## **On the Infectious Diseases in Networks and TheirControl**

Lilia Leticia Ramírez-Ramírez and Mary E. Thompson**University of Waterloo**

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To obtain <sup>a</sup> more realistic epidemic model that generalize the most commonly used epidemic models that assume that

 susceptible individuals are equally likely to acquire the infection during an outbreak, and

the infectious process is markovian.

The last two assumptions the models are analyzed with well knownalgorithms as the Euler and Euler-Maruyama, from differential equationsand Markov process methodology.

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#### r. **Background**

- r. Infectious diseases in simple random networks Andersson (1998), Molloy and Reed (1995), Newman(2002)
- Hierarchical networks with two populations Ramirez-Ramirez and Thompson (a)
- Evolution of infectious process in discrete time
- Control Measures

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### **Law of mass action**

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#### **Network of contacts**

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Some infectious diseases are very much affected by the detailed structureof the contacts that can result in transmission.



### **Individual evolution**

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### **Networks**

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#### **Networks**









#### **Epidemic in Networks**





R code visualize with Social Network Image Animator (Sonia)

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### **Control Measures**

#### **Mass vaccination**

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A fraction  $\boldsymbol{\nu}$  of susceptible individuals is randomly selected and vaccinated prior to an outbreak.



#### **Acquaintance vaccination**

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First a fraction  $\boldsymbol{\omega}$  of susceptible nodes is randomly chosen. Then a neighbor of each of those nodes is randomly selected and vaccinated.



### **Ring vaccination**

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#### F. With probability  $\gamma$  any infectious node i is detected.

- F. With probability q each of its neighbors is identified for vaccination (and vaccinated).
- r. **This detection is done after m units of time since i is infectious.**

#### Infectious contact



### **Ring vaccination**

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- F. With probability  $\gamma$  any infectious node i is detected.
- F. With probability q each of its neighbors is identified for vaccination (and vaccinated).
- r. **This detection is done after m units of time since i is infectious.**



### **Quarantine-Isolation**

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- F. With probability  $\gamma$  any infectious node i is detected.
- F. Once detected  $i$  is "isolated", reducing its transmission rate by 1 $1 - \alpha \times 100\%.$
- r. **This detection is done after m units of time since i is infectious.**



# InfNet-Simulación de redes aleatorias y brotes epidémicos en ellas

Lilia Leticia Ramírez-Ramírez and Mary E. Thompson

March 30, <sup>2012</sup>

# ¿Como simular una red aleatoria?

# Algoritmos

■ Havel-Hakim

- Modified Havel-Hakim
- Molloy-Reed
- $\blacksquare$  Selección aleatoria

# InfNet

# Local.network

- n number of individuals (susceptible and infective)
- $\blacksquare$  distrib is the degree distribution
- $\blacksquare$  param is the distribution parameter (if the function is "fixed" it is a vector of degrees)
- distrib can be "fixed" or "pois" or "ztpois" or "geom" or "nbinom","ztgeom" or "poly.log" or "logarithmic"or "power.law" or "full" (fully connected) or "none" (no element connected)
- one.connection is TRUE when only one connection is allowed between two nodes.
- method specifies the algorithm to build the network: Havel-Hakim ("HH") or Modified Havel-Hakim ("MHH"), Molloy-Reed ("MR") or Random ("Random")

# Local.network

...

local.network<-function(n, distrib, param=NULL, one.connection= TRUE, method="MHH",degree=NULL)

list(edges=edges,degree=degree,degree.left=degree.left)

## connect.two.ln

■ Connects two populations (local networks) with the specified degrees using the "MHH" algorithm

 $\blacksquare$   $\blacksquare$   $\blacksquare$   $\blacksquare$  is the number of individuals in each local network

 $\blacksquare$  degree is in the format of list

connect.two.ln<-function(p,degree){

...

list(edges=edges,degree=degree,degree.left=degree.left)}

# epidemic.sim

- $\blacksquare$  network: network structure (output of global.network)
- ini.infected: number or initial cases in each network
- seir: it is FALSE if model is sir
- $\blacksquare$  ini.infective: number of initially infectives in each network
- obs.time: observation period
- BETA1: parameter of transmission to susceptible
- BETA2: parameter of transmission from an infective
- distrib.lat: distribution of the latent period. The default is poisson resulting in an overal Markov process
- LAMBDA: parameter for the latent period
- distrib.inf: distribution of the infective period. The default is poisson resulting in an overal Markov process

# epidemic.sim

- other options are Normal "norm" and lognormal "Inorm".
- GAMMA: parameter of distrib.inf. When distrib.inf is Poisson, it is the reciprocal of the removal rate of infectives. GAMMA is <sup>a</sup> scalar or vector of length <sup>n</sup> (Poisson), or <sup>a</sup> vector of length <sup>2</sup> (normal, lognormal  $list(c(1,2))$  ) or a matrix of nX2

# epidemic.sim

....

epidemic.sim<- function(network=NULL,seir=F,ini.infected=0, ini.infective=0,obs.time=0,BETA1,BETA2=1,distrib.lat="norm",LAMBDA=NULL,distrib.inf="exp",GAMMA,vac.par=NULL,eff.exp=0, $eff.inf=0)$ 

 list(indexes=indexes,time.hist=time.hist,nodes.hist=nodes.hist,edges.hist=edges.hist,p.hist=p.hist,susp.hist=susp.hist, infectedp.hist=infectedp.hist,infectivep.hist=infectivep.hist,remp.hist=remp.hist, n=n)}

# **Ejemplos**

tempnet<-global.network(1,p=500,distrib="pois",param=2) epid1<-epidemic.sim(tempnet,ini.infective=1,obs.time=10,BETA=.2,distrib.inf="lnorm",GAMMA=list(c(1,.2))) epid2<-epidemic.sim(tempnet,seir=T,ini.infected=1,obs.time=10,BETA=.2,LAMBDA=list(c(1,.1)),GAMMA=.1) epid3<-epidemic.sim(tempnet,seir=T,ini.infected=1,obs.time=10,BETA=.2,distrib.lat="lnorm",LAMBDA=list(c(1,.1)),GAMMA=.1)

```
tempnet2<-global.network(n=2,p=20,distrib="pois",param=c(2,3),distrib.among="pois",param.among=1)
epid21<-epidemic.sim(tempnet2,seir=T,ini.infected=c(2,3),obs.time=10,BETA=.2, distrib.lat="norm", LAMBDA=
  list(c(1,.1),c(2,.1)),GAMMA=.2)
epid22<-epidemic.sim(tempnet,ini.infective=1,obs.time=10,BETA1=.2,
BETA2=c(rep(1,10),rep(.1,10),rep(1000,10)),distrib.inf="lnorm",GAMMA=list(c(1,.2)))
```


## SIMID: *SIMulation of Infectious Disease*

*A tool for simulation of infectious disease, enabling spatio-temporal visualization of the dynamics of influenza outbreaks.* 

Eileen de Villa, Peel Public Health Matt McPherson, Infonaut Lilia Leticia Ramírez Ramírez, University of Waterloo

**The Ontario Public Health Convention (TOPHC)** April 7, 2011

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### **Overview**

- About SIMID
	- **Project Overview**
	- **How it Works**
- **Benefits & Challenges**
- What's Next

## **Project Overview**



## **What We Developed**

- A tool to enable **visualization of the dynamics of infectious disease outbreaks over time and space**. (starting with **influenza**, including pH1N1 and seasonal influenza)
- A tool that can **educate & inform** public health officials to **create a more effective local infectious disease outbreak planning process**
- A system design paradigm & technology platform **applicable to other health-related urgent responses**



## **Why We Developed It**

- To **provide public health officials and key decision makers** with more effective local infectious disease outbreak planning tools
- To **create a tool that can be modified** for other infectious disease outbreaks and health-related urgent response
- To **reduce the time required to prepare** for and **respond** to public health emergencies
- **Being better prepared for emergencies, including** infectious disease outbreaks, **mitigates their negative impact**



## **Our Vision**

"To create <sup>a</sup> more effective planning process for local infectious disease outbreaks and other disasters by allowing decision makers to visualize the dynamics of the outbreak over time within their own community"

### **How It Works**



### **Basic Workflow**





## **SIMID Under the Hood – How it Works (1)**

- 'Scenario Manager' parameterizes & tests a simulation in **DRAFT**
- Scenario Manager chooses to **RUN** a simulation
- A **Job Service** tells R software that a simulation is 'waiting' and executes the simulation in R
- When a simulation completes in R, data output is created

#### *SIMulation of Infectious Disease*



## **SIMID Under the Hood– How it Works (2)**

- Job service 'listens' to SIMID and determines whether the simulation has completed
- Once completed, Job Service takes a completed output generated by R and imports data into SIMID
- Simulation data is used by SIMID to generate map outputs & other simulation statistics.

*SIMulation of Infectious Disease*







The lifecycle of <sup>a</sup> simulation, from DRAFT, to RUNNING, to FINALIZED, to ARCHIVED status

#### **Simulation Lifecycle**











## **About the Model Used**

- Network Model developed in collaboration with University of Waterloo
- For details of the model see:

Ramírez Ramírez, L. L. (2008) "On the dynamics of infectious diseases in non-homogeneous populations" (http://uwspace.uwaterloo.ca/bitstream/10012/4054/1/llramireThesis.pdf)



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## **Overview**

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### **Benefits**



# **Benefits (1)**

### **Modeling for 'Non-Modelers'**

Complex disease models developed in 'R' language can be parameterized, run and visualized **by field personnel** who are designated 'Scenario Managers'

# **Collaborative Modeling**

Scenario Managers can explore simulations, change parameters and then **share outputs** with designated 'Participants'



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## **Potential End-Users ('Participants')**

- 0 **Public Health**
	- $\bullet$ **Epidemiologists**
	- $\bullet$ MOH Personnel
	- $\bullet$ Emergency Planners
	- $\qquad \qquad \bullet$ Infection Control Specialists
	- $\bullet$ Surveillance Managers
	- $\bullet$ Front Line Case Management Staff
- 0 **Institutions**
	- $\bullet$ Emergency Site Managers
	- $\bullet$ Infection Control (Acute care)
	- $\qquad \qquad \bullet$ Infection Control (LTC)
	- $\bullet$ Institutional Planners
	- $\bullet$  Emergency Room surge capacity **Managers**
	- $\qquad \qquad \bullet$ Hospital Administrators
	- $\bullet$ LTC Facility Administrators
- 0 **Municipalities**
	- $\bullet$ Emergency Services (911)
	- $\bullet$ Emergency Management
	- $\qquad \qquad \bullet$ Municipal Operations Centre
	- $\bullet$ Public Works
	- $\bullet$ Emergency Medical Services
	- $\bullet$ Municipal Planners
	- $\bullet$ RICN Coordinators



# **Benefits (2)**

# **Enhanced Table-Top Exercises**

Simulations can be developed to support 'table-top' exercises with high-fidelity scenarios

## **Cultivates a 'Learning Organization'**

Creates intra/inter-organizational knowledge by testing assumptions and exploring 'What if' scenarios, e.g.:

- •contrasting simulated interventions
- •• evaluating response times
- •• quantifying costs



# **Benefits (3)**

## **Expandable Platform**

Can be expanded to accommodate:

- Additional parameters for influenza, including interventions;
- Additional influenza models developed with R language
- $\bullet$  Models for additional disease types and others healthrelated emergency response

## **Challenges**



# **Challenges (1)**

## **"All Models Are Wrong, Some Are Useful"1**

Modeling by nature is an exercise in uncertainty… the model is not predictive

Challenge setting user expectations with respect to the limitations of the model vs. real world events

Our approach was to create a learning tool capable of creating a range of possible 'What if' scenarios for organizational learning

1. Attributed to George E.P. Box



# **Challenges (2)**

## **Complexity & Knowledge Translation**

Knowledge translation among a wide variety of highly specialized professionals: e.g. **mathematical modelers**, **developers**, **epidemiologists**, **public health practitioners**

## **Technological Changes**

Technological changes during the design phase dictated a change in some system components… impacting our workplan, resource requirements and the effort involved

*SIMulation of Infectious Disease*



# **Challenges (3)**

### **Pandemic H1N1 response**

Key project collaborators were occupied with the pandemic H1N1 response

Collaborators managing a real event find it hard to find time to plan for an imaginary event



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# **Next Steps (1)**

## **Further Testing**

Due to some of the challenges mentioned, our team was unable to complete thorough pilot testing

## **Consume Additional Influenza Models**

SIMID platform is designed for reuse including:

- System architecture
- 'Job service' that integrates GIS & 'R'
- System development methodology
- $\bullet$ Geo-processing service



# **Next Steps (2)**

## **Customize for Additional Disease Types**

Reuse capabilities offer a collaborative platform for public health emergency planning, e.g.:

- $\bullet$ Infectious Diseases
- Environmental Health
- CBRN (Chemical, Biological, Radiological & Nuclear)

**Opportunity for Expanding the Tool for Other Communicable Diseases / Urgent Response**



# **Next Steps (3)**

### **Identify Additional Partners, including:**

- **Hosting Partner –** *Need to 'Find a Home'*
- Additional Sources of Funding
- Additional End-Users / Emergency Planners at the local level

### **Opportunity for Deployment in Other Jurisdictions**



### **Acknowledgements**





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