

# Inference

Connecting models to data

# The problem with infection data

Often only observe a proportion of reality

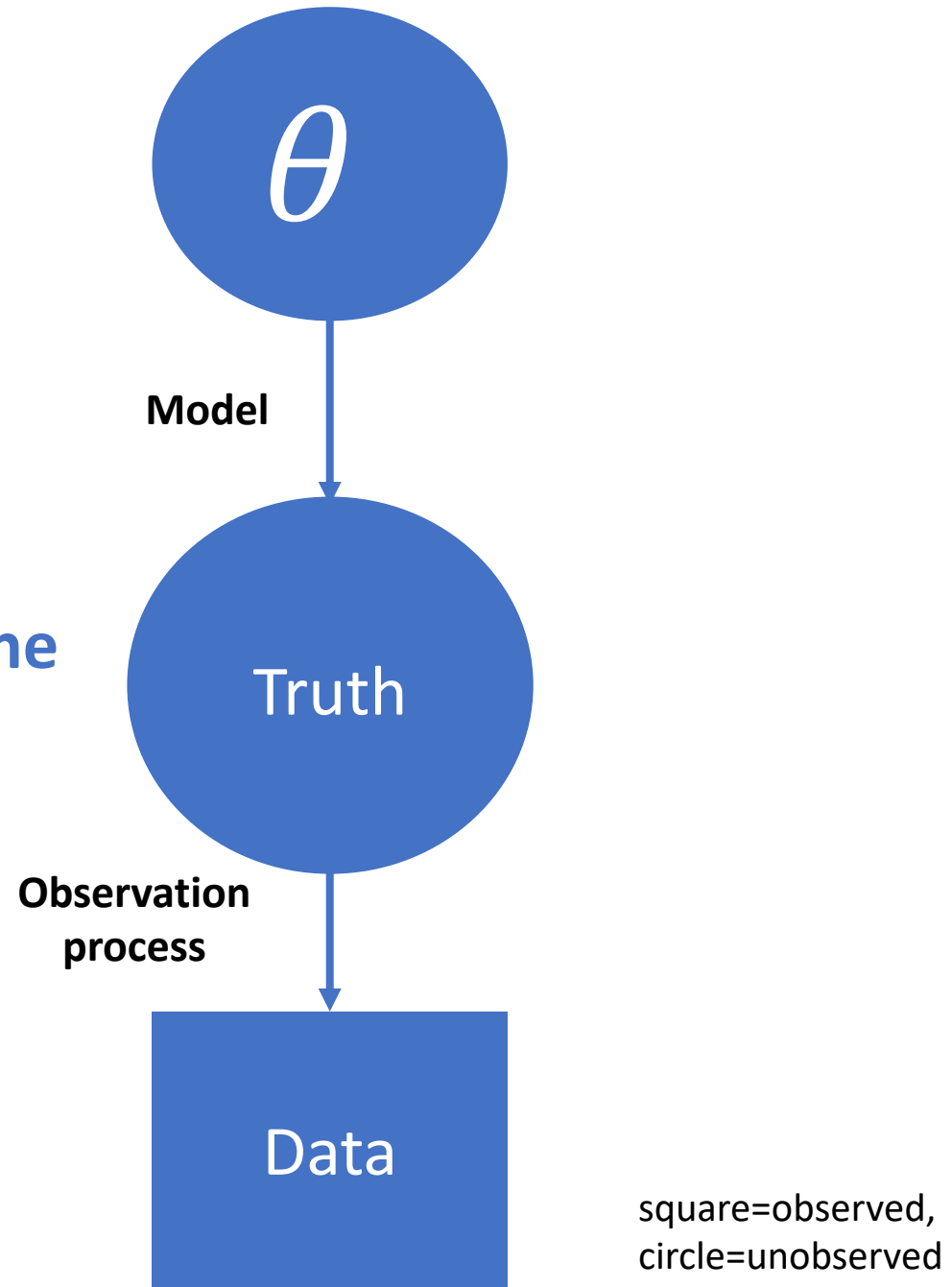
- Hospitalised case data gives you those who had severe infection
- Symptom onsets are observed but infection times are not

Or only observe a measure of infection

- antibody response at one time point
- result of imperfect diagnostic test

**We use this data to infer the ‘truth’.**

In a perfect world, we would directly observe the 'truth'.



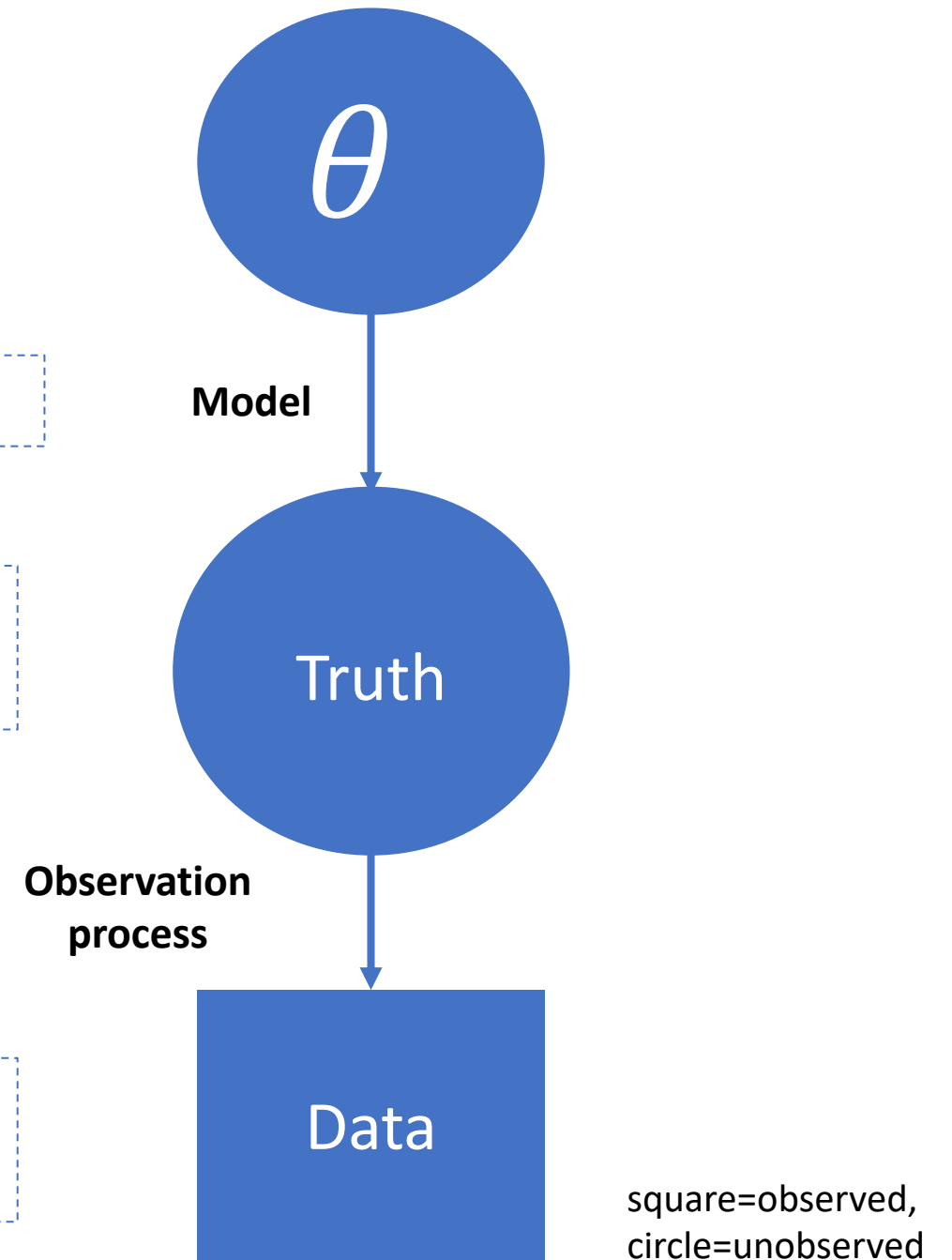
# Diagnostic testing results

- Susceptible-Infected model

- Predicted number of susceptible ( $S$ ) and infected ( $I$ ) animals

- $\text{Binomial}(I, \text{sensitivity}) \cdot \text{Binomial}(S, \text{specificity})$

- Diagnostic test results : test positives and test negatives



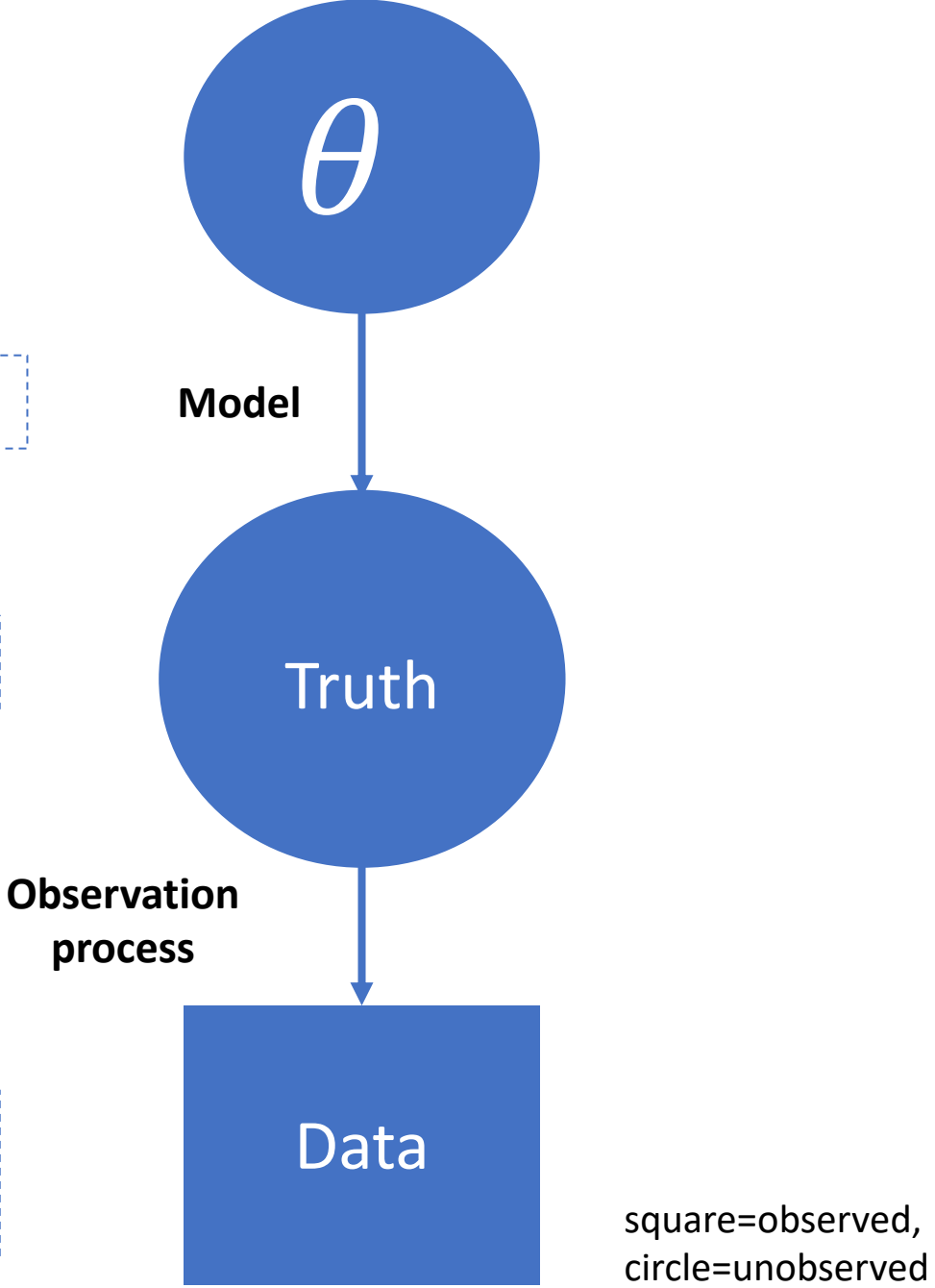
# Serological data

- Antibody process model

- Predicted log antibody titre

- Normally distributed error around predicted log antibody titre

- Laboratory based assay (measure of log antibody titre)



# Imperfect reporting of incidence data

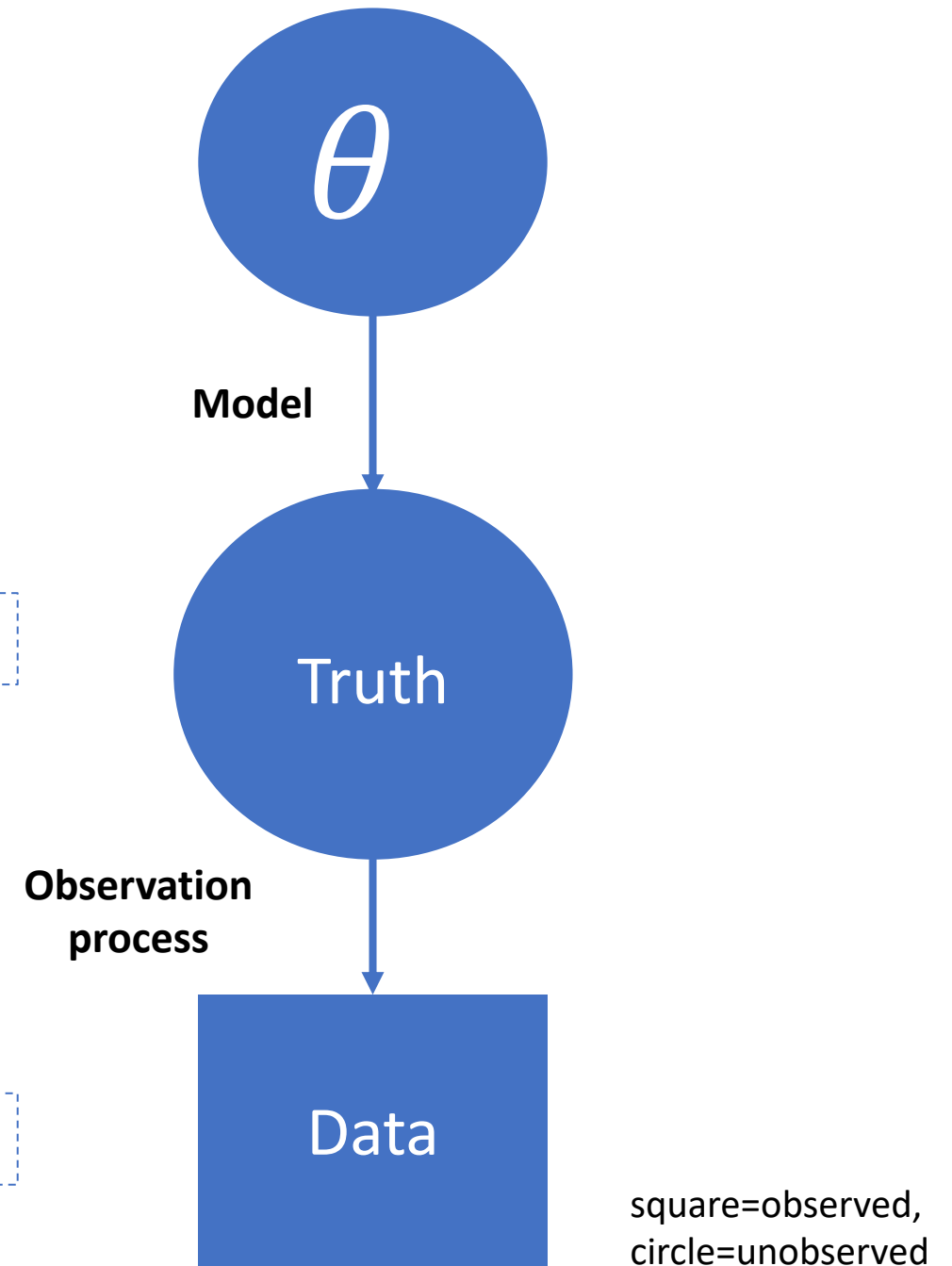
- $\theta = R_0, D_{lat}, D_{inf}, D_{imm}, \alpha, \rho$

- Deterministic/Stochastic SEITL model

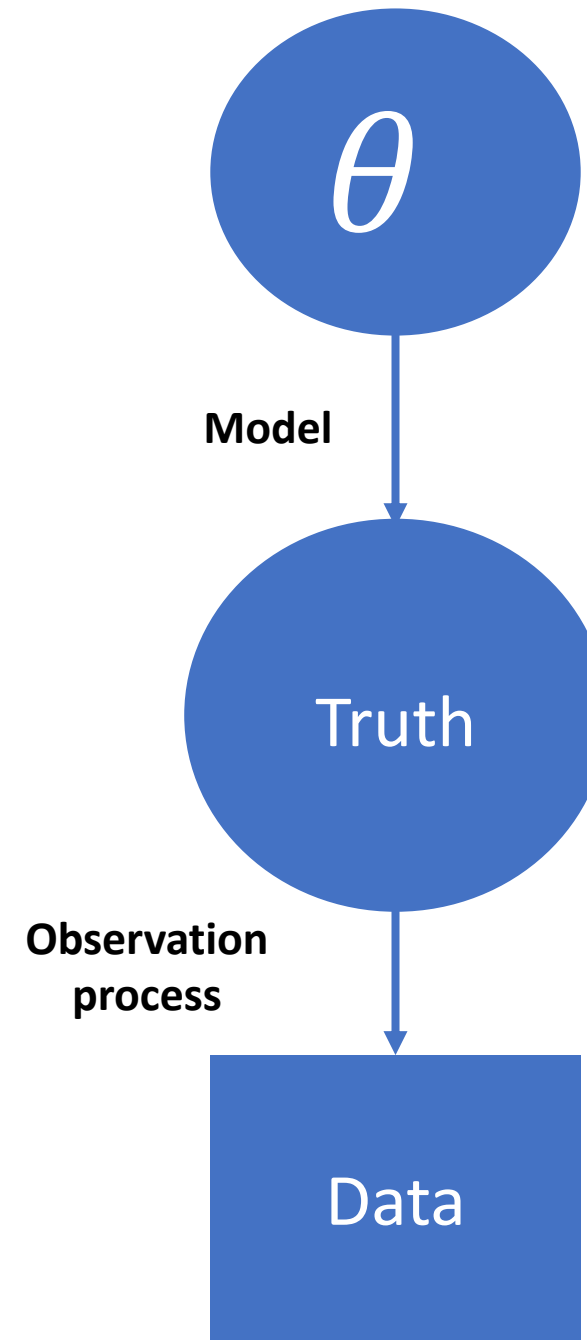
- Predicted incidence  $Inc$

- We assumed data were recorded according to a Poisson process :  $Poisson(\rho Inc)$  with reporting rate  $\rho$  and predicted incidence  $Inc$

- Reported incidence over time



Connecting your models to data relies on distinguishing how you predict the **'truth'** (*model*) and how you connect this 'truth' to your **data** (*observation process*).



# Examples

- Kucharski AJ, Lessler J, Cummings DAT, Riley S (2018) Timescales of influenza A/H3N2 antibody dynamics. PLOS Biology 16(8): e2004974. <https://doi.org/10.1371/journal.pbio.2004974>
- Brooks-Pollock E, Roberts G.O, Keeling, M.J (2014) A dynamic model of bovine tuberculosis spread and control in Great Britain. Nature, 511, pp. 228-231



# Approximate Bayesian Computation

# Outline

1. What is Approximate Bayesian Computation?
2. When do we use ABC instead of other methods?
3. How do we use it?
  - a) Choices in the ABC-rejection algorithm
  - b) Short introduction to more advanced ABC

1. What is Approximate Bayesian Computation?

Bayesian inference is based on the idea of updating belief with new evidence

- **Belief:** Prior distribution. Parameters are random variables instead of fixed quantities (they have their own distribution)
- **Evidence:** Likelihood function tells you the probability of the data given the parameters

# Bayesian inference

$\theta$  : Mathematical model parameter,  $D$  : Data

$$P(\theta|D) = \frac{P(D|\theta)P(\theta)}{P(D)}$$

# Bayesian inference

$\theta$  : Mathematical model parameter,  $D$  : Data

$$P(\theta|D) \propto P(D|\theta)P(\theta)$$

# Bayesian inference

$\theta$  : Mathematical model parameter,  $D$  : Data

$$P(\theta|D) \propto P(D|\theta)P(\theta)$$



**Probability of data  
given  $\theta$  (likelihood)**

**EVIDENCE**

# Bayesian inference

$\theta$  : Mathematical model parameter,  $D$  : Data

**Prior  
probability**  
**BELIEF**

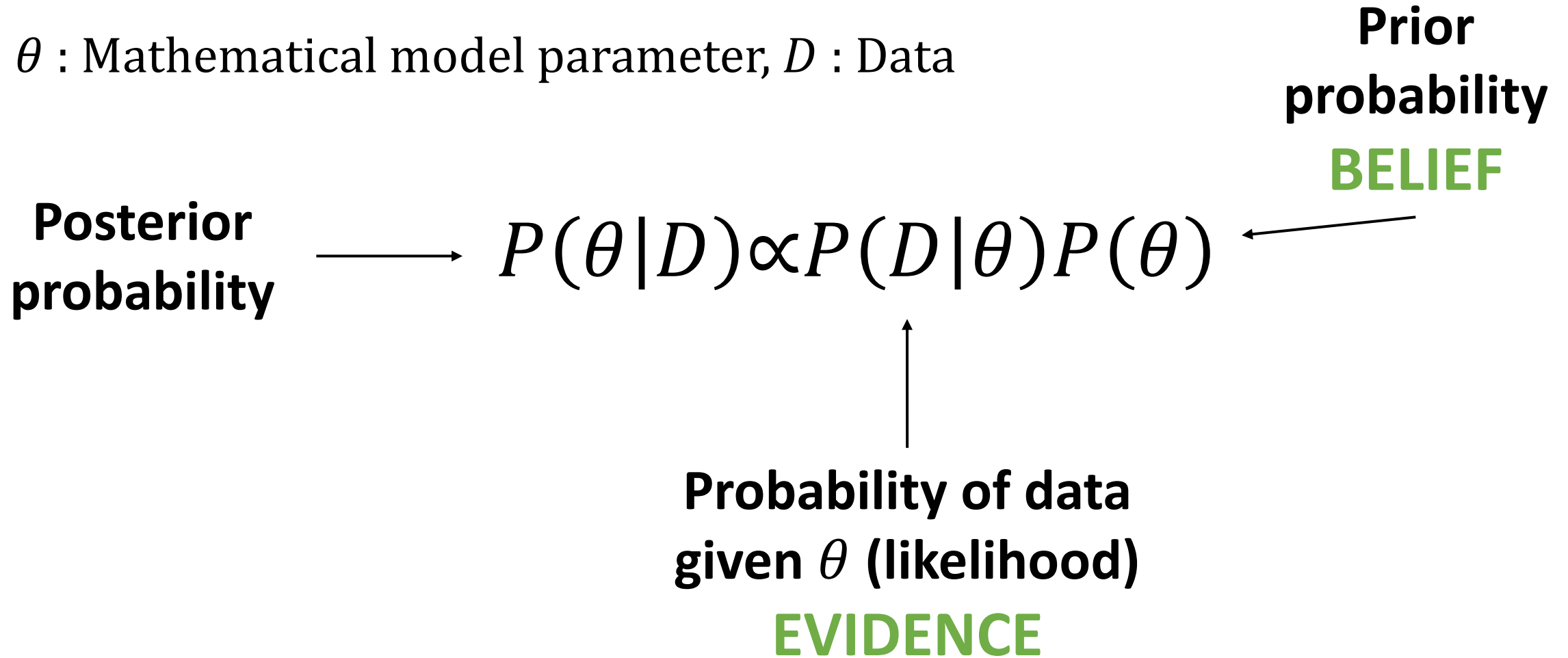
$$P(\theta|D) \propto P(D|\theta)P(\theta)$$

**Probability of data  
given  $\theta$  (likelihood)**  
**EVIDENCE**



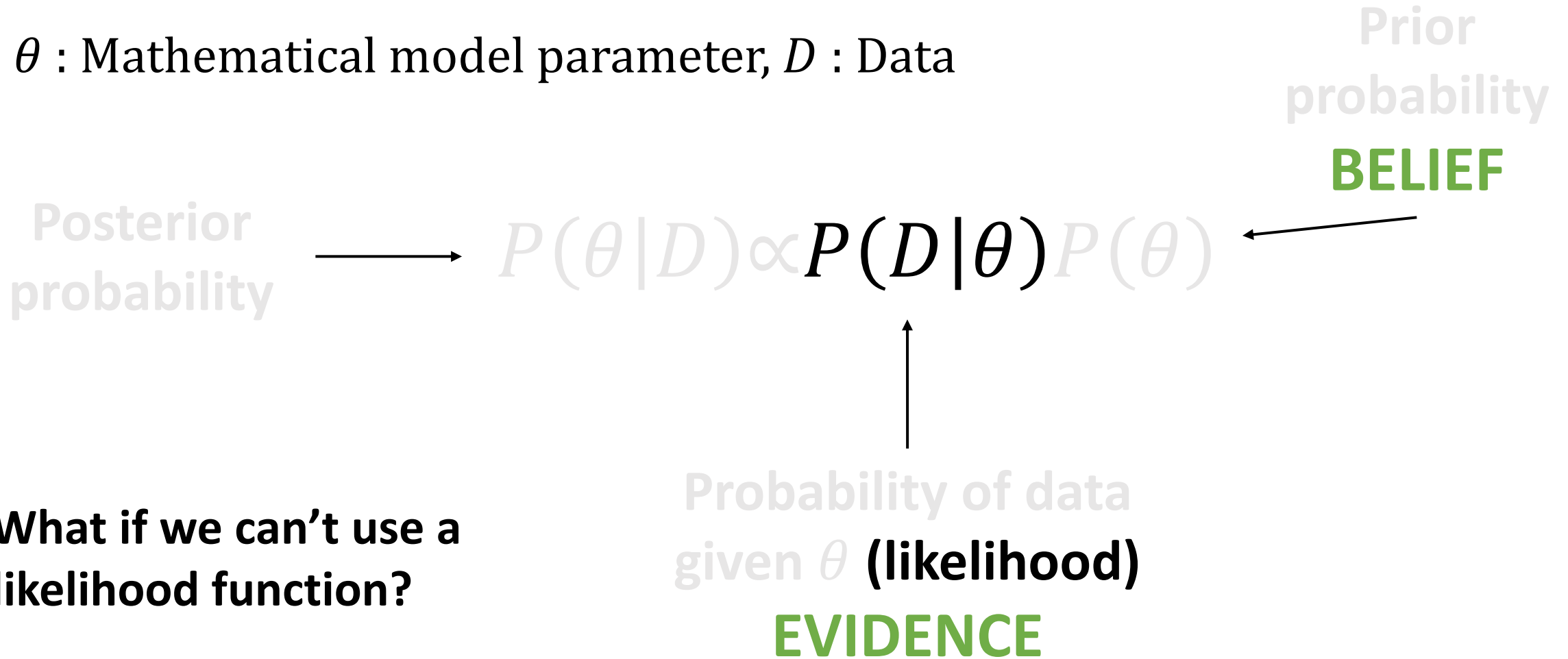
# Bayesian inference

$\theta$  : Mathematical model parameter,  $D$  : Data

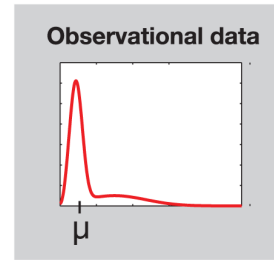


# Bayesian inference

$\theta$  : Mathematical model parameter,  $D$  : Data

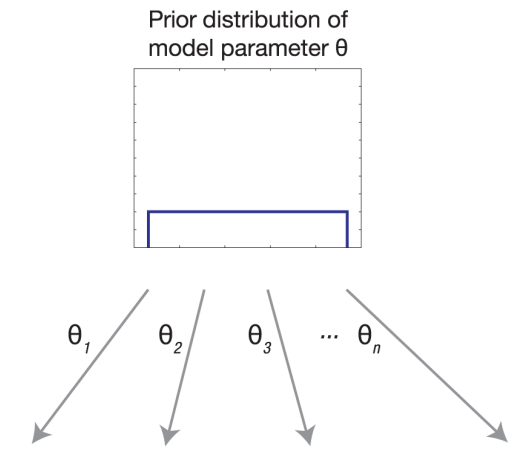
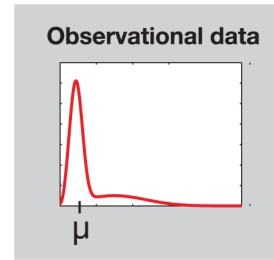


# ABC rejection algorithm



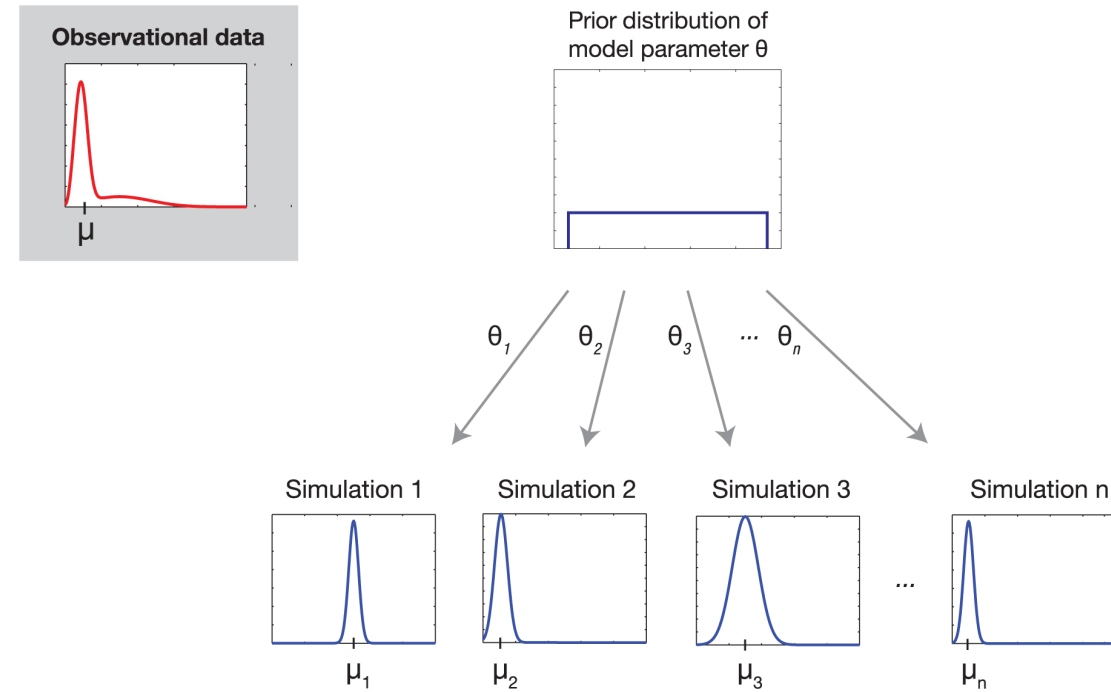
# ABC rejection algorithm

1. Sample  $\theta^*$  from the prior distribution  $P(\theta)$



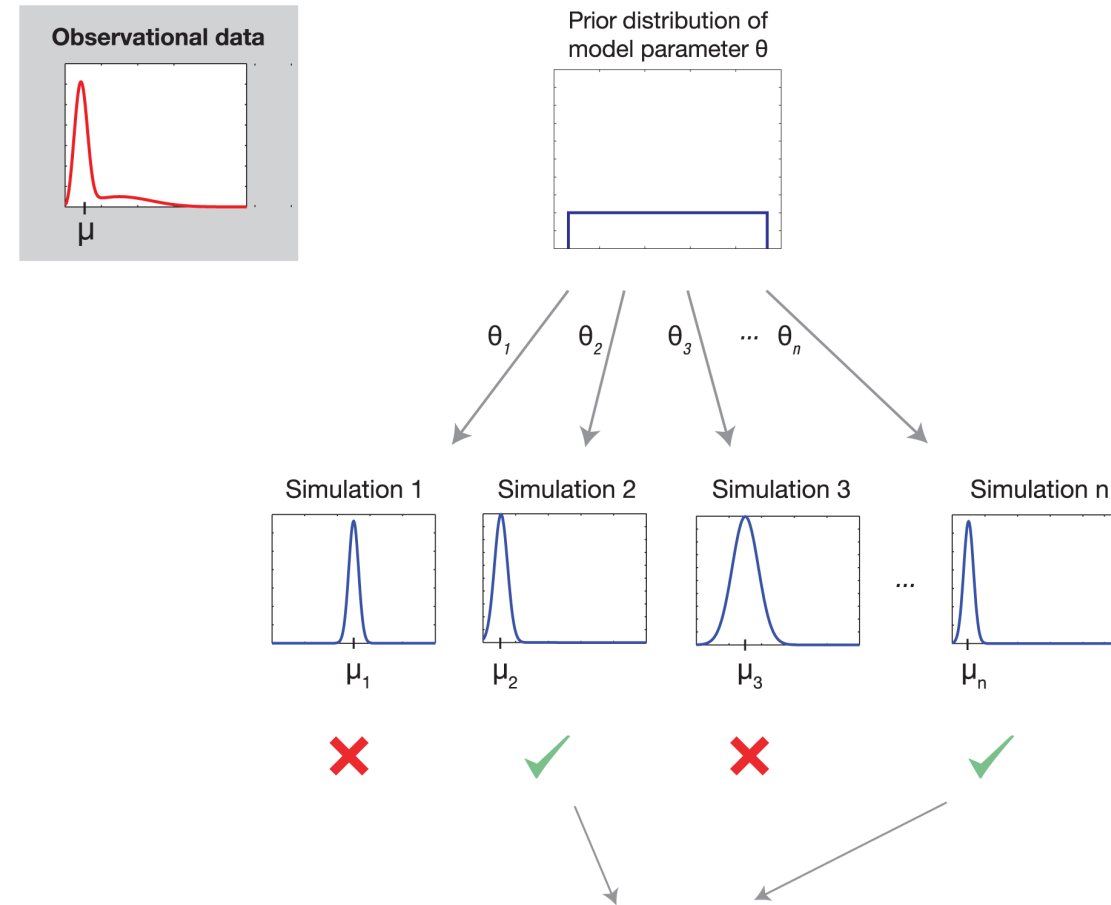
# ABC rejection algorithm

1. Sample  $\theta^*$  from the prior distribution  $P(\theta)$
2. Simulate a dataset  $D^*$  from your model using  $\theta^*$



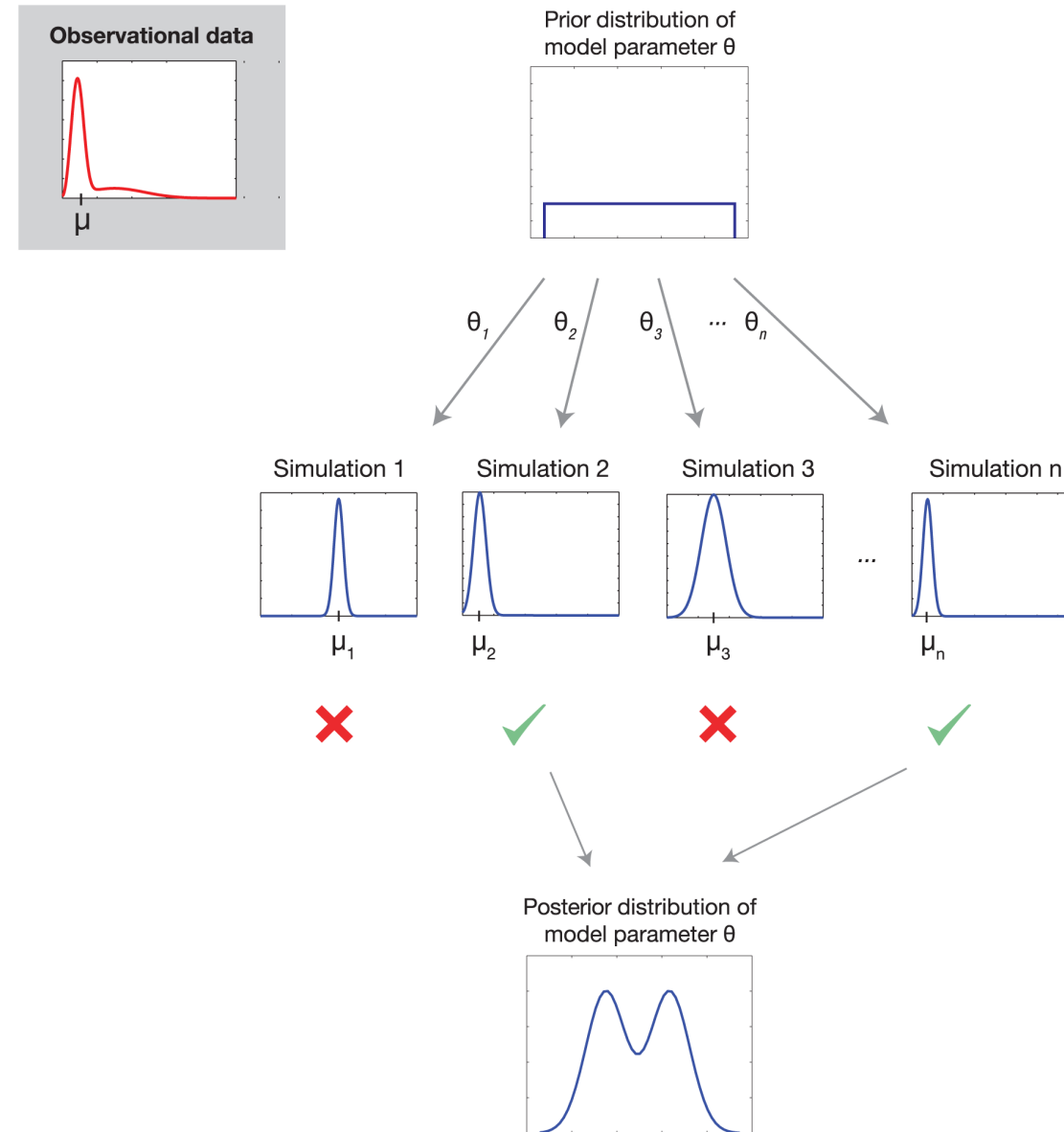
# ABC rejection algorithm

1. Sample  $\theta^*$  from the prior distribution  $P(\theta)$
2. Simulate a dataset  $D^*$  from your model using  $\theta^*$
3. If  $d(D, D^*) \leq \epsilon$  accept  $\theta^*$ , otherwise reject



# ABC rejection algorithm

1. Sample  $\theta^*$  from the prior distribution  $P(\theta)$
2. Simulate a dataset  $D^*$  from your model using  $\theta^*$
3. If  $d(D, D^*) \leq \epsilon$  accept  $\theta^*$ , otherwise reject
4. Repeat until you have  $N$  accepted samples



# ABC rejection algorithm

1. Sample  $\theta^*$  from  $P(\theta)$
2. Simulate a dataset  $D^*$  from your model using  $\theta^*$
3. **Calculate the summary statistic for the observed data  $\mu = S(D)$  and simulated data  $\mu = S(D^*)$**
4. **If  $d(S(D), S(D^*)) \leq \epsilon$  accept  $\theta^*$ , otherwise reject**
5. Repeat until you have  $N$  accepted samples



# ABC rejection algorithm

1. Sample  $\theta^*$  from  $P(\theta)$
2. Simulate a dataset  $D^*$  from your model using  $\theta^*$
3. **Calculate the summary statistic for the observed data  $\mu = S(D)$  and simulated data  $\mu = S(D^*)$**
4. **If  $d(S(D), S(D^*)) \leq \epsilon$  accept  $\theta^*$ , otherwise reject**
5. Repeat until you have  $N$  accepted samples

Summary statistic for model trajectory

Distance measure between summary statistic and data

# 1. What is Approximate Bayesian Computation?

A method to approximate the posterior distribution  $P(\theta|D)$  without a likelihood function

$$P(\theta|D) \approx P(\theta|d(S(D), S(D^*))) \leq \epsilon$$

2. When do we use ABC instead of other methods?

## 2. When do we use ABC instead of other methods?

- Data quality is poor, which means we have to aggregate it
  - Model: stochastic model of epidemic across a network of 6 small villages  
Data: one village had >50% attack rate; 4 villages had 10-50% attack rate; one village was not affected
- The likelihood function might be costly to evaluate (it takes a long time)
  - Large data sets / complicated likelihood function
- We want to reproduce patterns for which it is difficult to express a likelihood
  - Model: individual-based viral transmission model with explicit RNA sequence evolution  
Pattern: a particular binding motif (AAG CUG GGA U) appears within the virus

3. How do we use ABC?

a. Choices in the ABC- rejection algorithm

# Choice of summary statistic(s) $\mathbf{S}(\mathbf{D})$

- This is how we choose whether to accept or reject parameter values
- Sufficient summary statistic will give the same result as the likelihood
- "no other statistic that can be calculated from the same sample provides any additional information as to the value of the parameter"

# Choice of summary statistic(s) $\mathbf{S}(\mathbf{D})$

- This is how we choose whether to accept or reject parameter values
- Sufficient summary statistic will give the same result as the likelihood
- "no other statistic that can be calculated from the same sample provides any additional information as to the value of the parameter"
  
- If we haven't written down a likelihood then we can't know if our summary statistics are sufficient...

# Choice of summary statistic(s) $\mathbf{S}(\mathbf{D})$

- This is how we choose whether to accept or reject parameter values
- Sufficient summary statistic will give the same result as the likelihood
- "no other statistic that can be calculated from the same sample provides any additional information as to the value of the parameter"
- In practice
  - Look at published model fitting studies using ABC methods for ideas for sufficient statistics
  - **Check with simulated data!**



# Number of particles (N)

- The more the better, but computation time must be taken into account

# Tolerance value $\epsilon$

- Determines whether you accept or reject parameter(s) based on how closely the model prediction matches you data
  - Too small and the algorithm will take a long time to run
  - Too big and the final distribution of particles will be too wide

# Tolerance value $\epsilon$

- Determines whether you accept or reject parameter(s) based on how closely the model prediction matches you data
  - Too small and the algorithm will take a long time to run
  - Too big and the final distribution of particles will be too wide
- The magnitude of the tolerance value  $\epsilon$  will depend on your distance measure

For example, if the summary of the data  $S(D)$  is the cumulative number of cases, we could have:

- $S(D) = 100\,000$  (from the data)
- $S(D^*) = 99\,900$  (model prediction)
- If the distance measure  $d()$  is the sum of squared difference the,  
$$d(S(D), S(D^*)) = (100\,000 - 99\,900)^2 = (100)^2 = 10\,000$$

The prediction was 100 people short of the data, distance measure is 10 000. Hence here a reasonable choice of tolerance might be  $\epsilon = 10\,000$ .

3. How do we use ABC?

b. Short introduction to more advanced ABC

# Improvements to ABC rejection algorithm: ABC-Sequential Monte Carlo (ABC-SMC)

- Instead of one tolerance  $\epsilon$ , there is a vector of tolerances  $\epsilon_1, \dots, \epsilon_T$
1. We perform ABC rejection with a very large tolerance  $\epsilon_1$  and store our  $N$  accepted parameter values as population 1.

# Improvements to ABC rejection algorithm: ABC-Sequential Monte Carlo (ABC-SMC)

- Instead of one tolerance  $\epsilon$ , there is a vector of tolerances  $\epsilon_1, \dots, \epsilon_T$ 
  1. We perform ABC rejection with a very large tolerance  $\epsilon_1$  and store our  $N$  accepted parameter values as population 1.
  2. Then we propose parameters by re-sampling parameters from population 1 and perturb the parameters by a small amount. Accept/reject according  $\epsilon_2$ .
  3. Add **weight** to each parameter value according to the prior distribution, how likely you were to obtain that value from perturbation and the previous weights.

# Improvements to ABC rejection algorithm: ABC-Sequential Monte Carlo (ABC-SMC)

- Instead of one tolerance  $\epsilon$ , there is a vector of tolerances  $\epsilon_1, \dots, \epsilon_T$ 
  1. We perform ABC rejection with a very large tolerance  $\epsilon_1$  and store our  $N$  accepted parameter values as population 1.
  2. Then we propose parameters by re-sampling parameters from population 1 and perturb the parameters by a small amount. Accept/reject according  $\epsilon_2$ .
  3. Add **weight** to each parameter value according to the prior distribution, how likely you were to obtain that value from perturbation and the previous weights.
    - Repeat steps 2-3  $T$  times, sampling from the previous population. Each time decrease the tolerance value.



Practical

# In summary: ABC

- Can be used when data quality is poor, likelihood is complex or unknown and is an intuitive model fitting technique
- ***But*** you have to specify a suitable summary statistic(s)
- ABC can be slow, there are many extensions: ABC-SMC, ABC-PMC etc.

# Reading

## General introductions

- McKinley, Trevelyan J.; Vernon, Ian; Andrianakis, Ioannis; McCreesh, Nicky; Oakley, Jeremy E.; Nsubuga, Rebecca N.; Goldstein, Michael; White, Richard G. Approximate Bayesian Computation and Simulation-Based Inference for Complex Stochastic Epidemic Models. *Statist. Sci.* 33 (2018), no. 1, 4--18. doi:10.1214/17-STS618.  
<https://projecteuclid.org/euclid.ss/1517562021>
- Sunnåker M, Busetto AG, Numminen E, Corander J, Foll M, et al. (2013) Approximate Bayesian Computation. *PLOS Computational Biology* 9(1): e1002803. <https://doi.org/10.1371/journal.pcbi.1002803>
- Hartig, F. , Calabrese, J. M., Reineking, B. , Wiegand, T. and Huth, A. (2011), Statistical inference for stochastic simulation models – theory and application. *Ecology Letters*, 14: 816-827. doi:[10.1111/j.1461-0248.2011.01640.x](https://doi.org/10.1111/j.1461-0248.2011.01640.x)
- **Toni T, Welch D, Strelkowa N, Ipsen A, Stumpf MPH. (2009). Approximate Bayesian computation scheme for parameter inference and model selection in dynamical systems. *J. R. Soc. Interface* 6 187-202; DOI: 10.1098/rsif.2008.0172.**

# Reading

## Examples of ABC

- Conlan, A.J., McKinley, T.J., Karolemeas, K., Pollock, E.B., Goodchild, A.V., Mitchell, A.P., Birch, C.P., Clifton-Hadley, R.S. and Wood, J.L., (2012). Estimating the hidden burden of bovine tuberculosis in Great Britain. *PLoS Computational Biology*, 8(10), p.e1002730.
- McKinley, T., Cook, A. R. and Deardon, R. (2009). Inference in epidemic models without likelihoods. *Int. J. Biostat.* 5.
- Beaumont MA, Zhang W, and Balding DJ. (2002) Approximate Bayesian Computation in Population Genetics. *GENETICS*. 162 (4) 2025-2035.