Modeling the population dynamics of zoonoses, and the Allee effect

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Modeling zoonoses

Common characteristics:

- Spread between hosts (species, populations)
- Spread geographically,
- Mutate,

• ...

Two possible approaches: modeling the hosts vs. modeling the pathogen



Host-centered modeling

Example: compartmental models, like SIR

The base SIR model

$$\frac{dS}{dt} = -\beta IS \quad ; \quad \frac{dI}{dt} = \beta IS - \gamma I \quad ; \quad \frac{dR}{dt} = \gamma I$$

Image: A mathematical states and a mathem

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Host-centered modeling

Example: compartmental models, like SIR





Many extensions: V (vaccinated), D (deceased), C (carrier), E (exposed), ...

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Host-centered modeling

Example: compartmental models, like SIR





Many extensions: V (vaccinated), D (deceased), C (carrier), E (exposed), ...

- Directly models quantities relevant to public health
- Loses information about the pathogen (e.g. how it interacts with different hosts)

Host-centered models

System of equations for a pathogen-centered model



- $u_i(t, x, \theta)$: population density of the pathogen on host *i*,
- $\rho_i(t,x) = \int_{\Theta} u_i(t,x,\theta) d\theta$,
- $r_i(\theta)$: fitness of the pathogen on host *i*,
- $K_i(x)$: population density of host *i*.



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Allee effect



It may arise due to random fluctuations, which affect the survival of a species more when there are fewer individuals.

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System of equations with Allee effect

$$\partial_t u_i = d\Delta_x u_i + \mu \Delta_\theta u_i + r_i(\theta) u_i - \frac{u_i \rho_i}{\kappa_i(x)} + \alpha \kappa_i(x) \sum_{j \neq i} u_j + f(u_i) \quad (1)$$

 $f(u_i)$ is such that u_i decreases when it is below a threshold ε . When u_i is large, $f(u_i)$ is negligible.

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A reassuring result

There exists a unique solution to the system (1).

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Does adding the Allee effect make the model more useful?

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An argument why:

Will a given pathogen persist or go extinct?

Without the Allee effect: the answer depends only on the parameters and not on the initial sample i.e. if a pathogen can persist, it will always do so, not matter how small the starting population is.



Figure: A host population concentrated in two locations. In the model without the Allee effect, if the pathogen spreads in one location, it will also spread in the other.

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Useful for accurately testing the efficacy of public health measures.

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What's next?

- Do stochastic models recover the same consequences?
- · Combine the host-centered and pathogen-centered models,
- Develop a framework for simulation and parameter estimation,
- Test the models against real datasets.

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Thank you for your attention

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