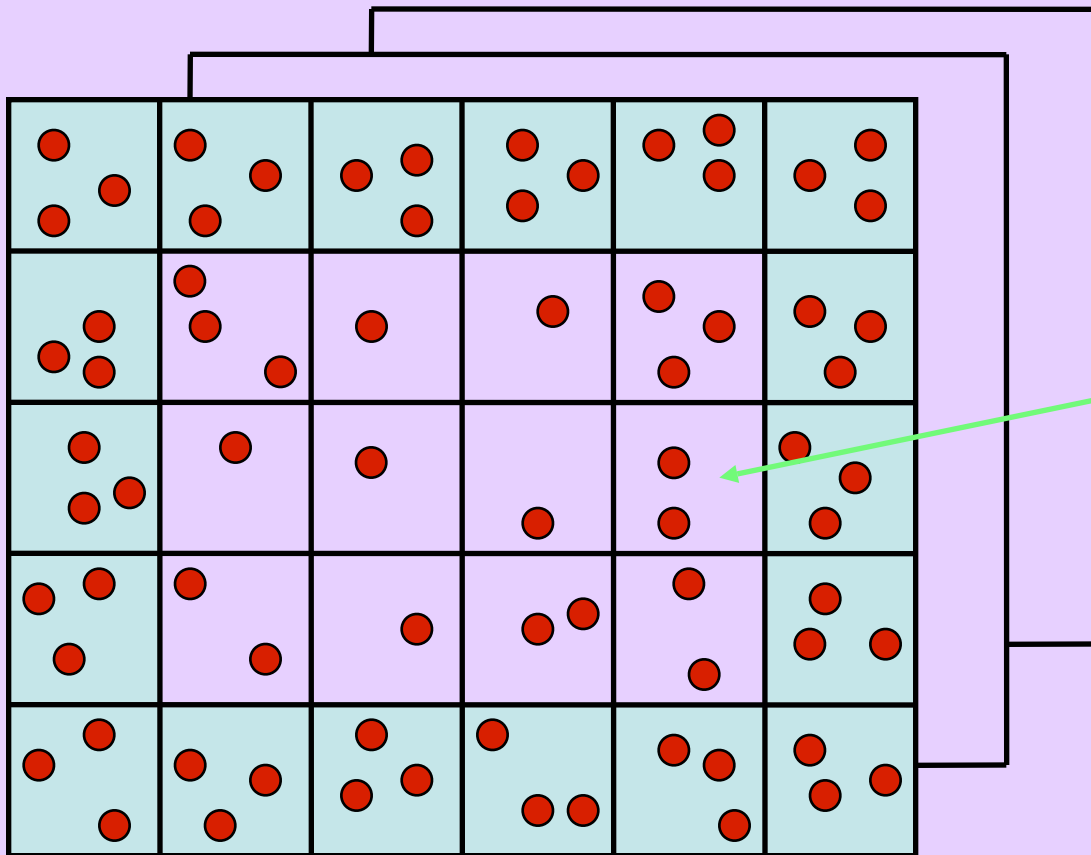


# Second Model:

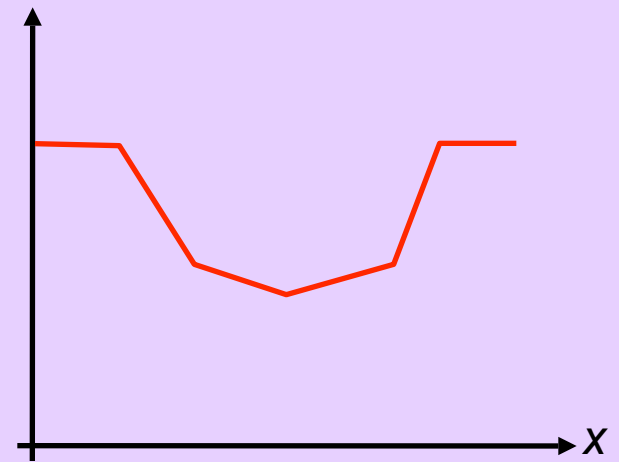
## ROM

# ROM

Monte Carlo Kawasaki dynamics on stack of coarse grained superconducting planes



$$n(x, y, z, t) = n_i$$



ROM

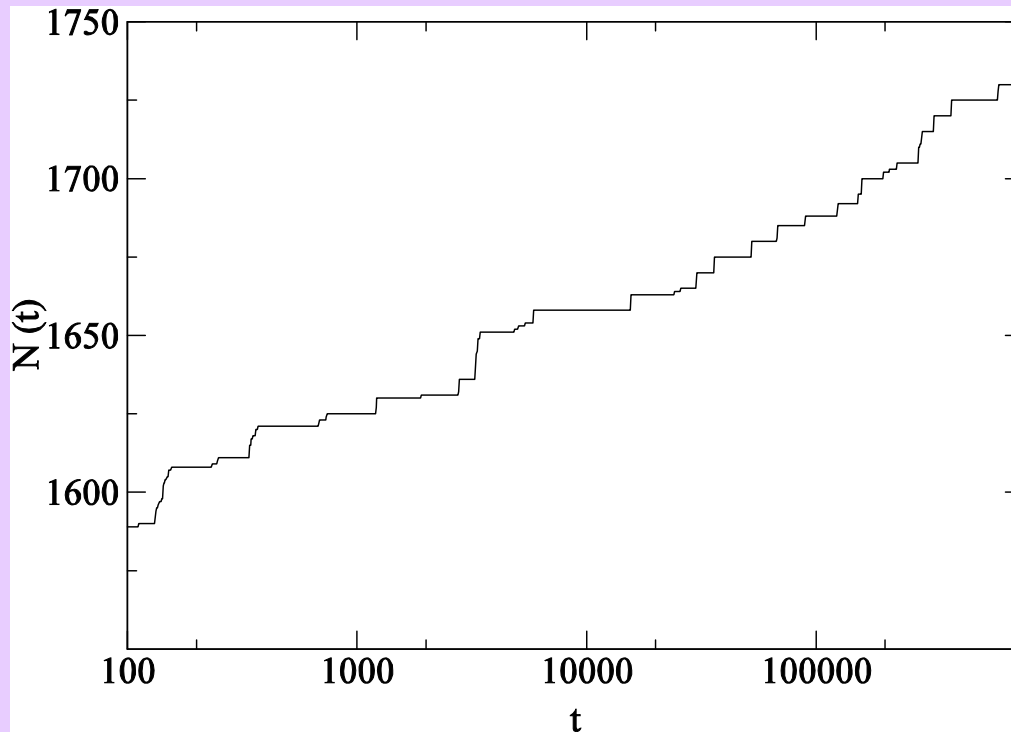
Hamiltonian

$$H = \sum_i n_i A_{ij} n_j - \sum_i A_{ii} n_i - \sum_i A_i^p n_i + \sum_{\langle ij \rangle_z} A_2 (n_i - n_j)^2$$

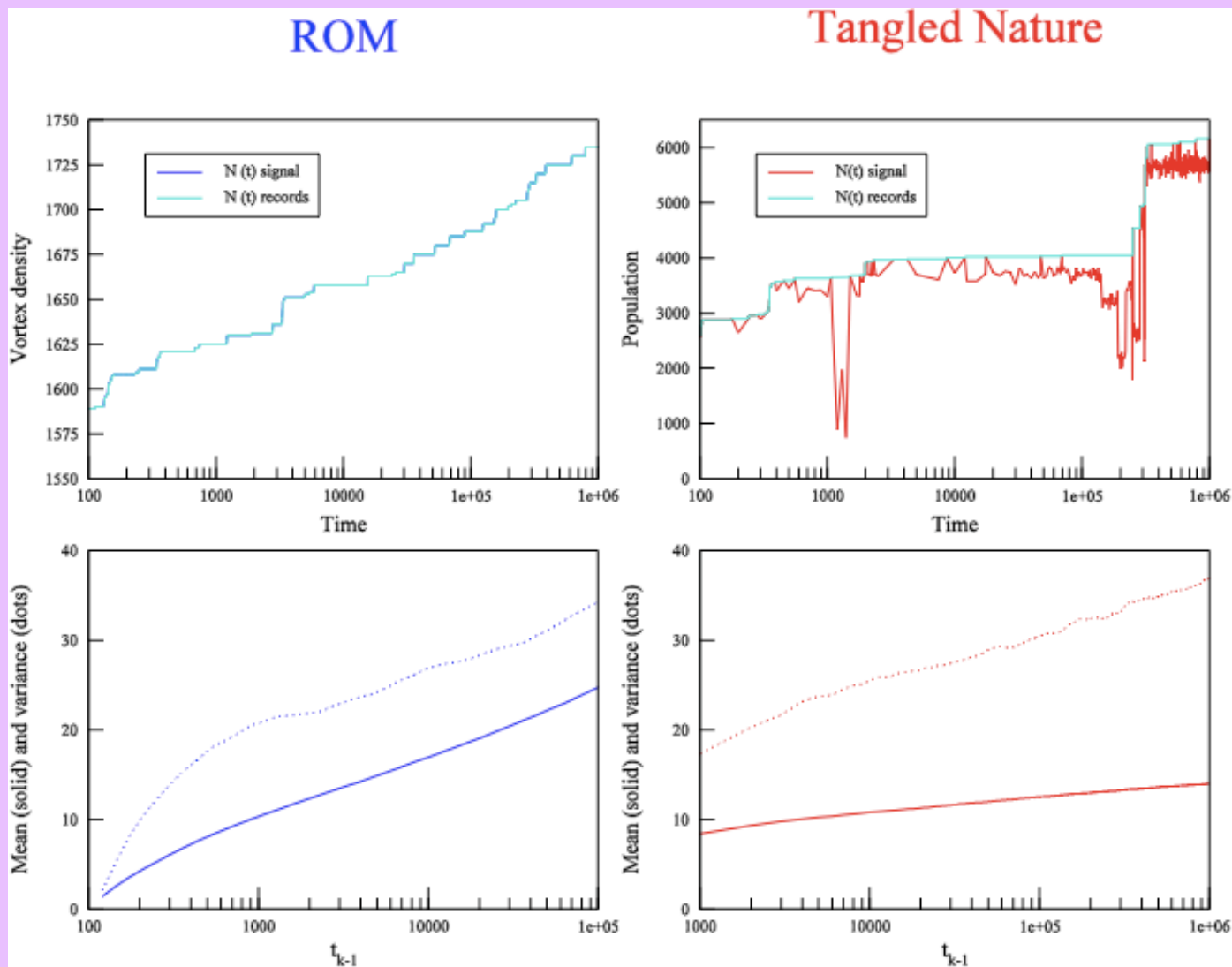
Here

$$0 \leq n_i < N_{c2} = \frac{B_{c2} l_0^2}{\varphi_0}$$

# ROM: Temperature independent creep



# Realisations of record dynamics

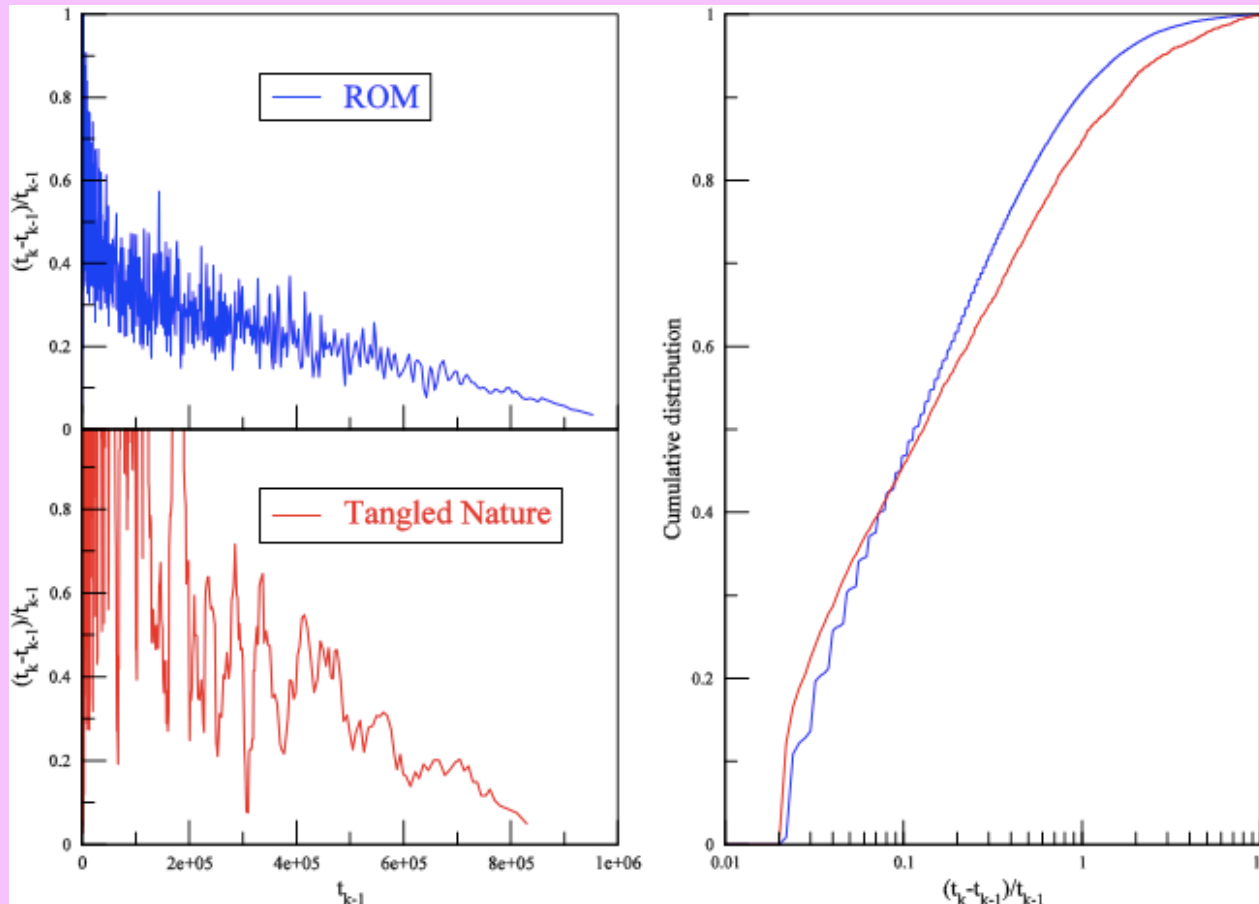


# Manifestation of the decelerating activity.

For stationary process

$$\frac{t_k - t_{k-1}}{t_{k-1}} \approx \text{const}$$

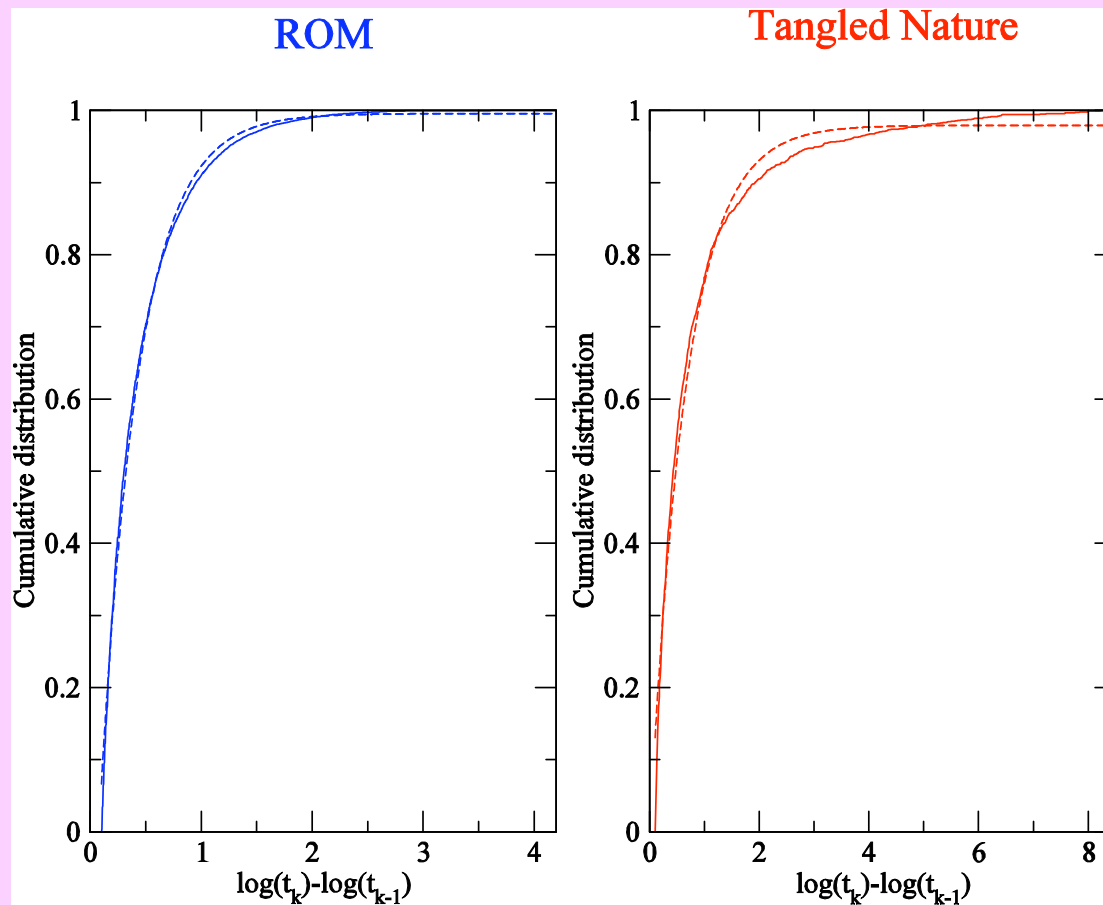
$$P\left(\frac{t_k - t_{k-1}}{t_{k-1}} = x\right) = \delta(x)$$



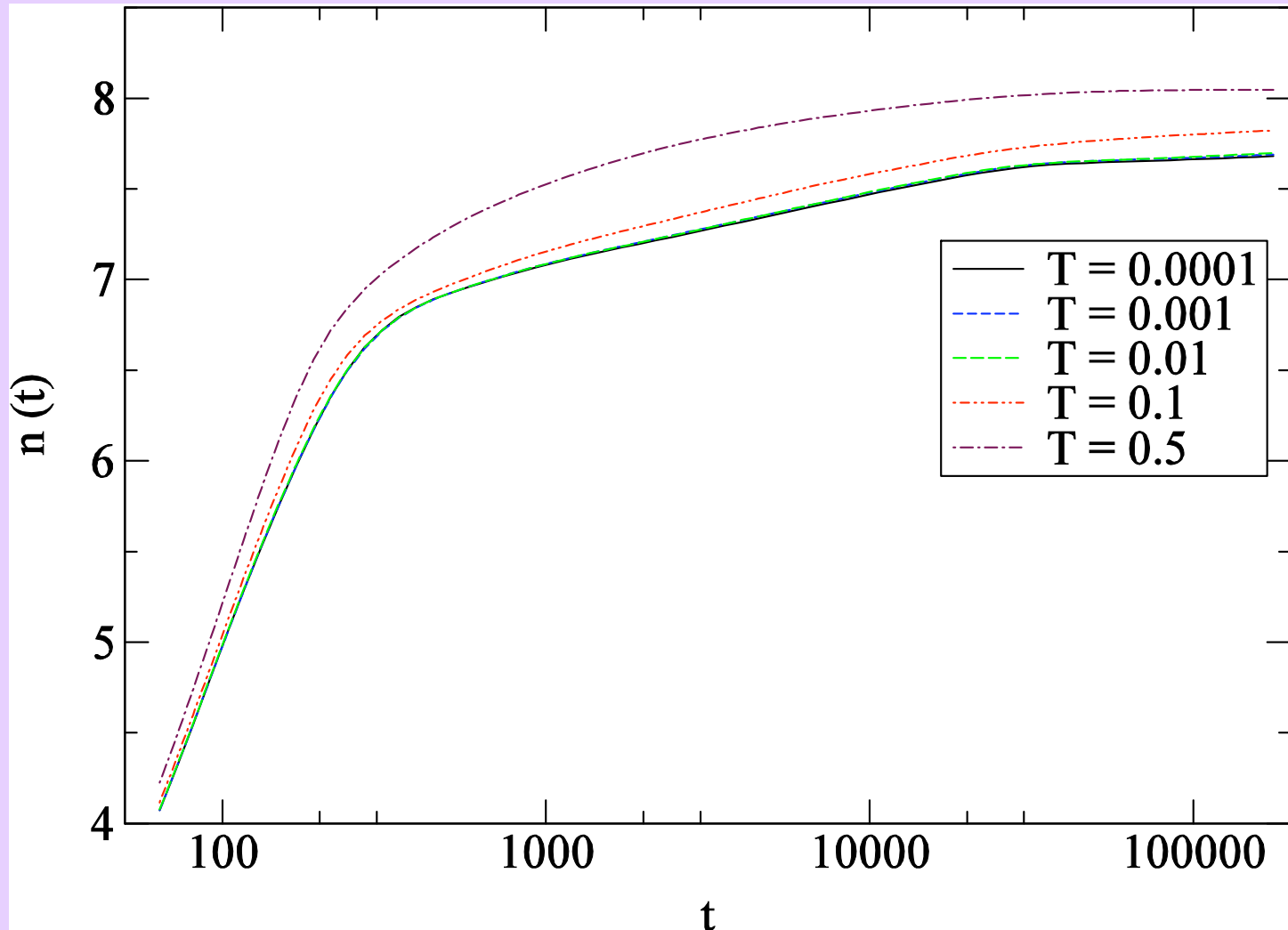


## Further evidence

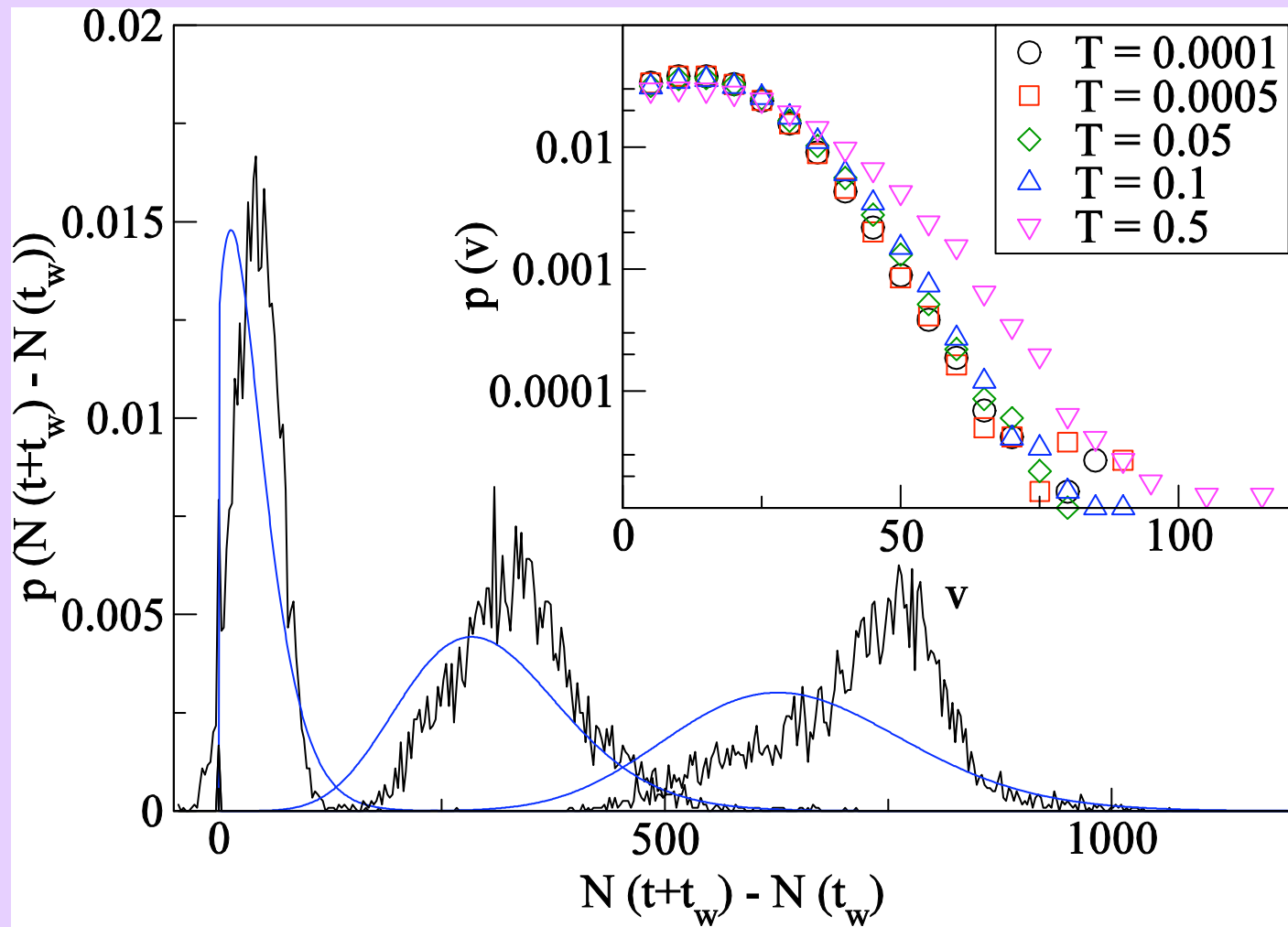
The cumulative distribution of the log waiting times.  
Comparison with exponential distribution.



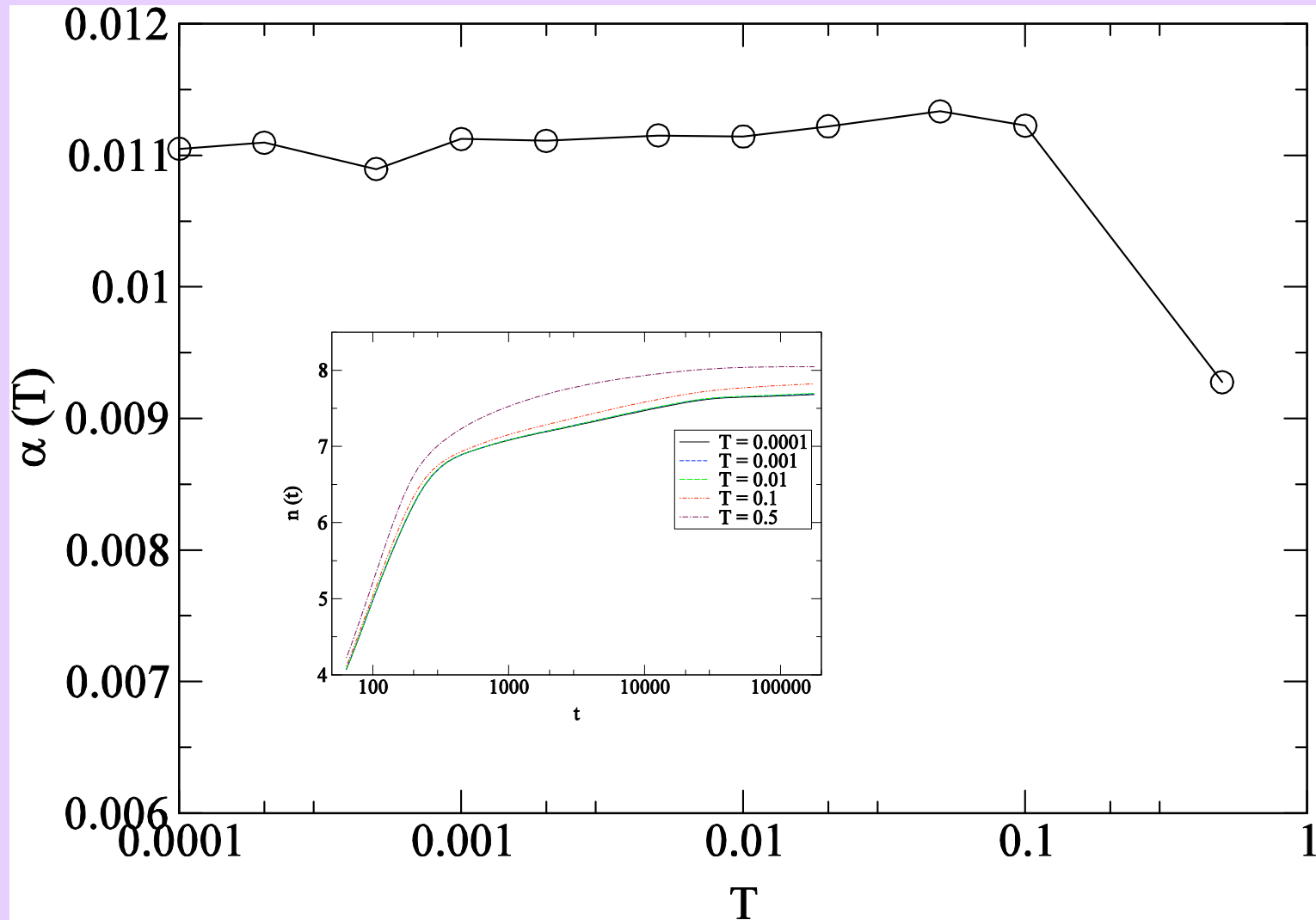
# Number of vortices in the bulk as function of time



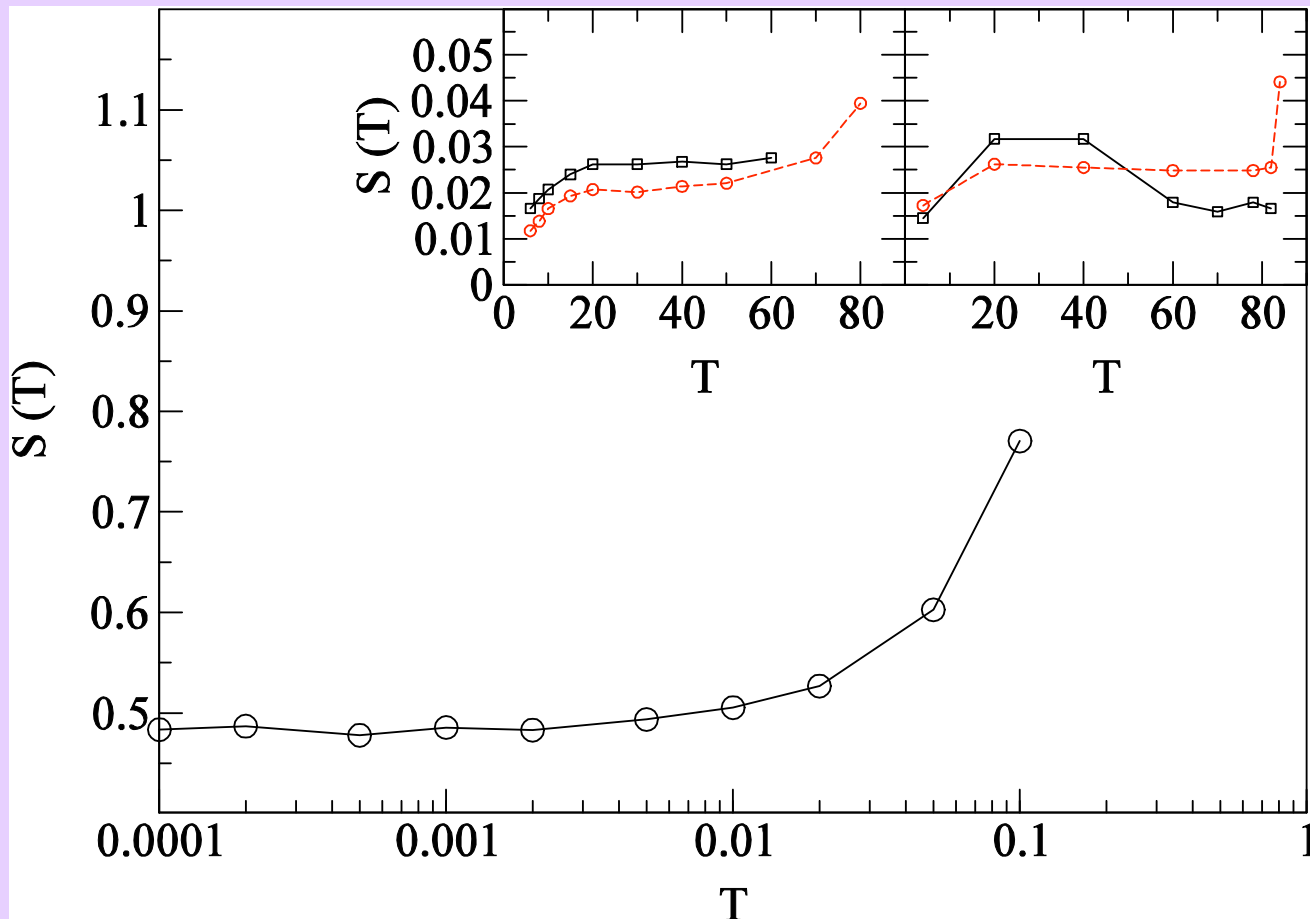
# Quake statistics and the total number vortices entering.



# The temperature in-dependence of the quake rate.



The magnetic creep rate:  $S = \frac{d \ln(M)}{d \ln(t)}$  where  $M(t) = |N(t) - N_{ext}|$   
 comparison with experiment



# Third Model:

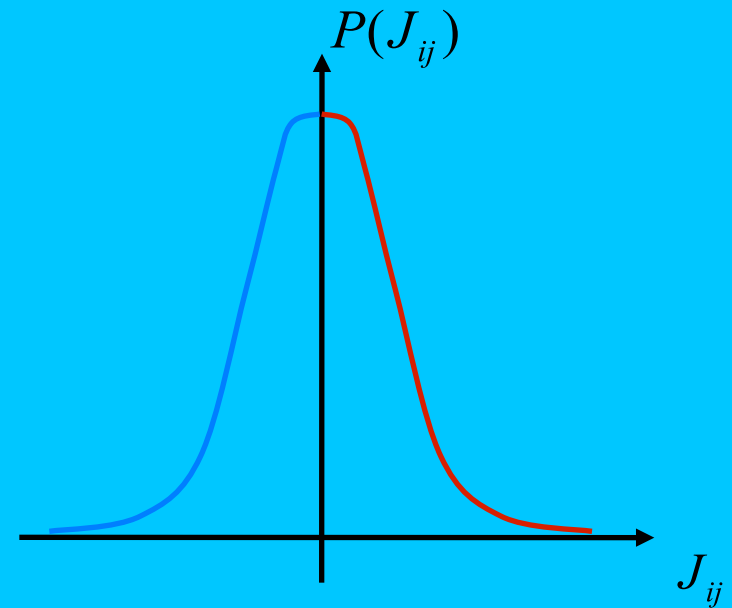
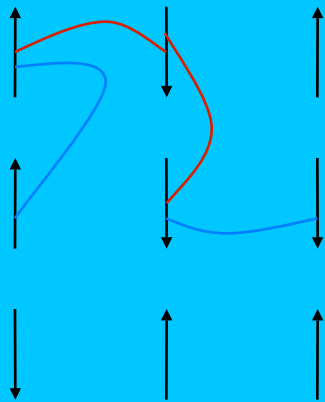
## Spin Glass

# Spin glass

Microscopic magnetic moments – or spins – coupled together with random coupling constants.

The Hamiltonian:

$$H = -\frac{1}{2} \sum_{ij} J_{ij} \mathbf{S}_i \cdot \mathbf{S}_j \text{ where } \mathbf{S}_i, \mathbf{S}_j = \pm 1$$

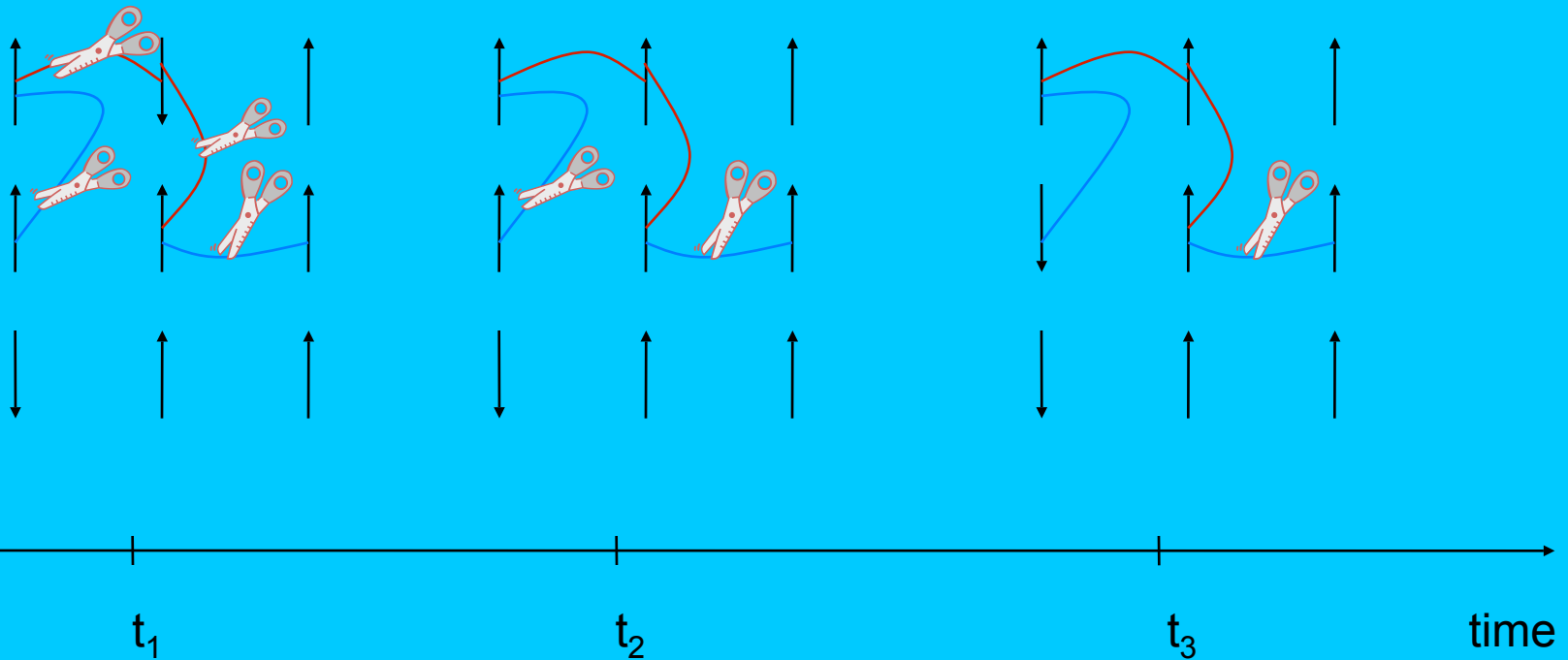


# Spin glass

Quench from high temperature:

time  $< 0$ :  $T = \text{high}$

time  $> 0$ :  $T = \text{very low}$





## Spin glass: heat transfer

Protocol: Quench from high temp. at time  $t=0$ .

Measure heat transfer,  $H$ , between spin glass and reservoir during time interval

$$[t_w, t_w + \delta t]$$

- If  $\delta t \ll t_w$  Gaussian  $p(H)$
- If  $\delta t \approx t_w$  exponential tail

# Spin glass: heat transfer

$$\delta t \ll t_w$$

$$\delta t \approx t_w$$

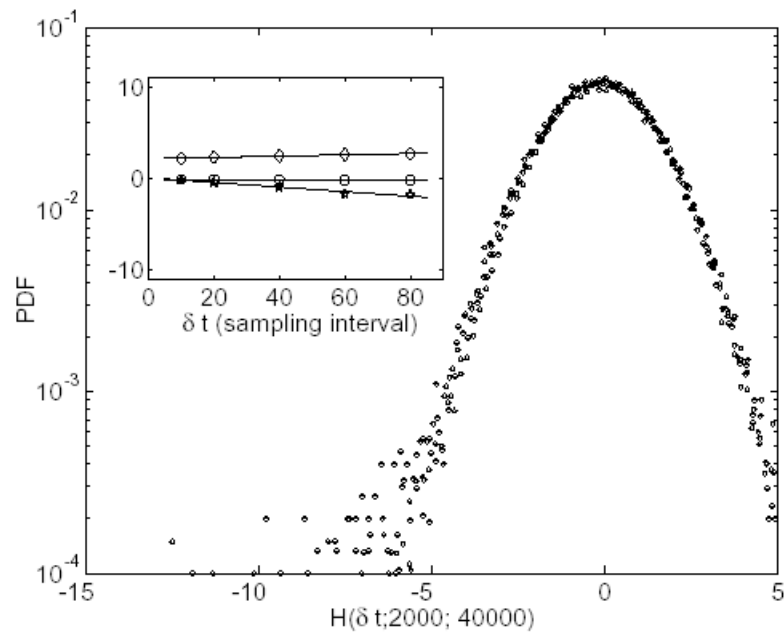


Fig. 1

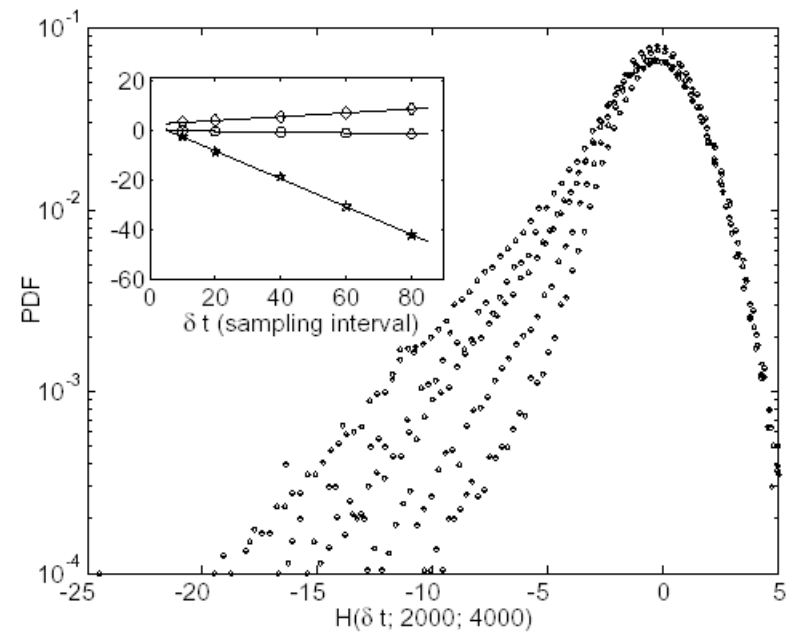


Fig. 2

## Consequences of record dynamics.

Statistics of quake times independent of underlying "noise mechanism".

- **Biology:** same intermittent dynamics in micro as in macro evolution.  
Decreasing transition rate.
- **Magnetic relaxation:** temperature independent creep rate
- **Spin glass:** exponential tails

## Conclusion/Summary

Considered spin-glasses, superconductors and biological evolution as typical complex systems.

Generic dynamics of complex systems:

- Non-stationary
- Intermittent record dynamics - quakes
- Rate of activity  $\sim 1/t$
- Stationary as function of  $\log(t)$



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