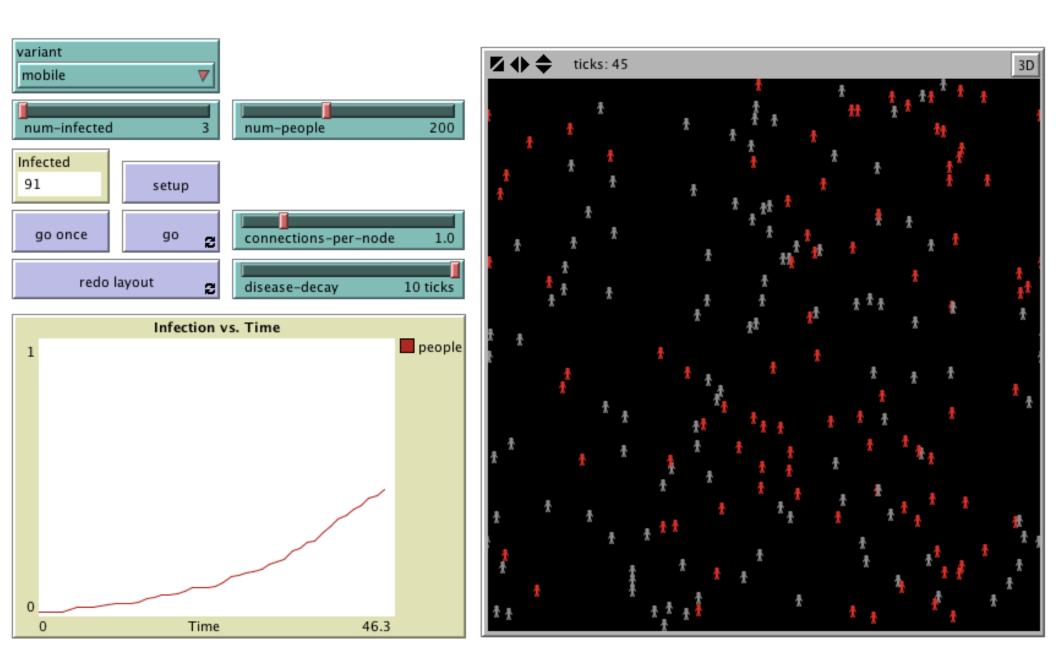
Analyzing ABMs

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Now that I have an Agent-Based Model how can I use it?

The Spread of Disease

- If a disease infects 5 people before the original person recovers (r0 = 5), then in three steps of the infection, the disease will infect 156 people (fast growth).
- On the other hand, the number of possible infections will quickly tail off because the disease may run out of new hosts (resource limits).
- Other Complications
 - No truly uniform mixing
 - Different susceptibilities to disease
 - Environmental interactions



Spread of Disease

An Initial Investigation

- What if we decide to investigate how population density affects the spread of disease?
- A couple of researchers decide to run experiments to investigate this
- We examine how population density affects the time to 100% infection

Different Results

First Run

Population	50	100	150	200
Time to 100% Infection	419	188 Second Run		127
Population	50	100	150	200
Time to 100% Infection	305	263	118	126
	Maybe we r	need to do	more runs?	

Raw Data

Population	Run 1	Run 2	Run 3	Run 4	Run 5	Run 6	Run 7	Run 8	Run 9	Run 10
50	419	365	305	318	323	337	432	380	430	359
100	188	263	256	205	206	205	201	181	202	231
150	169	118	163	146	143	167	137	121	140	140
200	127	126	113	111	133	129	109	101	105	133

The Necessity of Multiple Runs

- Since many of these models are stochastic, we need to run the model several times to understand the behavior
- There are several different ways to execute multiple runs in NetLogo or in any ABM platform

Within the Model

```
repeat 10 [
set num-people 50
setup
while [count turtles with [not infected?] > 0]
  [go]
 print ticks
```

	Experiment		Experiment			
Experiment name experi	ment	Experiment name population	on-density			
Vary variables as follows	(note brackets and quotation marks):	Vary variables as follows (no	ote brackets and quotation marks)			
["connections-per-node	" 1]	["variant" "mobile"]				
["disease-decay" 10]		["connections-per-node"	4.1]			
["num-infected" 3]		["num-people" [50 50 200]]]			
["variant" "mobile"]		["num-infected" 1]				
		["disease-decay" 0]				
Either list values to use, for exampl ["my-slider" 1 2 7 8]	le:	Either list values to use, for example: ["my-slider" 1 2 7 8]				
or specify start, increment, and end		or specify start, increment, and end, for	or example:			
["my-slider" [0 1 10]] (note addition to go from 0, 1 at a time, to 10.	nal brackets)	["my-slider" [0 1 10]] (note additional to go from 0, 1 at a time, to 10.	["my-slider" [0 1 10]] (note additional brackets)			
	-pxcor, max-pycor, min-pycor, random-seed.		cor, max-pycor, min-pycor, random-seed.			
Repetitions 1		Repetitions 10				
run each combination this many tir	nes	run each combination this many times				
Measure runs using these	reporters:	Measure runs using these re	eporters:			
one reporter per line; you may not	split a reporter	one reporter per line; you may not spl	it a reporter			
across multiple lines	spin a reporter	across multiple lines	it a reporter			
✓ Measure runs at every	step	Measure runs at every ste	ep			
if unchecked, runs are measured or	nly when they are over	if unchecked, runs are measured only				
Setup commands:	Go commands:	Setup commands:	Go commands:			
setup	go	setup	go			
► Stop condition:	▶ Final commands:	▶ Stop condition:	▶ Final commands:			
the run stops if this reporter becon		the run stops if this reporter becomes				
Time limit 0		Time limit 0				
stop after this many steps (0 = no I	imit)	stop after this many steps (0 = no limi	stop after this many steps (0 = no limit)			
	Cancel OK)	Cancel			

ОК

Raw Data

BehaviorSpace results (NetLogo 5.3.1)						
Spread of Disease.nlogo						
population-density						
09/26/2016 15:07:32:	:848 -0400					
min-pxcor	max-pxcor	min-pycor	max-pycor			
-20	20	-20	20			
[run number]	variant	connections-per- node	num-people	num-infected	disease-decay	[step]
2	"mobile"	4.1	50	1	0	
7	"mobile"	4.1	50	1	0	
5	"mobile"	4.1	50	1	0	
8	"mobile"	4.1	50	1	0	
6	"mobile"	4.1	50	1	0	
1	"mobile"	4.1	50	1	0	
4	"mobile"	4.1	50	1	0	
11	"mobile"	4.1	100	1	0	
10	"mobile"	4.1	50	1	0	
3	"mobile"	4.1	50	1	0	

Summary Statistics

Summary statistics provide us with a way of characterizing a large data set.

Easy to do in R.

Population	Mean	Std. Dev.
50	366.8	47.39385802
100	213.8	27.40154091
150	144.4	17.65219533
200	118.7	12.12939497

Mean

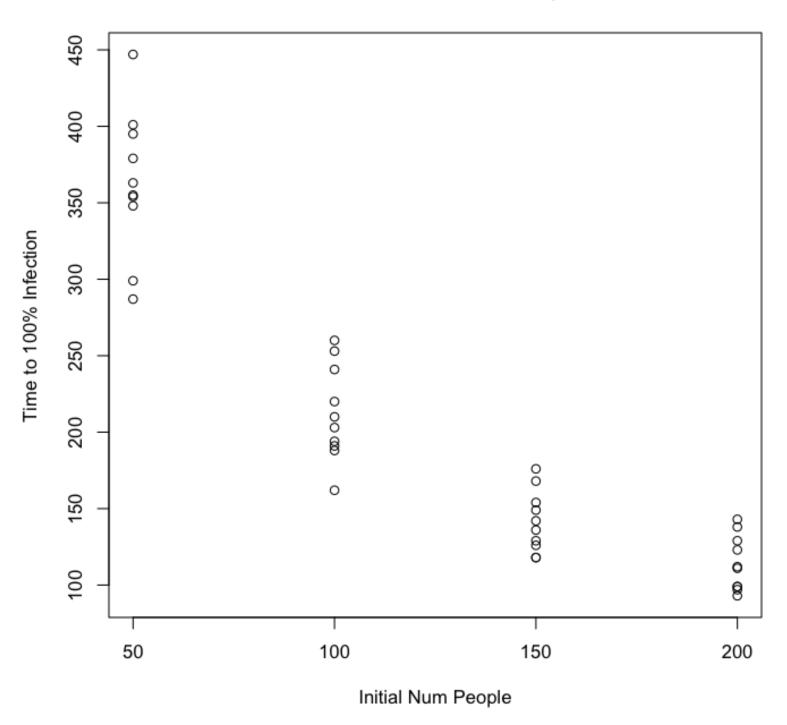
Standard Deviation

$$\mu = \frac{x_1 + x_1 + \dots + x_n}{N} \qquad \sigma = \sqrt{\frac{1}{N}[(x_1 - \mu)^2 + (x_2 - \mu)^2 + \dots + (x_n - \mu)^2]}$$

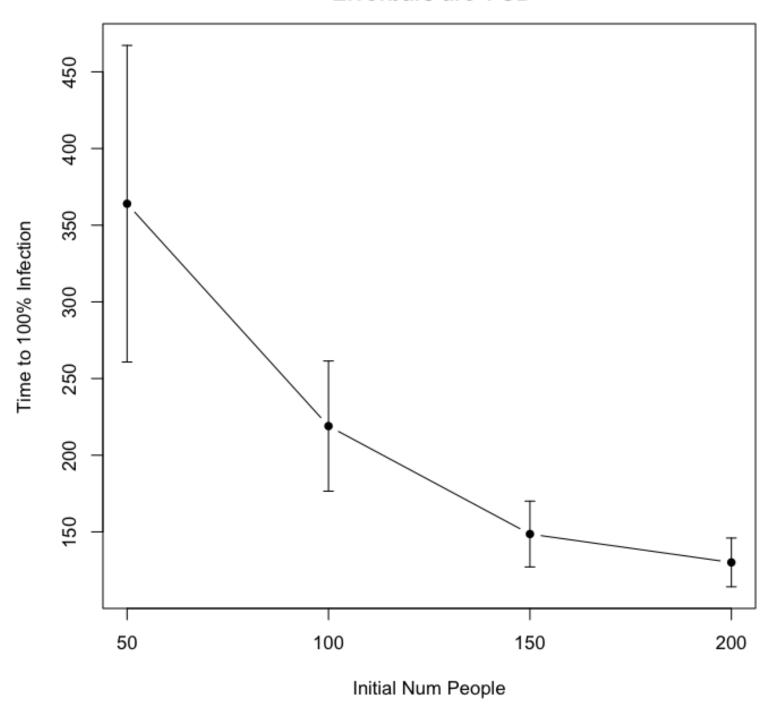
Creating Graphs

- Graphs provide a visual way to summarize the data
- They are not better or worse than summary statistics just different
- Many different graphs have been created
- Always useful to start by looking at all data
- Again, easy to do in R

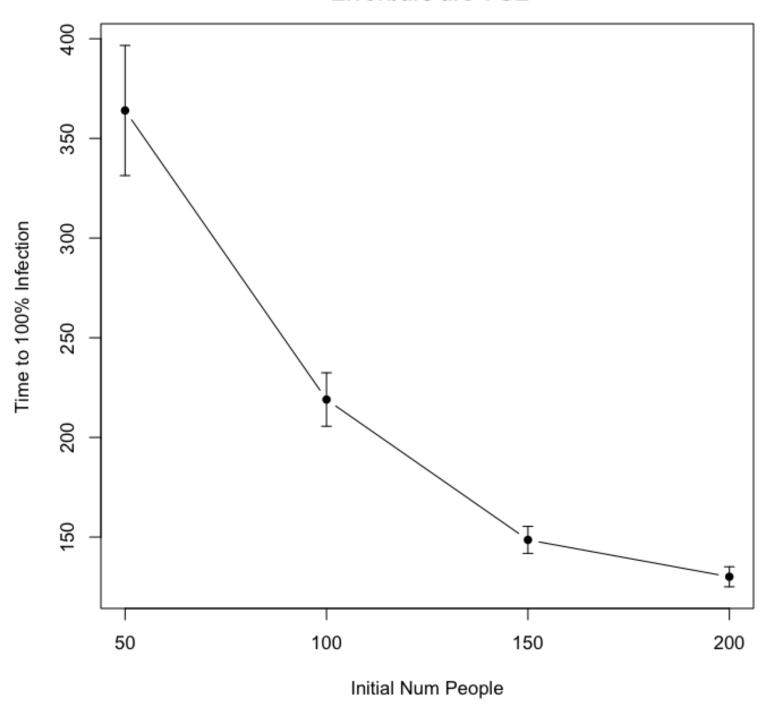
Time to 100% Infection vs. Population



Time to 100% Infection vs. Initial Population Errorbars are 1 SD



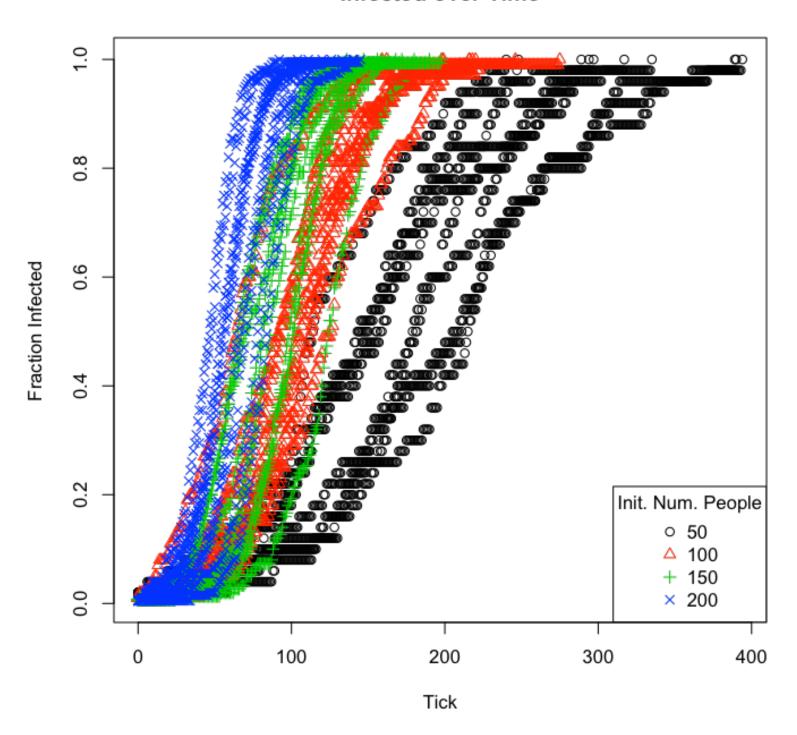
Time to 100% Infection vs. Initial Population Errorbars are 1 SE



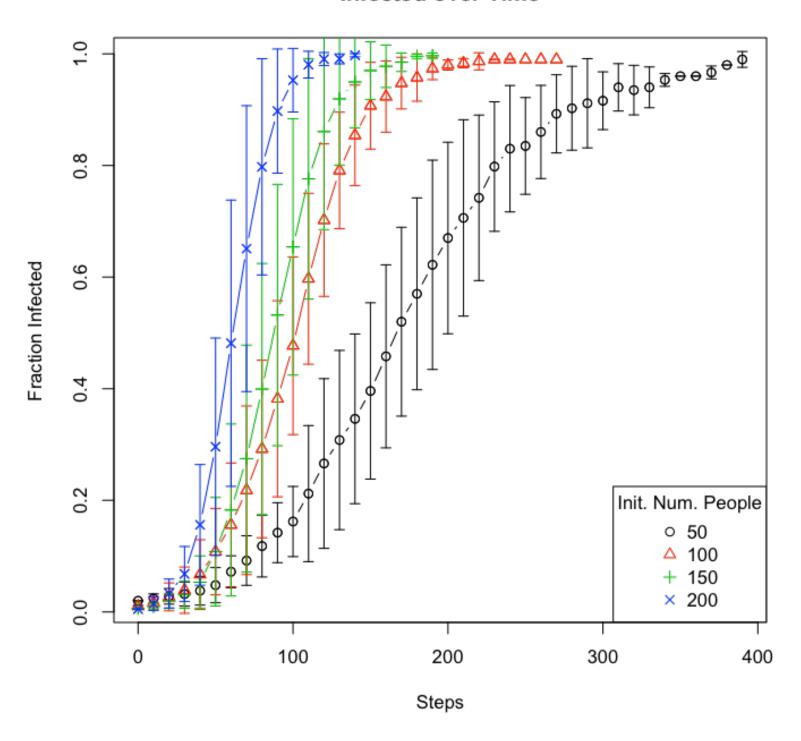
Time Series Graphs

- The graphs so far are all end of run results
- One important aspect of ABM is that you also have transitionary data recorded
- Runtime graphs are useful to look at, but they require lots of data

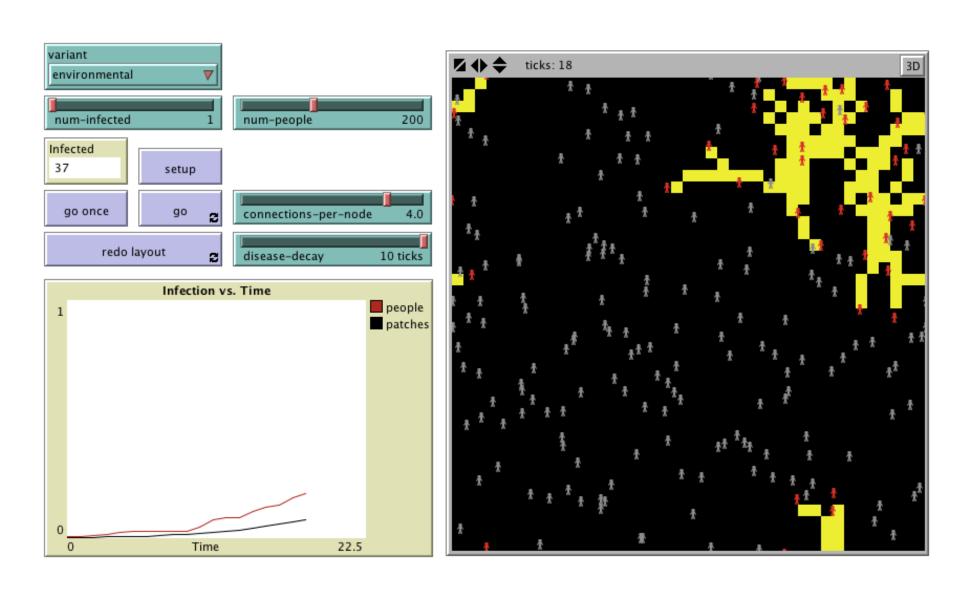
Infected over Time



Infected over Time

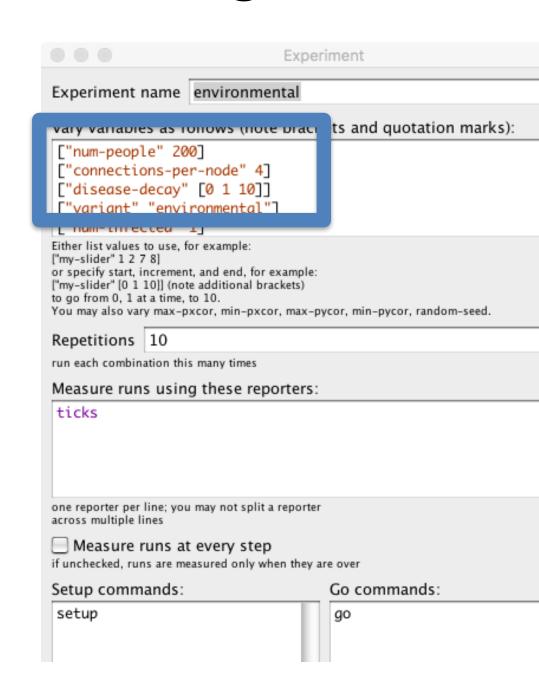


An Environmental Variant

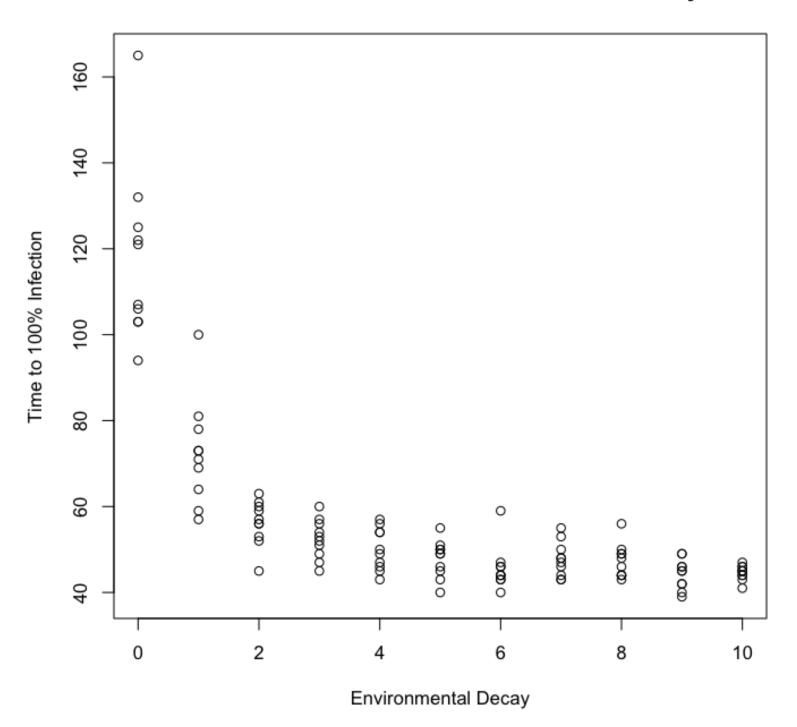


Environmental Changes

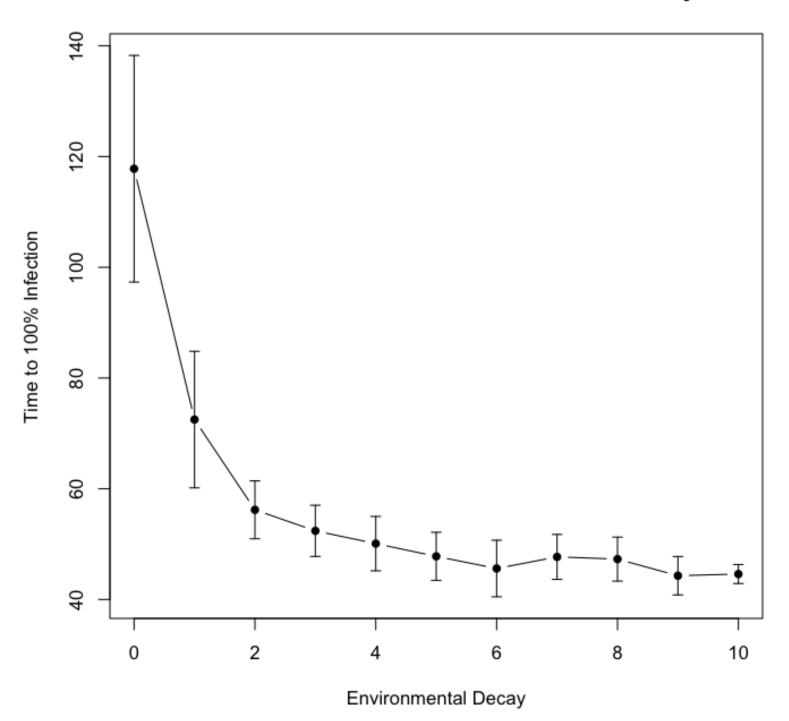
- Set number of people to a constant
- Add new variable called disease decay
- Anyone entering an area where the disease still is will be infected



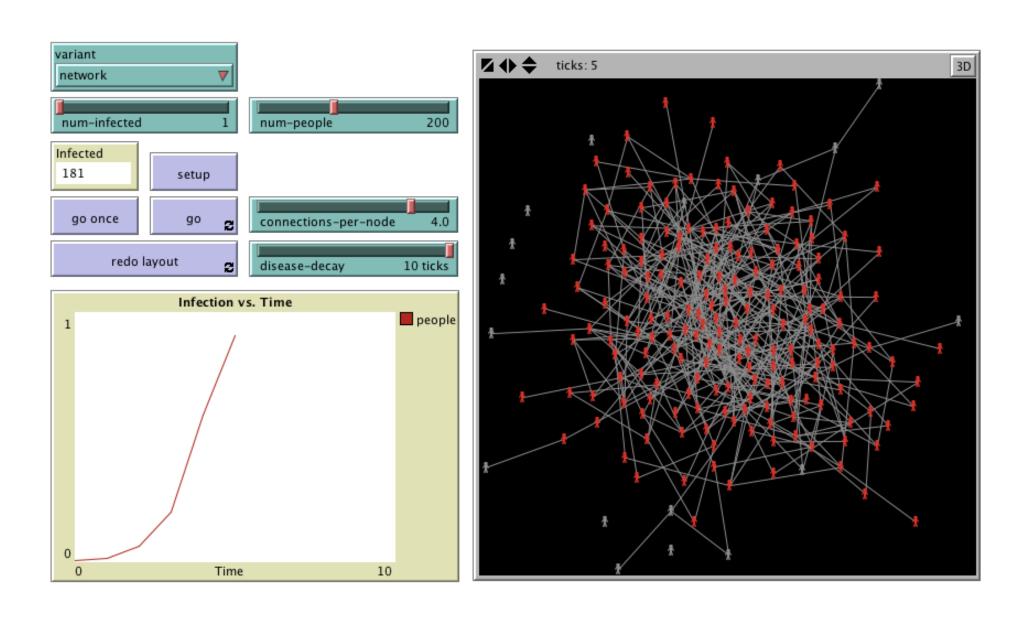
Time to 100% Infection vs. Environmental Decay



Time to 100% Infection vs. Environmental Decay

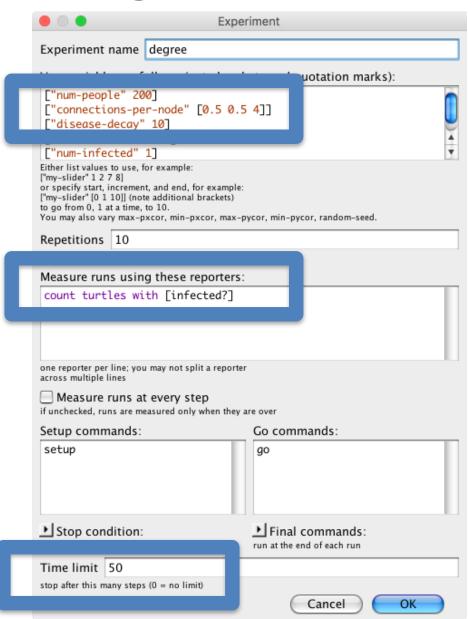


A Network Variant

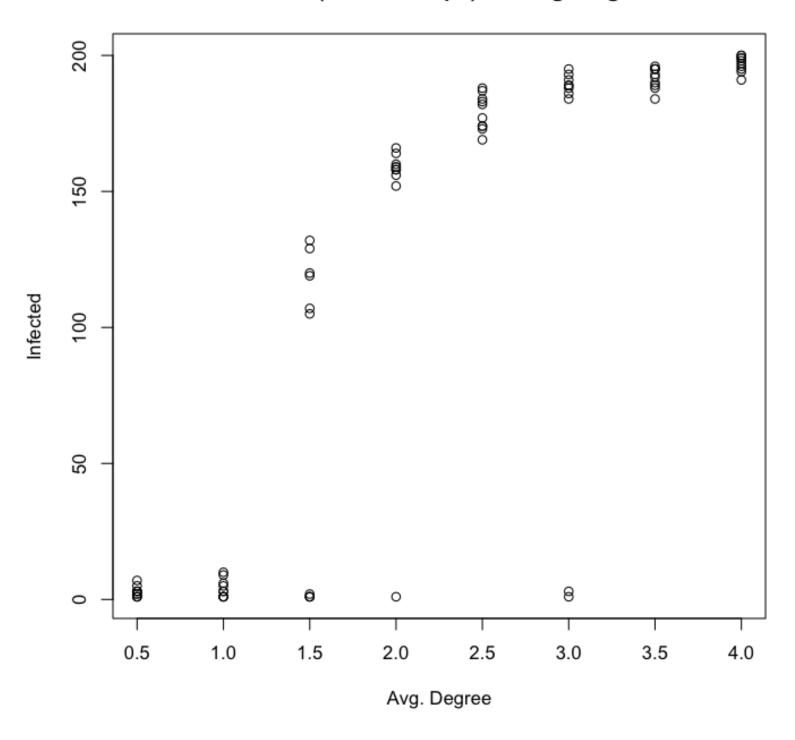


Network Changes

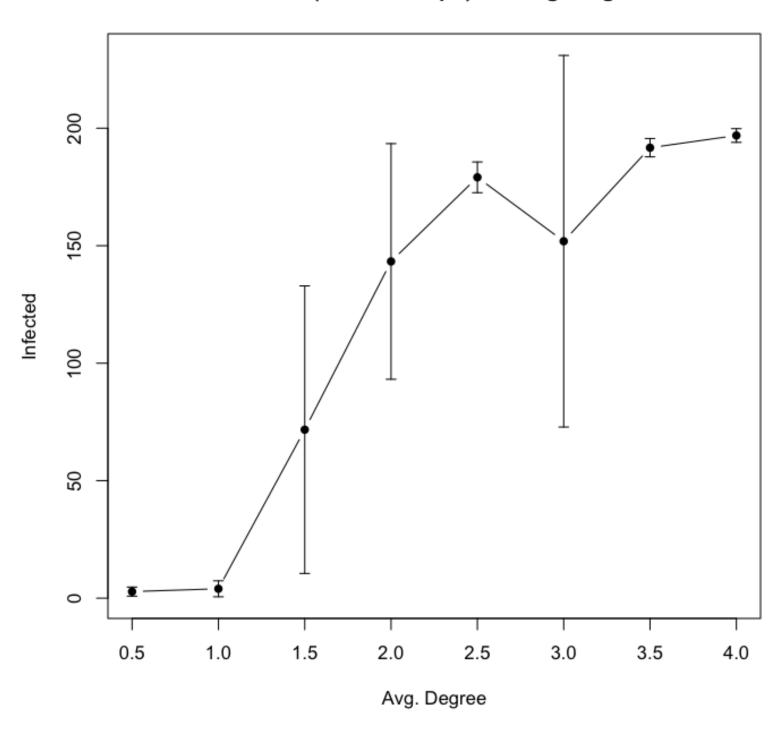
- 100% Infection is no longer guaranteed
- Measure spread until a certain time
- Record number infected at that time



Infected (after 50 steps) vs. Avg. Degree



Infected (after 50 steps) vs. Avg. Degree

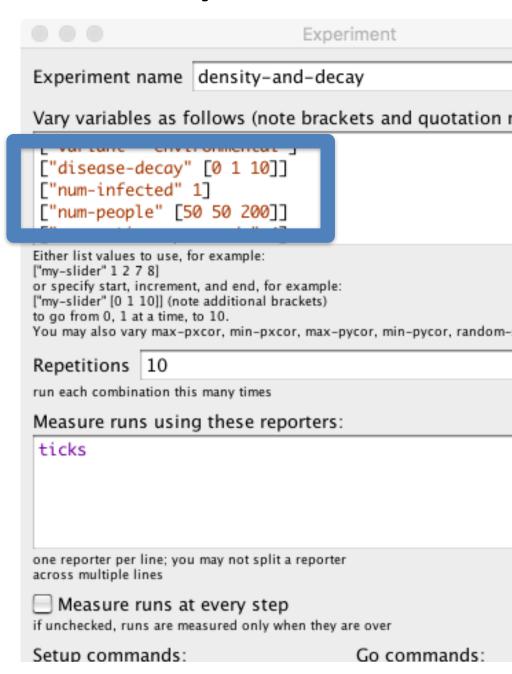


Multivariate Analysis

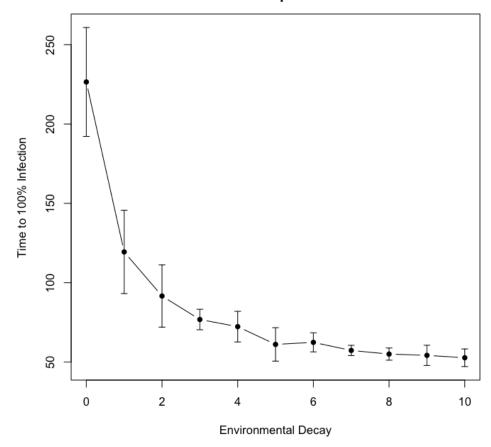
- Our analysis so far has been about one input and one output
- What if you wanted to look at multiple inputs simultaneously?
 - 1. Establish base settings and vary from that
 - 2. Aggregating over all other variables
 - 3. 3D Charts of combinations
 - 4. Regression

Density and Decay

- We will vary population density and disease decay simultaneously
- Observe the results
- 440 Runs

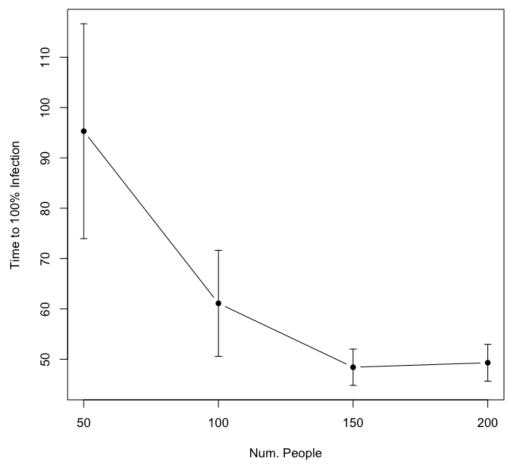


Time to 100% Infection vs. Environmental Decay Num. People = 100

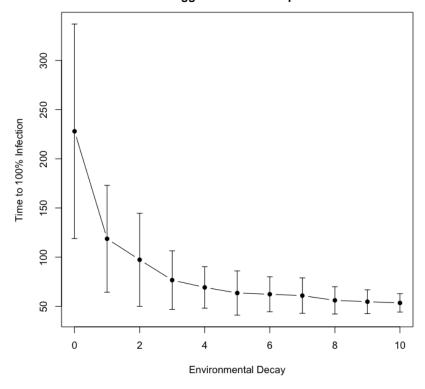


1. Base Case and Variants

Time to 100% Infection vs. Num. People Disease Decay = 5

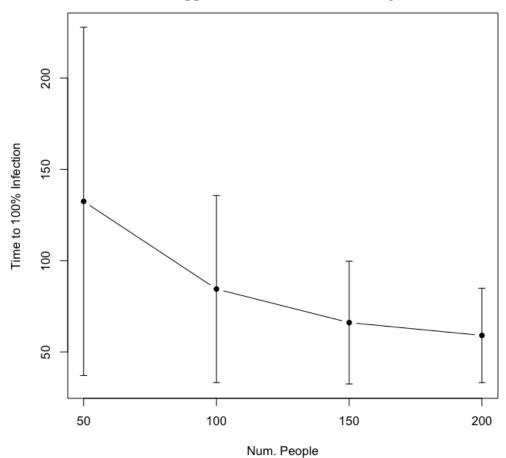


Time to 100% Infection vs. Environmental Decay Agg. over Num. People

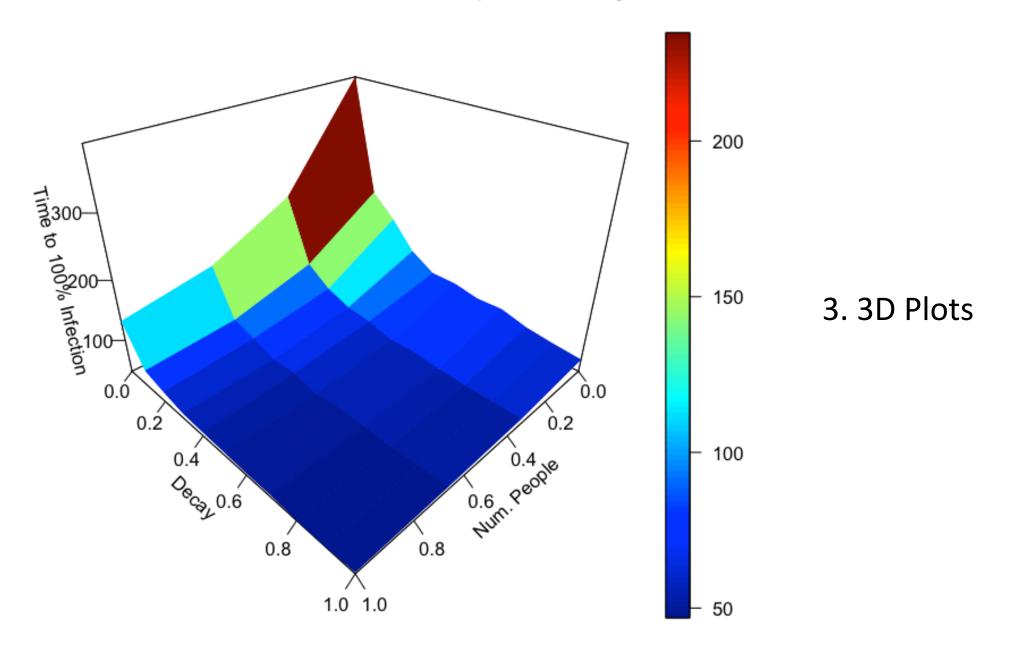


2. Aggregating over Variation

Time to 100% Infection vs. Num. People Agg. over Environmental Decay



Time to 100% Infection vs. Num. People and Decay Ticks



4. Regression

- If the model is really complex we can regress our output variable on all of our inputs
- This gives us a sense of how critical each input is to driving the output
- We can also look at the interactions

Coefficients:

```
Estimate Std. Error t value Pr(>|t|)

(Intercept) -3.428e-14 1.143e-14 -2.998e+00 0.00465 **

agg$num.people 1.000e+00 8.350e-17 1.198e+16 < 2e-16 ***

agg$disease.decay 2.643e-15 1.933e-15 1.367e+00 0.17912

agg$num.people:agg$disease.decay -1.586e-17 1.411e-17 -1.123e+00 0.26793

---

Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```