

# Bootstrap percolation and interacting particle systems with kinetic constraints

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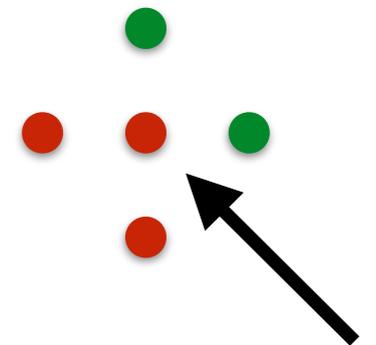
(University Roma Tre)

(a joint project with L. Mareche, R. Morris and C. Toninelli)

ERC grant “Malig”

# Bootstrap percolation: a first example

- At time  $t=0$  vertices of  $\mathbb{Z}^2$  are i.i.d. infected  with prob  $p$  and healthy  with prob  $1-p$ .
- At time  $t=1$  healthy vertices with at least 2 infected neighbours become infected.
- Iterate...



The bootstrap process is monotone (infection never heals)

## 2-neighbor process

(A. Holroyd)

Colors measure  
# iterations of  
BM necessary  
to infect a given  
vertex.

Black < Blu < Azur < Green < Yellow < White

# Some basic questions

**Q<sub>1</sub>:** Will the entire lattice become infected eventually ?

**A<sub>1</sub>:** Yes (van Enter '87)

**Q<sub>2</sub>:** How long does it take (w.h.p. or in mean) to infect the origin ?

**A<sub>2</sub>:** Approx  $\text{Exp}(\pi^2/18p)$  steps for  $p \ll 1$  (A. Holroyd '03)

**Q<sub>3</sub>:** Compute the scaling as  $n \rightarrow \infty$  of

$$p_c(n) := \inf\{p: \text{Prob}_p(\Lambda_n \text{ becomes eventually infected}) \geq 1/2\}$$

**A<sub>3</sub>:**  $p_c(n) = \Theta(1/\log n)$ .

# A general setting

★ A bootstrap rule  $\mathcal{U}$  is a collection  $(X_1, X_2, \dots, X_m)$  of finite subsets of  $\mathbb{Z}^2 \setminus \{0\}$ .

★  $A_t \subseteq \mathbb{Z}^2 \iff$  Set of infected sites at time  $t$ .

★  $\mathcal{U}$ -bootstrap process:

$$A_{t+1} = A_t \cup \{x \in \mathbb{Z}^2 : x + X \subseteq A_t \text{ for some } X \in \mathcal{U}\}$$

$[A]_{\mathcal{U}} := \bigcup_{t \geq 0} A_t$  the “closure of  $A$ ” (or final infected set).

# Examples

The Duarte  
model

$\mathcal{U}$  = family of 2-subsets of  
the South, North and West neighbours  
of the origin

The “South or West”  
process  
(known as the East model 😊)

$\mathcal{U}$  = family of 1-subsets of  
the South and North neighbours

The North-East process

$\mathcal{U}$  = North & East neighbours

# Critical Probability and Critical Length

- Let  $\Lambda_n$  be the  $n$ -torus in  $\mathbb{Z}^2$ .
- Take the initial infected set  $A := A_{t=0}$   $p$ -random.
- Critical probability:

$$p_c(n) = \inf \{ p: \text{Prob}_p( [A]_{\mathcal{U}} = \Lambda_n ) \geq 1/2 \}$$

The scaling of  $p_c(n)$  as  $n \rightarrow \infty$  is dictated by the action of the  $\mathcal{U}$ -process on discrete half-planes (Bollobas, Smith and Uzzell '15)

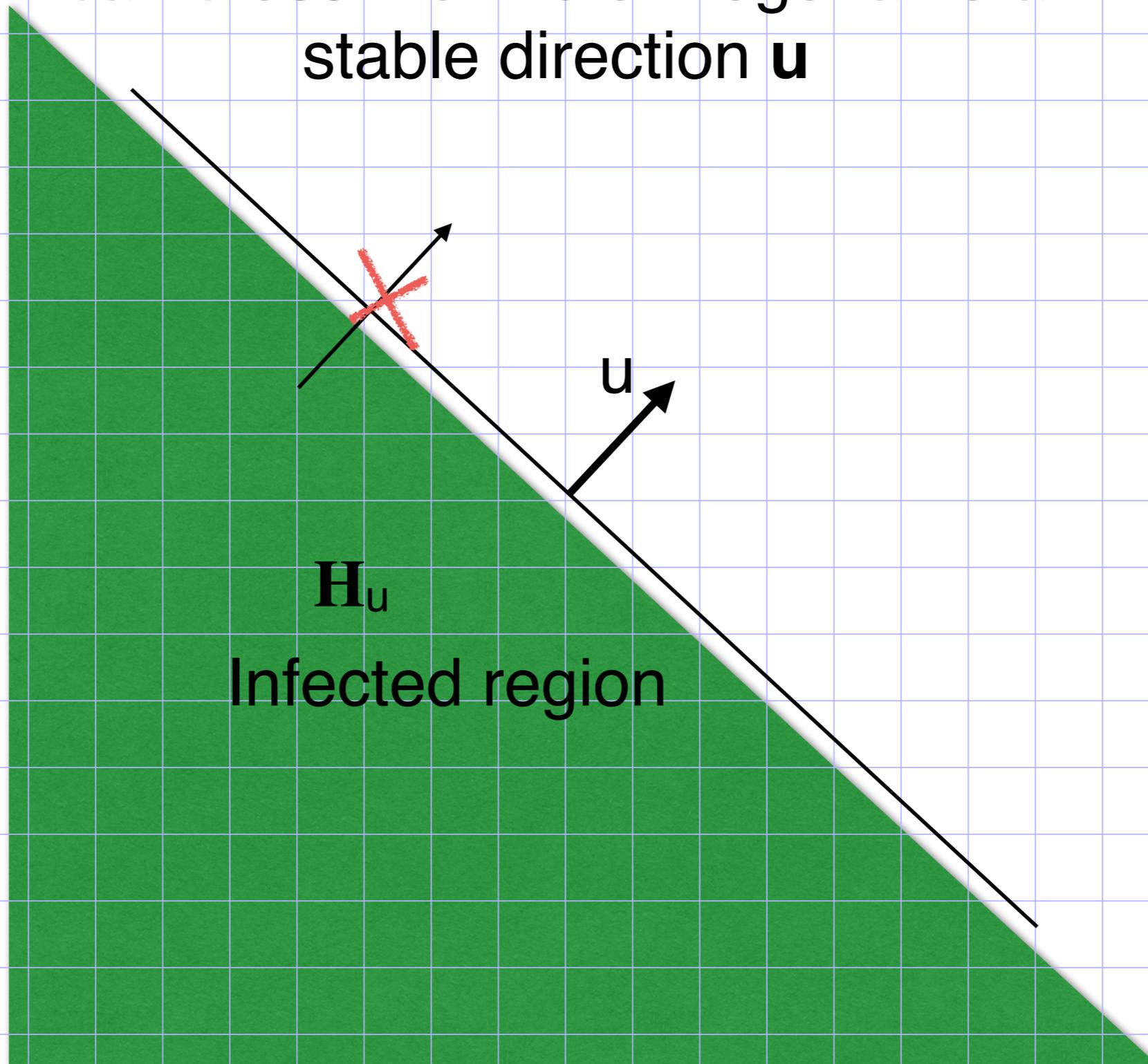
# Stable Directions

A direction  $u \in S^1$  is **stable** if the “ $\mathcal{U}$ -closure” of the half-plane

$$\mathbf{H}_u = \{x \in \mathbb{Z}^2: \langle x, u \rangle < 0\}$$

is  $\mathbf{H}_u$  itself:  $[\mathbf{H}_u]_{\mathcal{U}} = \mathbf{H}_u$ .

No infection  
can cross the line orthogonal to a  
stable direction  $\mathbf{u}$



$H_u$

Infected region

# A general classification

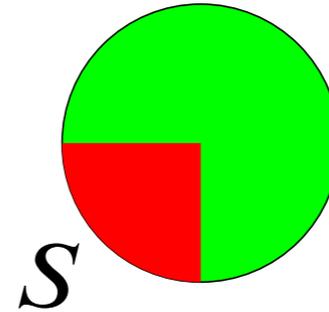
(Bollobas-Smith-Uzzell '15)

Let  $S$  be the set of stable directions. The  $\mathcal{U}$ -process is:

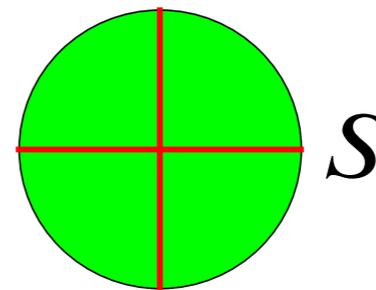
- **Supercritical** :  $\exists$  open semicircle  $S$  with no stable directions.
- **Critical** : every semicircle is such that  $S \cap S \neq \emptyset$  and  $\exists$  one semicircle with finite intersection with  $S$ .
- **Subcritical** : every semicircle has  $\infty$  intersection with  $S$ .

# Examples

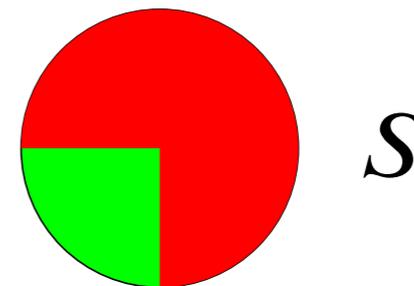
The South-or-West process  
(supercritical)



2-neighbour process (critical)



The North-East process (subcritical)



# A universality result for $p_c(n)$

(Bollobas-Smith-Uzzell '15, Balister-Bollobas-Przykucki-Smith '16)

Theorem Let  $\mathcal{U}$  be an update family.

$\mathcal{U}$  supercritical  $\Rightarrow p_c(n) = 1/n^{\Theta(1)}$

$\mathcal{U}$  critical  $\Rightarrow p_c(n) = 1/(\log n)^{\Theta(1)}$

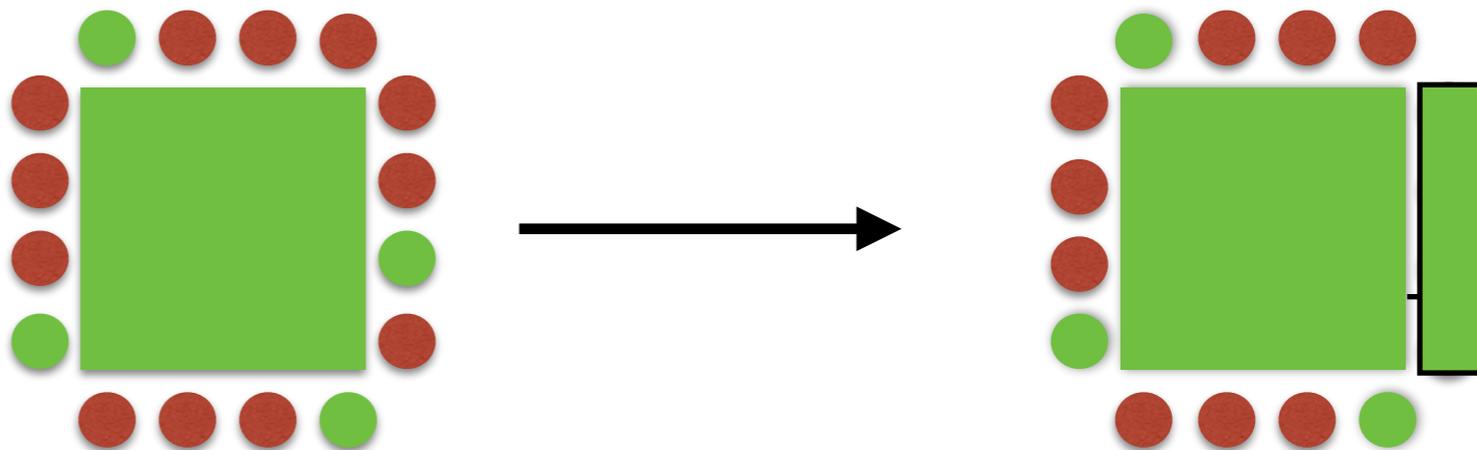
$\mathcal{U}$  subcritical  $\Rightarrow \liminf p_c(n) > 0.$

# A refinement for critical systems

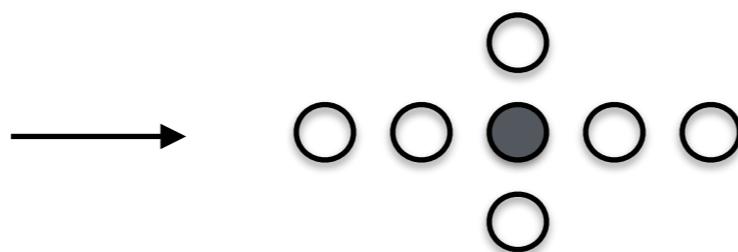
- In '16 Bollobas, Duminil-Copin, Morris and Smith introduced the notion of “difficulty”  $\alpha$  for critical systems and proved  $p_c(n) = \Theta(1/(\log n)^{1/\alpha})$ .
- Roughly,  $\mathbf{u} \in S^1$  has difficulty  $\alpha$  if, with the help of  $\alpha$  extra infected sites, the half-plane  $H_{\mathbf{u}}$  is able to grow in the  $\mathbf{u}$  direction.

Definition:  $\alpha(\mathcal{U}) = \min_S \max_{u \in S} \alpha(u)$

2-neighbor process  
 $\alpha = 1$

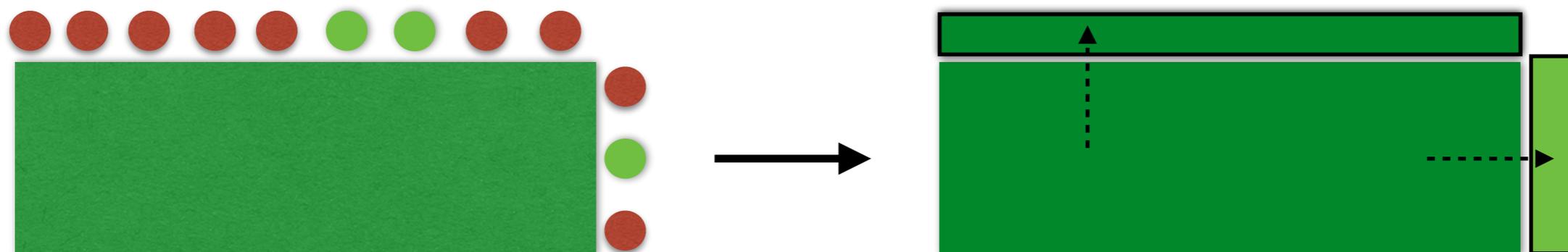


Anisotropic model: at least  
 3 infected sites among



$\alpha_v = 2$  in the  $\updownarrow$  direction  
 $\alpha_h = 1$  in the  $\leftrightarrow$  direction

$$\Rightarrow \alpha = \min(\alpha_v, \alpha_h) = 1$$



# Critical Lengths

Fix  $p \ll 1$  and let  $L_c = L_c(p)$  be such that

$$p_c(L_c) = p$$

- $\mathcal{U}$  supercritical :  $L_c(p) = 1/p^{\Theta(1)}$  as  $p \searrow 0$ .
- $\mathcal{U}$  critical :  $L_c(p) = \Theta(\text{Exp}(1/p^\alpha))$  as  $p \searrow 0$ .

# Kinetically constrained models

- ★ Fix a bootstrap rule  $\mathcal{U}$  and  $0 < p < 1$ .

The updating rule :

- ◆ Each vertex which could be infected by one step of the  $\mathcal{U}$ -process becomes **infected** with rate  $p$  and **non-infected** with rate  $1-p$  independently of the others.
- ◆ All other vertices don't change their state.

# KCM: general features

- Reversible interacting particle systems on  $\mathbb{Z}^2$ .
- Reversible measure  $\mu = \text{product Bernoulli}(p)$
- Not monotone (**infection can heal**).
- Not necessarily ergodic (there could be blocked configurations)

# Main obstacles for a mathematical analysis

- KCMs are **not monotone**: more infected vertices in the system may have **unpredictable** consequences.
- Powerful tools like **FKG** inequalities or **Censoring** are no longer available.
- **Worst case analysis** (w.r.t. the initial condition) usually too rough.
- Coercive inequalities (e.g. Poincare' and Log-Sobolev) behave **very differently** from e.g. the “high temperature Ising model”

# Origins of KCMs

(nice review by Evans & Sollich)

G. Fredrickson and H. Andersen in '84 introduced KCMs as simple models featuring *glassy dynamics* at low temperature :

- *Trivial equilibrium.*
- *Broad spectrum of relaxation time scales.*
- *Heterogenous dynamics.*
- *Possible ergodicity breaking transition.*
- *Aging.*

In physical terms small values of the density  $p$  of the infected (facilitating) sites  $\longleftrightarrow$  *small temperature*

# The infection Time for the KCM

- $T(\mathcal{U}) = \inf \{t \geq 0: \text{the origin is infected at time } t\}$
- **Q1:** How does  $T(\mathcal{U})$  grow as  $p \searrow 0$  for the stationary process? Does it scale as  $L_c(p)$  ?
- General Bounds :
  - (a) W.h.p  $T(\mathcal{U}) \geq \varepsilon L_c(p)$  for some small  $\varepsilon > 0$  (**not sharp !!**)
  - (b)  $E_\mu(T(\mathcal{U})) < C \times (1/p) \times \text{“Relaxation Time”}$ .

# The Relaxation Time $T_{\text{rel}}(\mathcal{U})$

Quadratic form of the Markov generator of the KCM:

$$\mathcal{D}(f) = \sum_x \mu( c_x \text{Var}_x(f) )$$

$c_x$  = indicator of  $x$  being  
“infectable” by the BM

local variance=spin flip  
at  $x$

$$T_{\text{rel}} = \inf \{ \lambda : \text{Var}(f) \leq \lambda \mathcal{D}(f) \quad \forall f \}$$

A constrained  
Poincare' inequality

# Bounding the relaxation time

Lower bound: use the variational characterisation and find a good test function  $F$

$$\Rightarrow T_{\text{rel}} > \text{Var}(F)/\mathcal{D}(F)$$

Upper bound: find  $C$  s.t.  $\text{Var}(f) \leq C \mathcal{D}(f)$  for all  $f$

$$\Rightarrow T_{\text{rel}} \leq C$$

# Scaling of the infection time for some models

## 1-neighbor process

(Cancrini, M. Roberto, Toninelli '08  
Pillai & A. Smith '15)

$$E_{\mu}(T(\mathcal{U})) \asymp 1/p^{\Theta(1)}$$

explicit

## South-or-West process

(Chleboun, Faggionato, M. '15)

$$E_{\mu}(T(\mathcal{U})) \asymp p^{-\log(1/p)(\beta+o(1))}$$

$$(L_c(p) = 1/p^{\Theta(1)} !!)$$

## 2-neighbor process

(M & Toninelli '16)

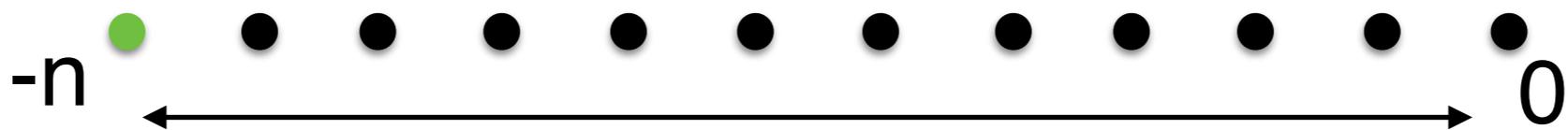
$$- E_{\mu}(T(\mathcal{U})) < (L_c)^{\beta(p)}$$

$$\beta(p) = O(\log \log L_c)$$

# On the $p^{-\Theta(\log 1/p)}$ growth for the S-or-W Model.

The system features “Logarithmic energy barriers”

- Consider the East process in 1D (infection goes from left to right BUT not viceversa !)
- Let Energy := # of infected vertices
- Assume that the first infected site to the left of  $x=0$  is at  $-n$



Theorem [*Chung, Diaconis, Graham '01*] At least  $\log_2(n)$  vertices must be infected before  $T(\mathcal{U})$ .

# Role of energy barriers

$$P_{\mu}(T(\mathcal{U}) < t) \leq O(t) \times \mu(\exists \log_2(n) \text{ infected sites})$$

$$\leq O(t) \times p^{\log_2(n)} \times n^{\log_2(n)}$$

Choose  $n \sim 1/p^{\varepsilon}$ ,  $\varepsilon \ll 1$ ,

( $1/p$  = equilibrium distance between 2 infected vertices)



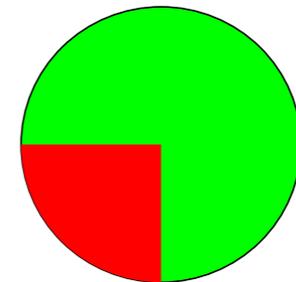
$$P_{\mu}(T(\mathcal{U}) < t) \leq O(t) \times \exp(-c(\varepsilon)\log(1/p)^2)$$

# A refined classification of supercritical systems

**Definition.** A supercritical  $\mathcal{U}$ -process is **rooted** if there exists two non-opposite stable directions. Otherwise it is **unrooted**.

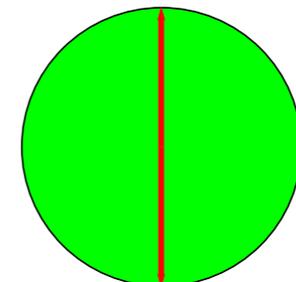
## Examples.

- **Rooted:** the S-or-W process.



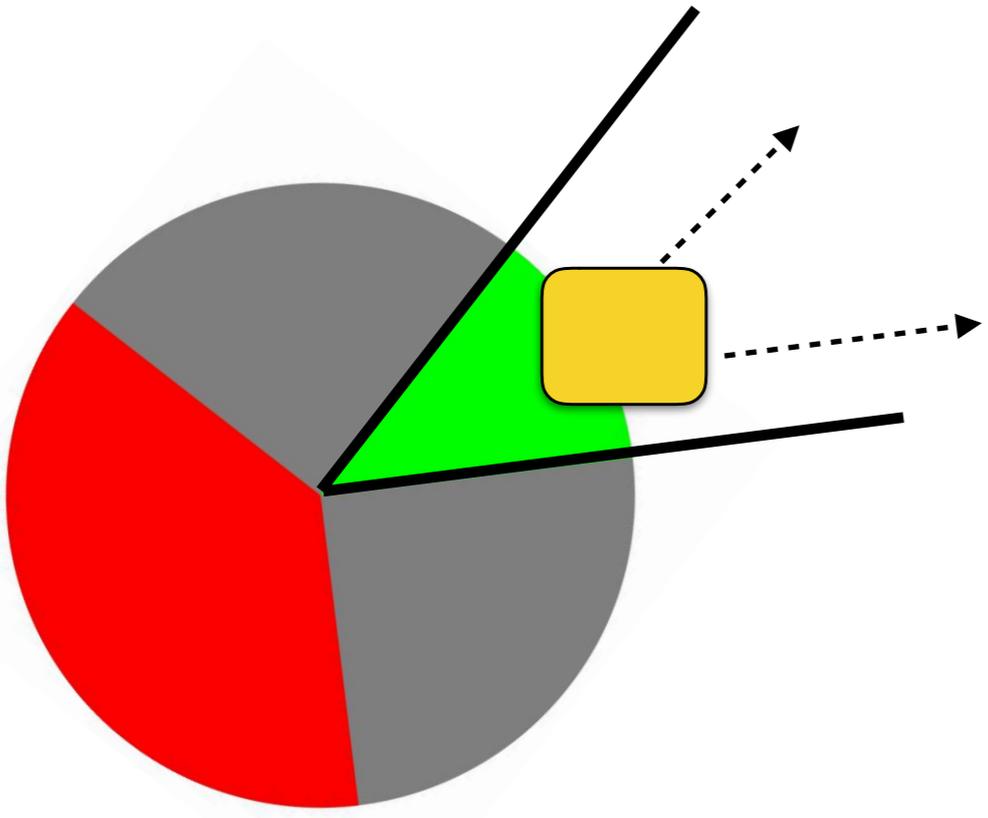
$S$

- **Unrooted:**  $\mathcal{U} \equiv$  an infected site between the left and right neighbours.

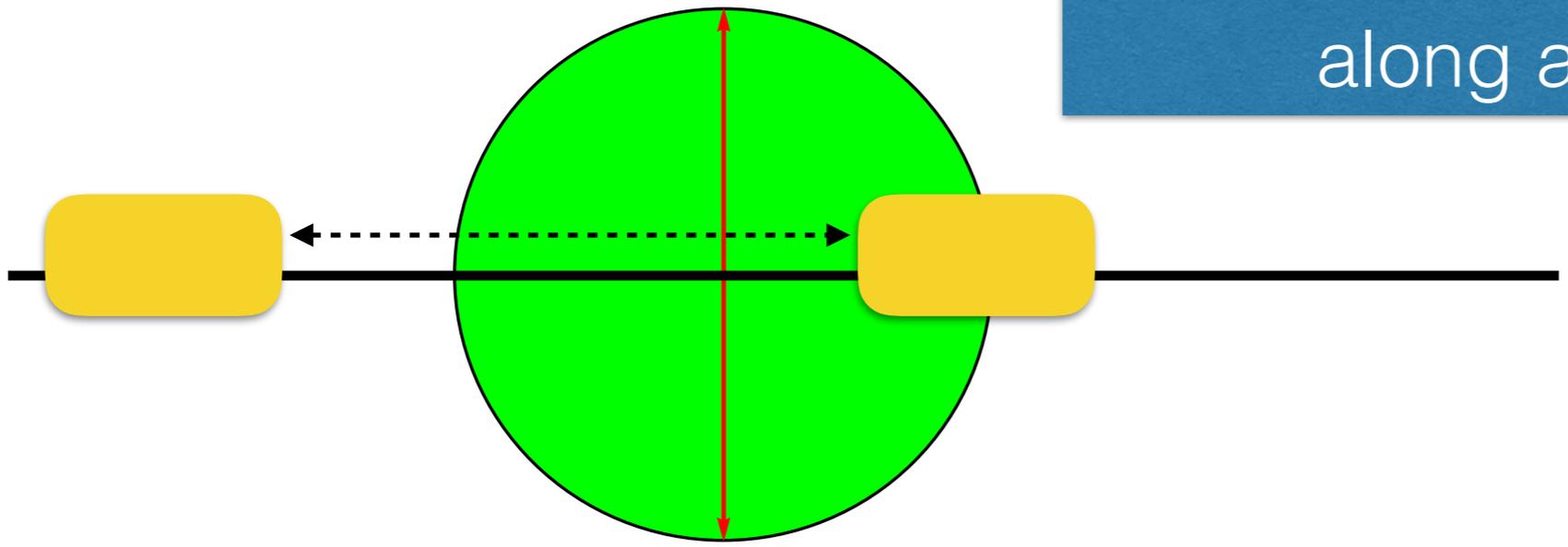


$S$

In a *rooted* supercritical system the droplet of infected sites can only move “forward” inside the cone delimited by black lines



In *unrooted* systems a large enough droplet can move back and forth along a given direction



# Growth of $T(\mathcal{U})$ for supercritical systems

(with R.Morris, C. Toninelli, L.Mareche)

Let  $\mathcal{U}$  be supercritical. As  $p \searrow 0$

$T(\mathcal{U}) = p^{-\Theta(1)}$  if  $\mathcal{U}$  is unrooted.

$T(\mathcal{U}) = p^{-\Theta(\log(1/p))}$  if  $\mathcal{U}$  is rooted.

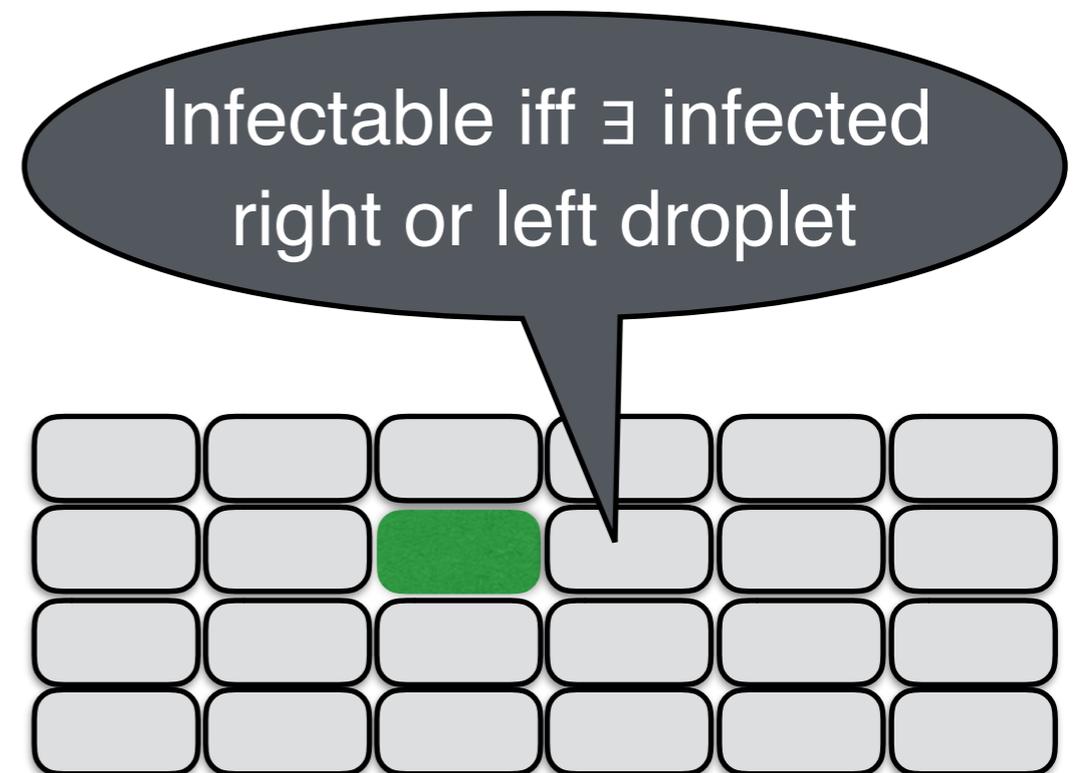
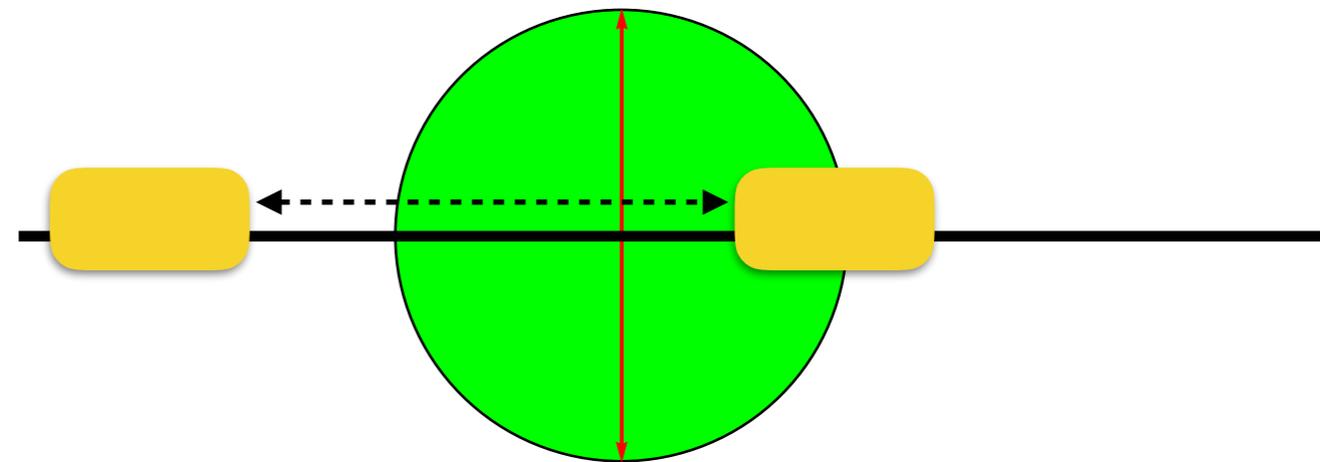


Unrooted case:  $T(\mathcal{U}) \leftrightarrow L_c(p)$

Rooted case:  $T(\mathcal{U}) \gg L_c(p)$

# The unrooted case

- ◆  $\exists$  a direction  $\mathbf{u}$  and a big enough rectangular droplet  $R$  parallel to  $\mathbf{u}$  which can shift in both directions  $\mathbf{u}$  and  $-\mathbf{u}$ .
- ◆ We can compare the  $\mathcal{U}$ -KCM to a **renormalised 1-neighbor** KCM model in which each vertex of  $\mathbb{Z}^2$  actually represents a **droplet  $R$** .



# Renormalised Droplet Model

## $\Leftrightarrow$ Original Model

In the renormalised model  $p_{\text{eff}} = p^{\text{Area}(\text{droplet})}$

For the renormalised droplet model

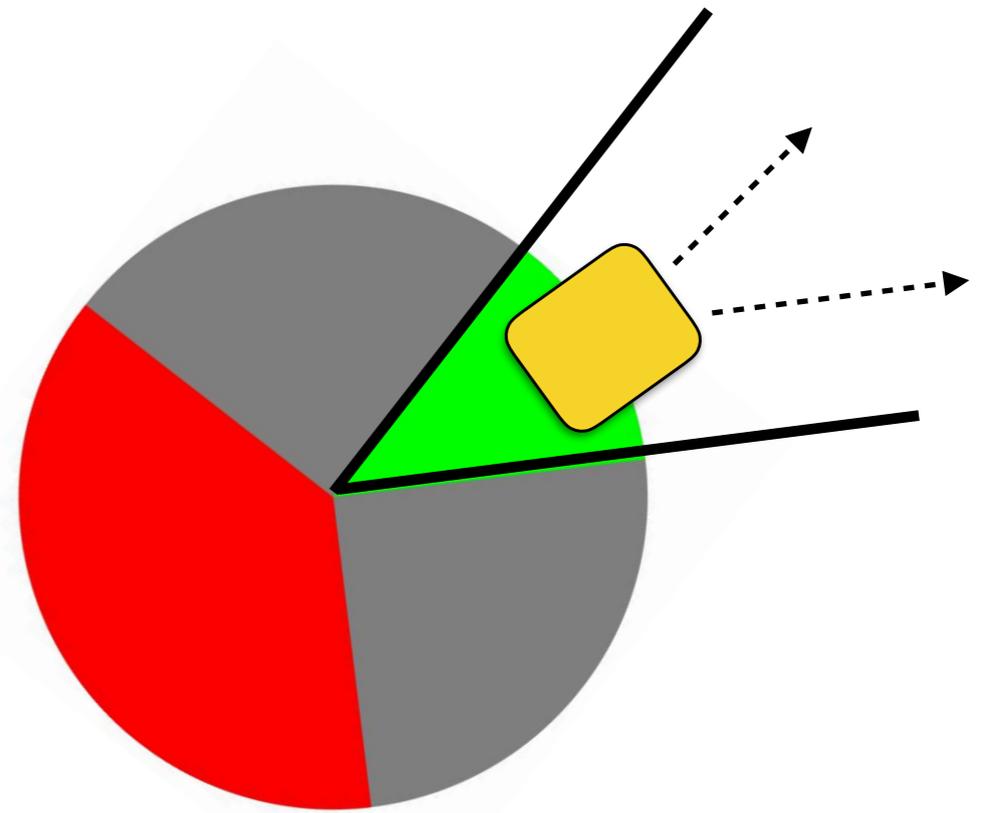
$$T_{\text{rel}}(\mathcal{U}_{\text{ren}}) \sim 1/p_{\text{eff}}^{\Theta(1)}$$

Comparisons of the Dirichlet forms of the renormalised and original Markov chains  $\Rightarrow$

$$T_{\text{rel}}(\mathcal{U}) = 1/p^{\Theta(1)}$$

# The rooted case

- (a) There exists a direction  $\mathbf{u}$  and a big enough rectangular droplet  $R$  parallel to  $\mathbf{u}$  which can shift in the  $\mathbf{u}$  direction.
- (b) L. Mareche has proved that there exist **logarithmic energy barriers**.
- (c) Again use renormalisation to compare the  $\mathcal{U}$ -KCM to an **effective** East model.



# Critical systems

Def. A critical system with difficulty  $\alpha$  is **rooted** if there exist two ***non-opposite stable*** directions with **difficulty**  $> \alpha$ . Otherwise it is **unrooted**

Theorem [M. Morris, Toninelli]

Let  $\mathcal{U}$  be unrooted critical.

$$c_1 L_c \leq E(T(\mathcal{U})) \leq L_c^\beta, \quad \beta = O(\log \log L_c).$$

Conjecture: Let  $\mathcal{U}$  be rooted critical.

$$T(\mathcal{U}) = \Theta(\exp(\beta/p^{2\alpha}))$$

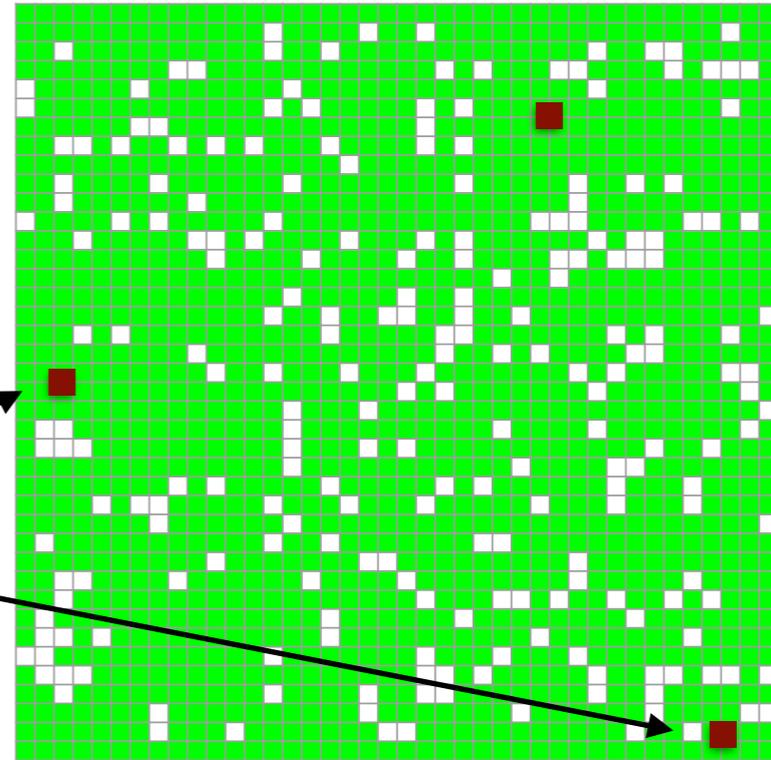
# Strategy of the proof

Def: a  $L \times L$  box is **good** if **every** column/row has an infected site.

If  $L \sim Cp^{-1} \times \log(1/p)$ ,  $C > 4$   
 $\Rightarrow P(\text{Good}) \sim 1$

Good boxes form a (unique)  $\infty$ -cluster with holes.

Among the good boxes there are (rare) super-good ones:

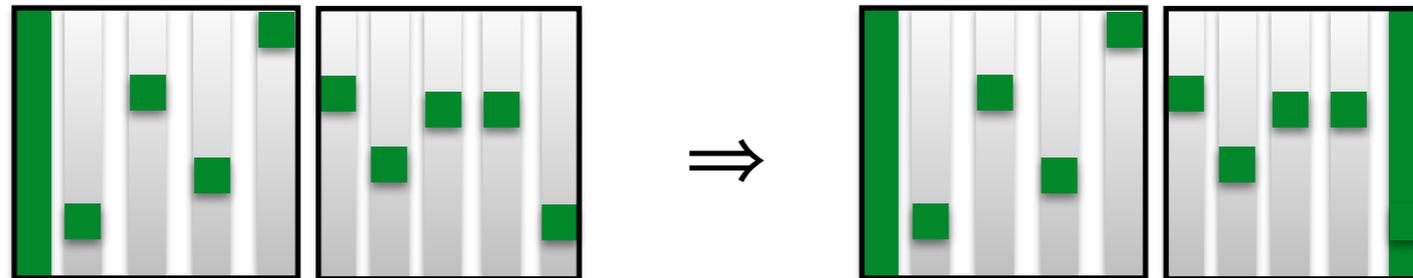


Def: A box is **super-good** if it is **good** and 1st row & column completely infected

$\text{Prob}(\text{SG}) \sim \text{Exp}(-1/p \times \text{polylog}(p))$

# Heuristics for $E(T(\mathcal{U})) \leq L_c^\beta$

A SG box can freely travel among good boxes



A. At  $t=0$  w.h.p. a SG box will be found at distance

$$L \sim 1/P(\text{SG})^{1/2}$$

B. In a diffusive time the SG box will move next to the origin.

C. In another time  $\propto \text{poly}(L)$  the bootstrap map will infect the origin.

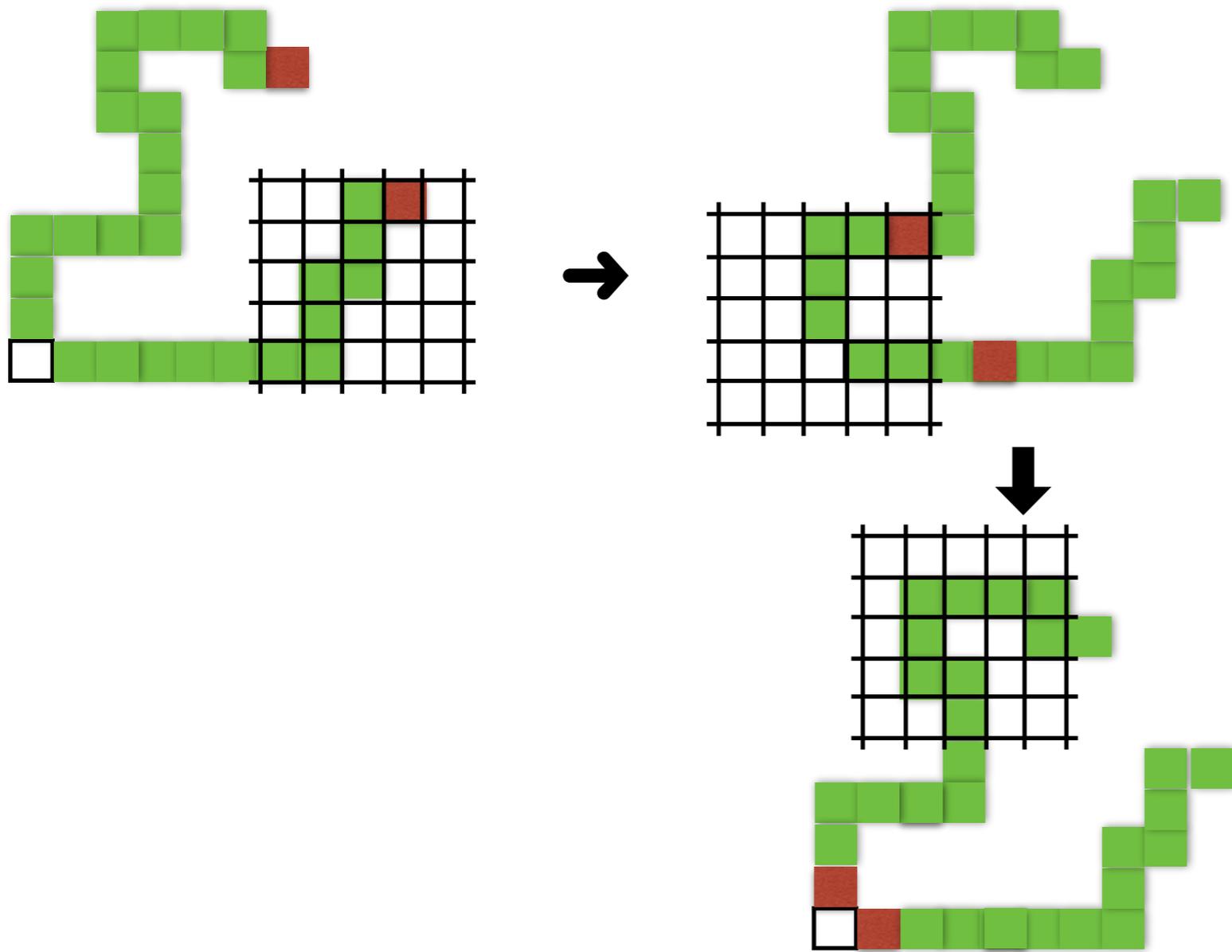
# Main Obstacles

1. The environment of the good and super-good boxes evolves in time;
2. No monotonicity argument to move infos at  $t=0$  to  $t > 0$ .

Solution:

Bound from above the relaxation time



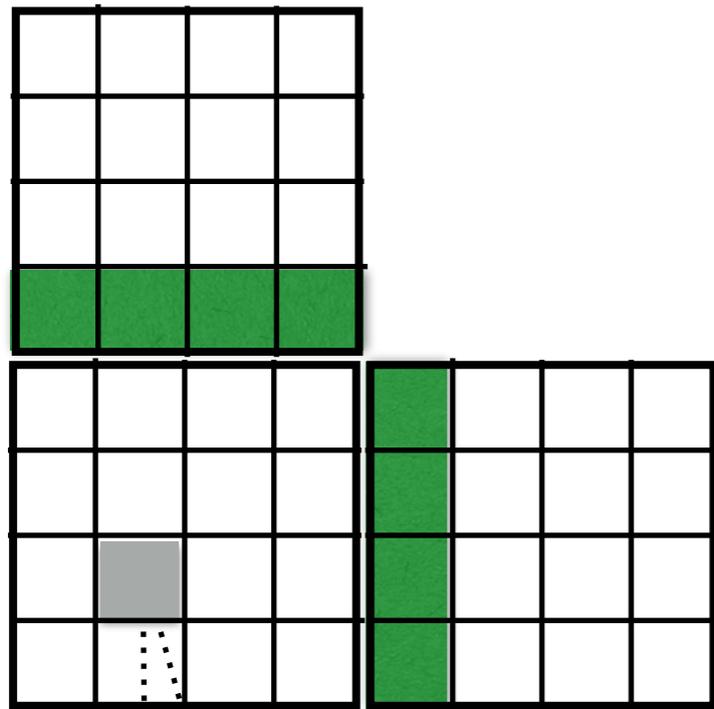


Using canonical paths techniques for reversible Markov chains

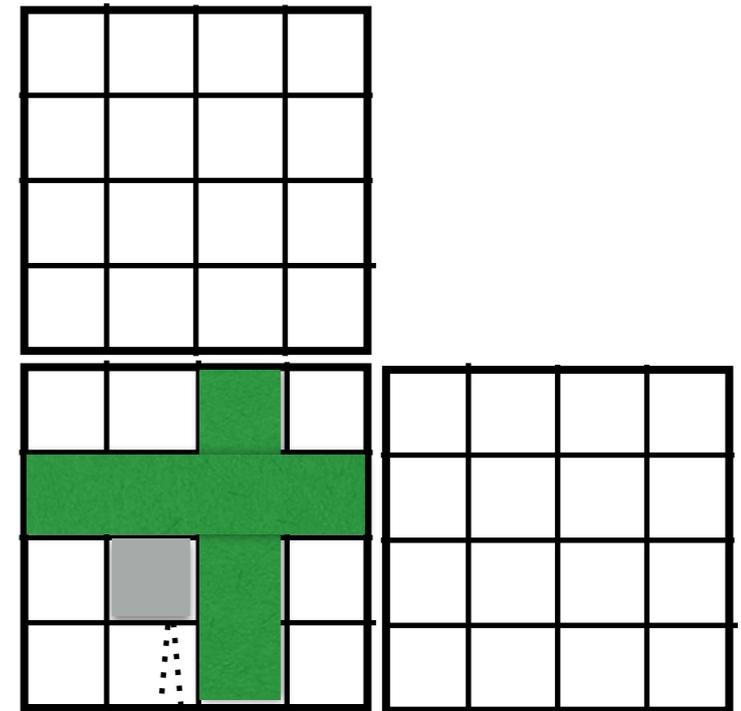
the SG boxes are there

$$\text{Var}(f) \leq \widetilde{\exp}(1/p) \sum_{\text{boxes}} \mu(\lambda_x \text{Var}_{B_x}(f))$$

# Conclusion



Want to flip here



Flip here is now legal

$$\Rightarrow \text{Var}(f) \leq \widetilde{\exp}(1/p) \sum_{\text{boxes}} \mu(c_x \text{Var}_{B_x}(f))$$

the true constraints

# Summary

- Characteristic **time scales** for KCM may scale very differently from the characteristic **length scales** of the bootstrap percolation model.

The new feature of KCM is the possible presence of “**energy barriers**” that the dynamics has to overcome.

A more **refined classification** seems to be able to characterise systems for which **energy barriers matter**.

A **universality** result on the scaling of the infection time emerges.



**Thank you**