

## Model evaluation and comparison

Jonathan Dushoff, McMaster University

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# Goals

- ▶ Discuss model types and model goals
- ▶ Explain the value of simulation for validating models
- ▶ Discuss metrics for evaluating fit
  - ▶ Put the Goodness of fit test in its place

# Do I have a good model?

- ▶ What is my model trying to accomplish?
  - ▶ Generating hypotheses
  - ▶ Evaluating plausibility
  - ▶ Prediction
  - ▶ Extrapolation
  - ▶ Mechanistic understanding

# Outline

Conceptual models

Prediction

Model Validation

Model Evaluation

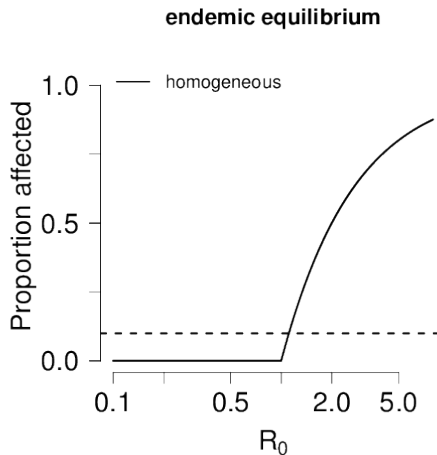
- Goodness of fit

- Capturing patterns

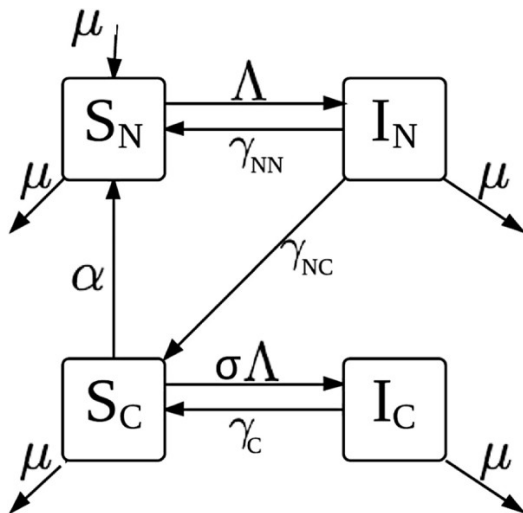
- Going beyond

Conclusion

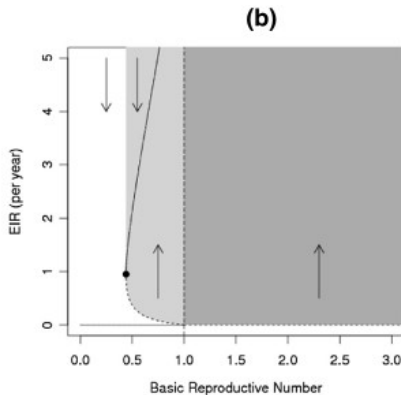
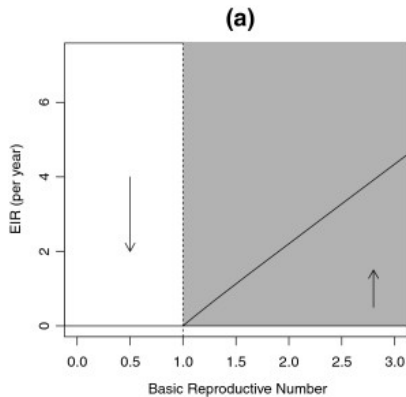
# Disease thresholds



## Effects of clinical immunity



# Bistability



# Outline

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- Goodness of fit

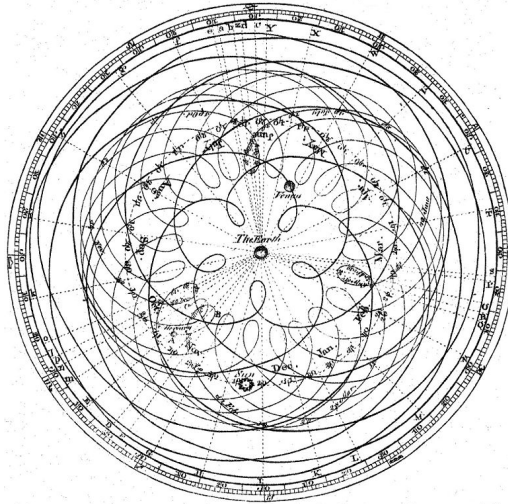
- Capturing patterns

- Going beyond

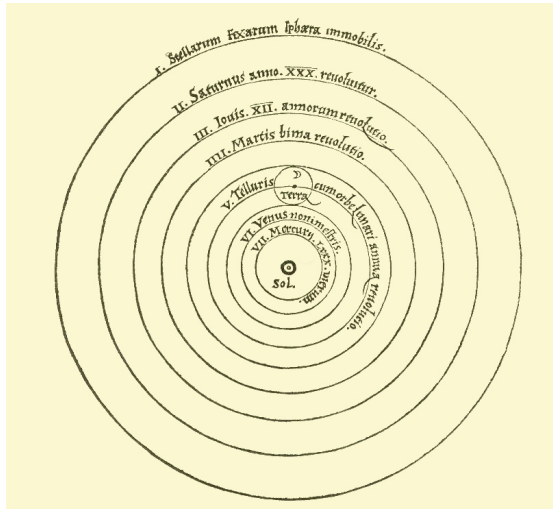
Conclusion



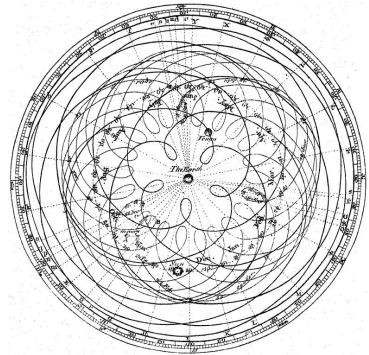
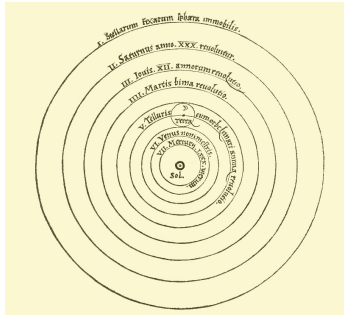
# Ptolemy v. Copernicus



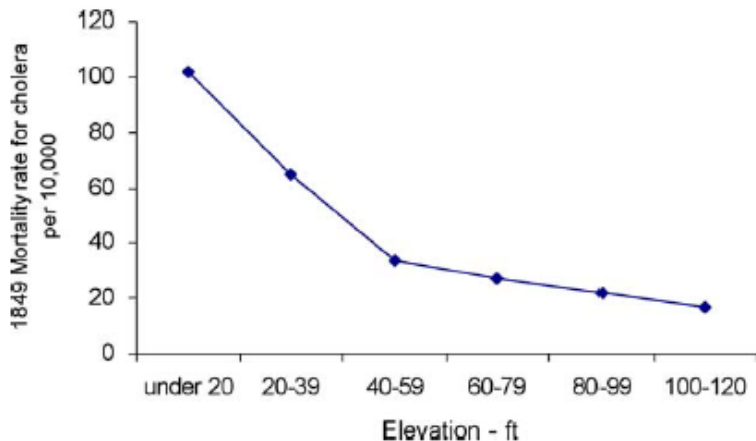
# Ptolemy v. Copernicus



# Ptolemy v. Copernicus



## Where will we see cholera cases?



# Where will we see cholera cases?



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Goodness of fit

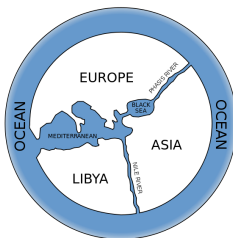
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# Model Validation

- Does your fitting algorithm match your *model world*?



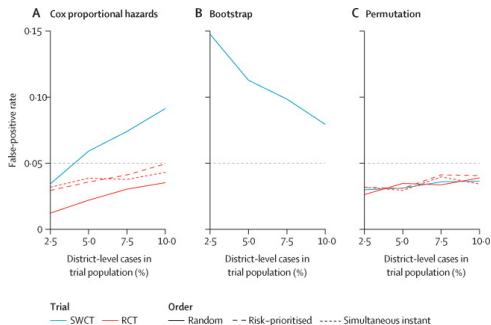
- If you use your fitting algorithm on simulations from your model world, then you *know the right answer!*

# Validation measures

- ▶ Coverage
- ▶ Precision
- ▶ Bias?
- ▶ Accuracy?



# Coverage



- ▶ The right answer should be inside your 95% confidence interval 95% of the time
  - ▶ If more, your model is *too conservative*
  - ▶ If less, your model is *invalid*
- ▶ In many cases it's good to look at the two tails separately:
  - ▶ How often do you overestimate? Underestimate?

# Precision

- ▶ You should aim to make your confidence intervals as narrow as possible
  - ▶ Provide as much information as possible
- ▶ As data increases, your precision should increase
  - ▶ CIs should approach zero width

# Bias?

- ▶ Nobody wants to be biased
- ▶ You *need* to be *asymptotically* unbiased
  - ▶ Good coverage and good precision assure this
- ▶ Not so clear you need to be *absolutely* unbiased
  - ▶ Bias is the difference between the *mean* expected prediction and the true value
  - ▶ Scale dependent: an unbiased estimate of  $\gamma$  is automatically a biased estimate of  $D$  (but not asymptotically biased)
- ▶ It may be better to evaluate using medians (instead of means)

# Accuracy?

- ▶ Nobody wants to be inaccurate
- ▶ Good coverage and good precision should guarantee good accuracy

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# Model Evaluation



- ▶ Does your model match the *real world*?
  - ▶ \* No!
- ▶ How well does your model match the real world?

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# Goodness of fit

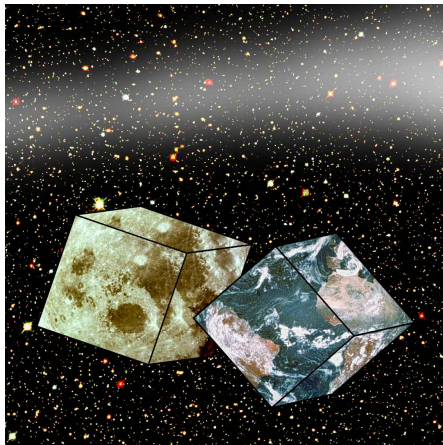
- ▶ Goodness of fit *statistics* describe how well a model prediction matches observed data
- ▶ Goodness of fit *tests* attempt to determine whether the observed difference between model and data is statistically significant



# Your model is false!

- ▶ A goodness of fit test won't make it true
- ▶ You can “pass” a goodness of fit test by:
  - ▶ having a good model
  - ▶ making very broad predictions
  - ▶ having bad data
  - ▶ choosing an inappropriate way to compare
- ▶ So why would we do this?
- ▶ For that matter, why do we use P values at all in biology?

# Passing goodness of fit tests



- ▶ I can make any model pass a goodness of fit test but broadening the uncertainty
- ▶ That doesn't make it a good model

# Low P values



## High P values



# What does the P value mean?

- ▶ Low: you are seeing something clearly
- ▶ High: you are seeing something unclearly

# Goodness of fit test

- ▶ Your model is *not* reality (null hypothesis is false)
- ▶ Can we see the difference clearly?
  - ▶ If no, model may be good or bad.
    - ▶ We probably can't add any more complexity based on current data
  - ▶ If yes, model may be good or bad. We *may* be able to add more complexity based on current data
    - ▶ But we may not need to

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# Capturing patterns

- ▶ You can ask:
  - ▶ Does your model do a reasonable job of capturing the data?
    - ▶ You can use a goodness of fit *statistic* for this, and not worry about the P value
  - ▶ Does your model capture patterns and relationships that you (or other experts) think are important?



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# Out-of-sample validation

- ▶ Does your model make predictions *outside* the range on which you calibrated it?
  - ▶ Predicting gravitational shifts in star positions from measurements in Earth laboratories
  - ▶ Predicting cholera outbreaks in Bangladesh from a model calibrated to Haiti
  - ▶ Predicting influenza patterns in 2010 from a model calibrated from 2000–2009

# Test sets

- ▶ What is **test set** spelled backwards?
- ▶ Hold some data out while fitting your model
- ▶ Or just *pretend* to do this as an evaluation method
  - ▶ In other words, test what would happen under various withholding scenarios

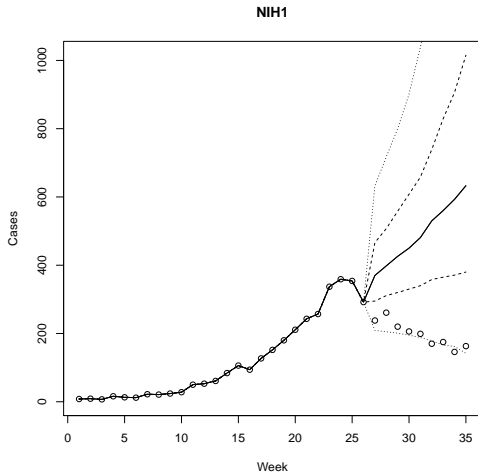
# Other model worlds

- ▶ The model you're *fitting* is probably pretty simple
- ▶ But you can *simulate* very complicated models, indeed

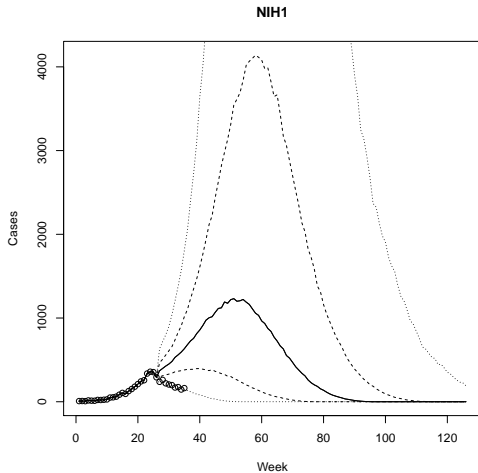


- ▶ How well can you do? Which details are important?

# Other model worlds



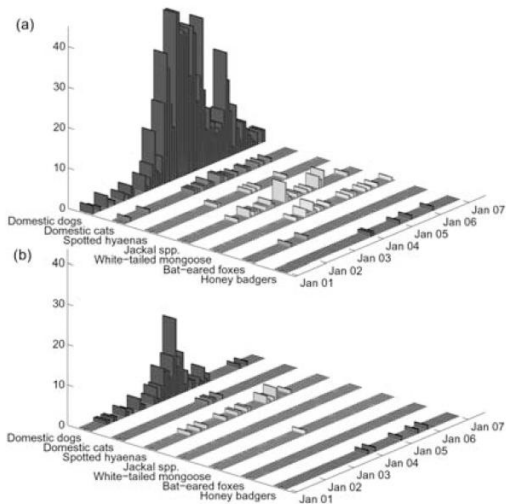
# Other model worlds



# Generating hypotheses

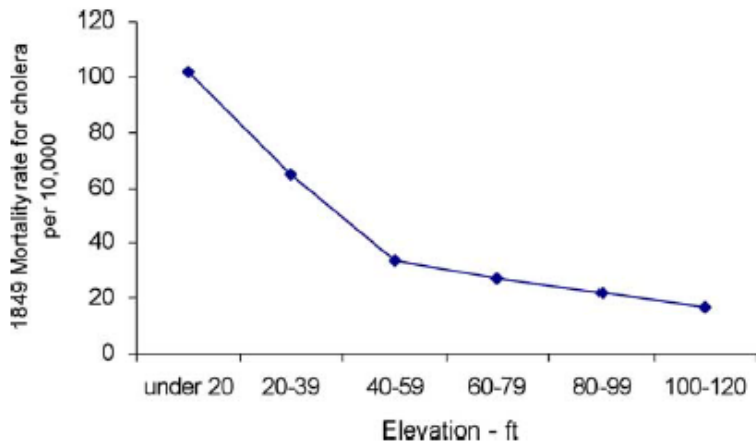


# Generating hypotheses





# Testing hypotheses



# Testing hypotheses



# Testing hypotheses



# Hard questions



Answers are not always easy

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- Capturing patterns

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# Summary

## Dynamic models

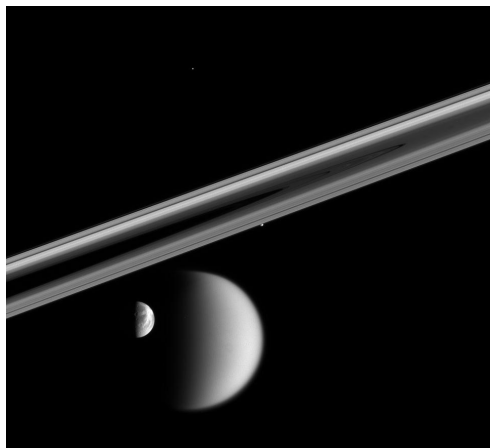
- ▶ Clarify thinking
  - ▶ What are our assumptions, what else do we need to know?
- ▶ Understand outcomes
  - ▶ Can heterogeneity explain the time course of HIV epidemics?
  - ▶ Is it possible that MDA could break the cycle of malaria transmission in some areas?
- ▶ Predict outcomes
  - ▶ What is the potential for a hepatitis A outbreak in Cape Town?
  - ▶ What might happen if I improve testing-and-treatment outreach in Jamaica?
- ▶ Find new mechanisms
  - ▶ Why can't I explain my data? What haven't I thought of?

# Summary

## Evaluation

- ▶ Validation (inside your model world)
  - ▶ Does my fitting method work (assuming my model is right)?
- ▶ Inspection (compare patterns)
- ▶ Prediction (and other out-of-sample comparison)
  - ▶ Can my model predict things I haven't told it yet?
- ▶ Generate and test mechanistic hypotheses

# Conclusion



Essentially, all models are wrong, but some are useful.  
– Box and Draper (1987), *Empirical Model Building* ...





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For further information please contact [admin@ici3d.org](mailto:admin@ici3d.org).



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