

# Comparing Compartment and Agent-based Models

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August 2, 2017

Thesis work with:

William F. Eddy (Chair)

Joel Greenhouse

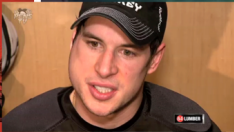
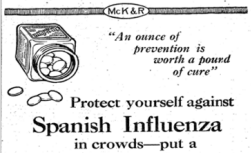
Howard Seltman

Cosma Shalizi

Samuel L. Ventura

Goal: Combine two good models into a better one

Studying infectious disease is important



# Compartment vs. Agent-based Models

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# Compartment models (CMs) describe how individuals evolve over time

Assumptions (Anderson and May 1992) :

1. Homogeneity of individuals

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Assumptions (Anderson and May 1992) :

1. Homogeneity of individuals

2. Law of mass action

$$I(t+1) \propto I(t)$$

# Agent-based models (AMs) simulate the spread of disease

Assumptions (Helbing 2002):

1. Heterogeneity of agents

# Agent-based models (AMs) simulate the spread of disease

## Assumptions (Helbing 2002):

1. Heterogeneity of agents
2. Model adequately reflects reality



## CMs

- Equation-based
- Computationally fast
- Homogeneous individuals
- No individual properties

## AMs

- Simulation-based
- Computationally slow
- Heterogeneous individuals
- Individual properties

(Bobashev 2007, Banos 2015, Wallentin 2017)

- ad hoc approaches
- perspective from non-statisticians

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**Goal: Create a statistically justified hybrid model**

## Current Work

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# There are two main avenues of improvement

1. Quantifying how similar CMs and AMs are
2. Speeding up AM run-time

(Kermack and McKendrick 1927)

$$\left\{ \begin{array}{lcl} \frac{dS}{dt} & = & -\frac{\beta SI}{N} \\ \frac{dI}{dt} & = & \frac{\beta SI}{N} - \gamma I \\ \frac{dR}{dt} & = & \gamma I \end{array} \right.$$

- $\beta$  – rate of infection
- $\gamma$  – rate of recovery
- $N$  – total population size

(Kermack and McKendrick 1927)

$$\left\{ \begin{array}{l} \frac{\Delta S}{\Delta t} = -\frac{\beta SI}{N} \\ \frac{\Delta I}{\Delta t} = \frac{\beta SI}{N} - \gamma I \\ \frac{\Delta R}{\Delta t} = \gamma I \end{array} \right.$$

- $\beta$  – rate of infection
- $\gamma$  – rate of recovery
- $N$  – total population size

$$\hat{S}(t+1) = \hat{S}(t) - s_t$$

$$\hat{R}(t+1) = \hat{R}(t) + r_t$$

$$\hat{I}(t+1) = N - \hat{S}(t+1) - \hat{R}(t+1),$$

with

$$s_{t+1} \sim \text{Binomial} \left( \hat{S}(t), \frac{\beta I(t)}{N} \right)$$

$$r_{t+1} \sim \text{Binomial} \left( \hat{I}(t), \gamma \right).$$



For an agent  $x_n(t)$ ,  $n = 1, 2, \dots, N$ , the forward operator for  $t > 0$  is

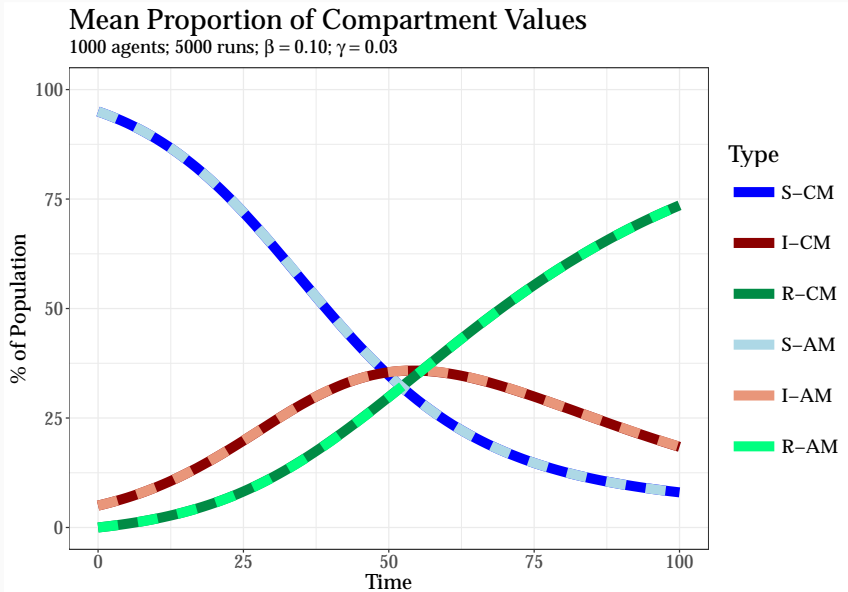
$$x_n(t+1) = \begin{cases} x_n(t) + \text{Bernoulli}\left(\frac{\beta I(t)}{N}\right) & \text{if } x_n(t) = 1 \\ x_n(t) + \text{Bernoulli}(\gamma) & \text{if } x_n(t) = 2 \\ x_n(t) & \text{otherwise} \end{cases}.$$

where  $x_n(t) = k$ ,  $k \in \{1, 2, 3\}$  corresponds to state S, I, and R, respectively

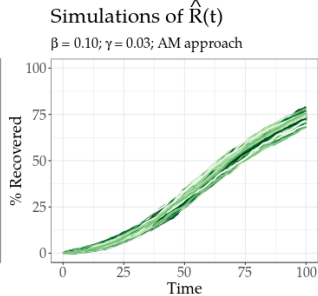
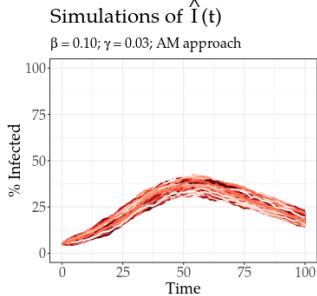
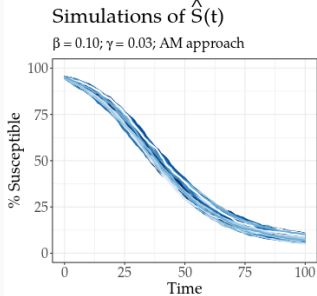
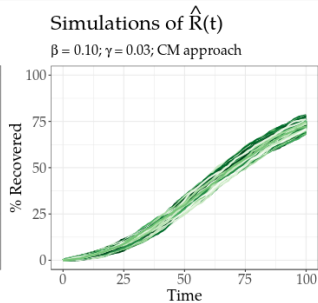
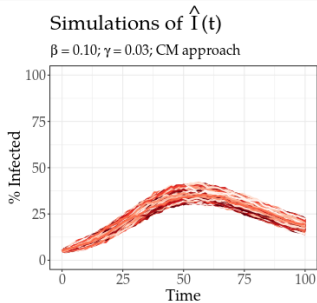
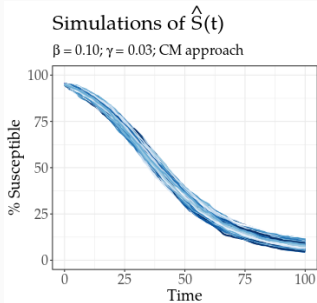
Let the **aggregate total** in each compartment be

$$\hat{X}_k(t) = \sum_{n=1}^N \mathcal{I}\{x_n(t) = k\}$$

# The means overlap



# The distributions look the same



## Theorem

Let the CM and AM be as previously described. Then for all  $t \in \{1, 2, \dots, T\}$ ,

$$\hat{S}(t) \stackrel{d}{=} \hat{X}_S(t) \tag{1}$$

$$\hat{I}(t) \stackrel{d}{=} \hat{X}_I(t)$$

$$\hat{R}(t) \stackrel{d}{=} \hat{X}_R(t).$$

## Theorem

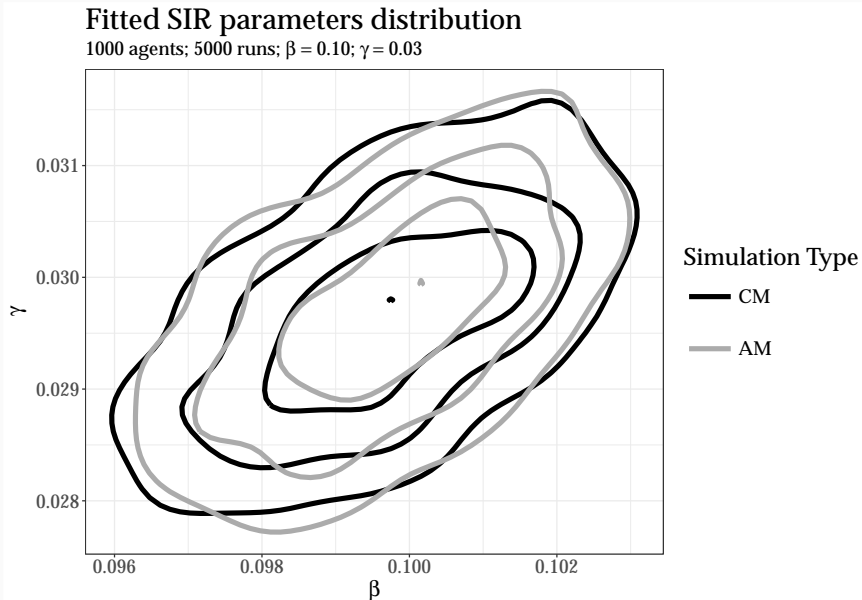
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We can compare CM/AM pairs and AM/AM pairs by fitting the underlying model



## AMs are appealing because they can be run multiple times

- Simulate an epidemic en masse!
- A **run** - same initial parameters, different random numbers
- Runs ( $L$ ) are independent of one another  $\implies$  **parallelization**
- Roughly, the variance of compartments  $\downarrow$  when  $N, L \uparrow$

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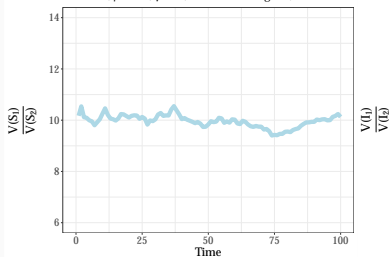
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**Goal:** Improve computation time without sacrificing statistical details

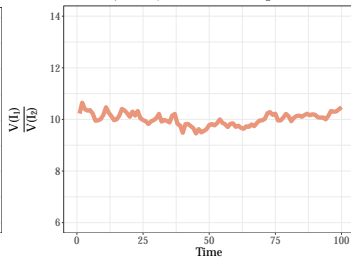


# There is a tradeoff between the number of agents and number of runs

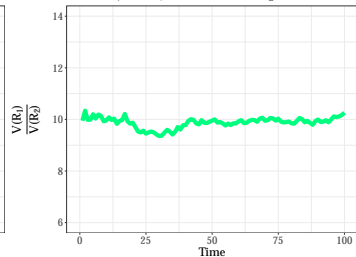
**Ratio of Variance of # Susceptibles**  
5000 runs;  $\beta = 0.10$ ;  $\gamma = 0.03$ ; Model 1–1000 agents, Model 2–100



**Ratio of Variance of # Infected**  
5000 runs;  $\beta = 0.10$ ;  $\gamma = 0.03$ ; Model 1–1000 agents, Model 2–100



**Ratio of Variance of # Recovered**  
5000 runs;  $\beta = 0.10$ ;  $\gamma = 0.03$ ; Model 1–1000 agents, Model 2–100



## The calculations show that the variance scales

- Note that for a given  $\beta$  and  $\gamma$ , if  $\frac{S_1(0)}{N_1} = \frac{S_2(0)}{N_2} \implies \frac{S_1(t)}{N_1} = \frac{S_2(t)}{N_2}$

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- $V \left[ \hat{S}(t+1) \right] = S(t)(1-p_t)p_t + (1-p_t)^2 V \left[ \hat{S}(t) \right]$

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$$\begin{aligned} \frac{V\left[\frac{1}{L_1} \sum_{\text{runs } \ell} \frac{\hat{S}_1(t)}{N_1}\right]}{V\left[\frac{1}{L_2} \sum_{\text{runs } \ell} \frac{\hat{S}_2(t)}{N_2}\right]} &= \frac{L_2 N_2^2}{L_1 N_1^2} \cdot \frac{V[\hat{S}_1(t)]}{V[\hat{S}_2(t)]} \\ &= \frac{L_2 N_2}{L_1 N_1}. \end{aligned}$$

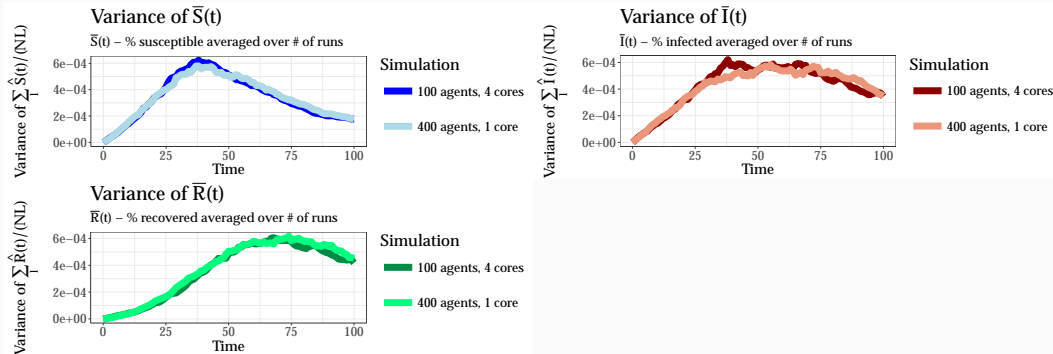
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We can replace agents with runs!

# Through parallelization, we can get a speed-up without losing statistical information



Simulation 1 (100 agents, 4 cores, 100 times): 3:30 minutes

Simulation 2 (400 agents, 1 core, 100 times): 4:05 minutes

## Future work

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## There is more work to be done: short-term

- Implementation of current methods in FRED
  - FRED - an open source, supported, flexible AM
  - Incorporate different levels of homogeneity
    1. Independent agents
    2. Agents go to one other activity (school, work, neighborhood)
    3. Multiple activities
  - Compare CM and AM parameters empirically
- Empirically determine when different regions can be combined

Thank you!

Questions?