# Unbiased on lattice domain growth

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## Motivation

Domain growth is an inherent feature of many biological systems. One such example is in the movement and proliferation of melanoblasts, the precursor to pigment producing melanocytes, on a growing embryo. If melanoblasts fail to fully populate the spatial domain, small regions of skin will contain no pigment [Mort et al., 2016]. This disease is called piebaldism, a proxy for many neurocristopathies — a class of pathogies caused by the similar failure of neural crest cells to proliferate and migrate during embryonic growth. Models for domain growth are of importance in order to simulate and predict biological and physical systems such as that of piebaldism.

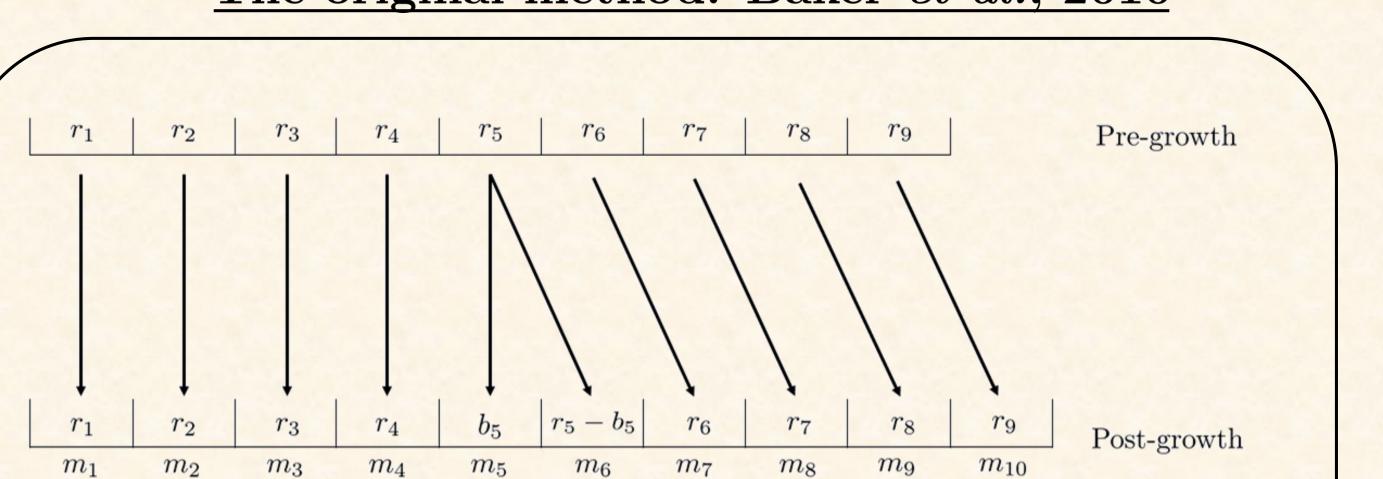
We demonstrate that a previously employed method for on-lattice domain growth causes a build up of particles at the boundaries of the spatial domain in "low-diffusion regimes".



A piebald mouse with a white belly spot. Image reproduced with permission from Mort *et al.*, 2016.

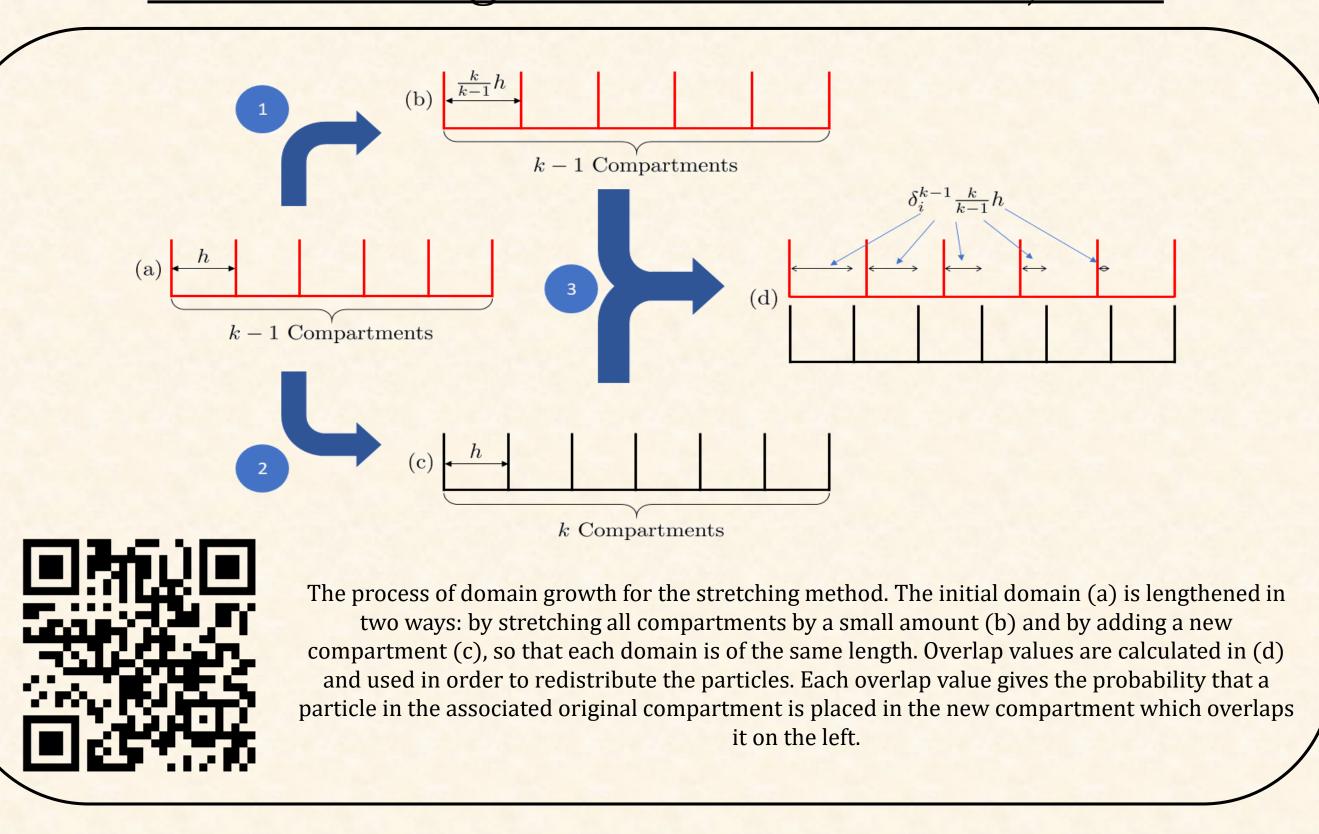
#### The original method: Baker et al., 2010

#### The stretching method: Smith et al., 2019



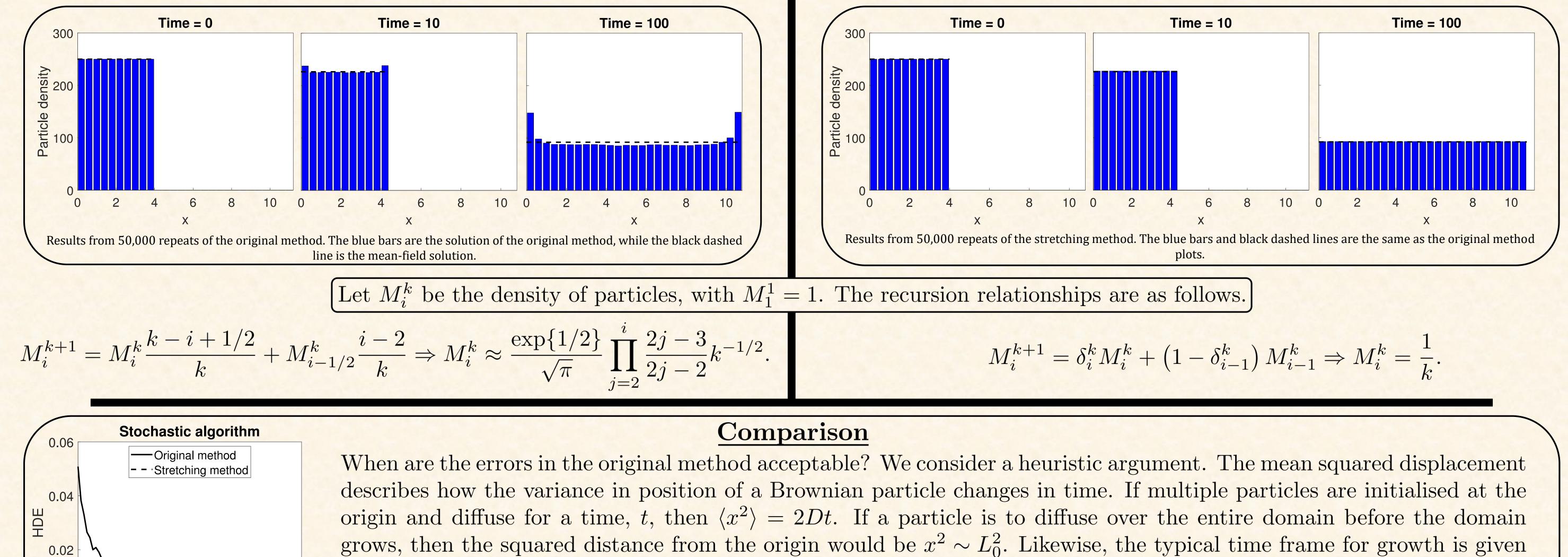
A schematic showing a possible splitting event when there are nine pre-growth compartments (compartments identified by the subscripts on the particle numbers). Pre-growth particle numbers are denoted  $r_i$ . In this case, compartment 5 is chosen to divide, and its contents are split binomially with probability of success 1/2 ( $b_5 \sim Bin(r_5, 1/2)$ ) between the compartment in the original position (compartment 5 in this case) and the new one that is created to its right. The compartments originally numbered 6, 7, 8 and 9 are moved one position to the right and become compartments 7, 8, 9 and 10, respectively. Post-growth particle numbers are denoted by the  $m_i$ 's.





**On-lattice domain growth can introduce** unexpected particle accumulation at the

# boundaries



by setting  $t \sim 1/\rho$ . Substituting these into the expression for the mean squared displacement yields  $D \sim L_0^2 \rho$ . Therefore,

minimise 0.5 Comparing the histogram distance error (HDE) for a non-dimensional stochastic simulation of the original and stretching methods

we say we are in a high diffusion parameter regime when  $D > L_0^2 \rho$ .

To the left is the result of several non-dimensional simulations of both the original and stretching methods, where the non-dimensional diffusion parameter is  $D^* = D/(L_0^2 \rho)$ . For  $D^* > 1$ , the two methods have similar histogram distance errors, which verifies the results of the heuristic argument.

### **References**

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