



# Epidemic renewal models and sequential Monte Carlo

Nicholas Steyn - 30 May 2025  
Bioinference

# Who is this talk for?

Public health  
epi

An intro/refresher on  
**SMC methods**

An intro/refresher on the  
renewal model

A (small) handful of real-  
world results

(Statistical)  
modeller

Statistical  
epidemiologist

## Renewal Models

A simple ***model*** of an epidemic

$$C_t \sim \text{Poisson} \left( R_t \sum_{u=1}^{\omega_{max}} C_{t-u} \omega_u \right)$$

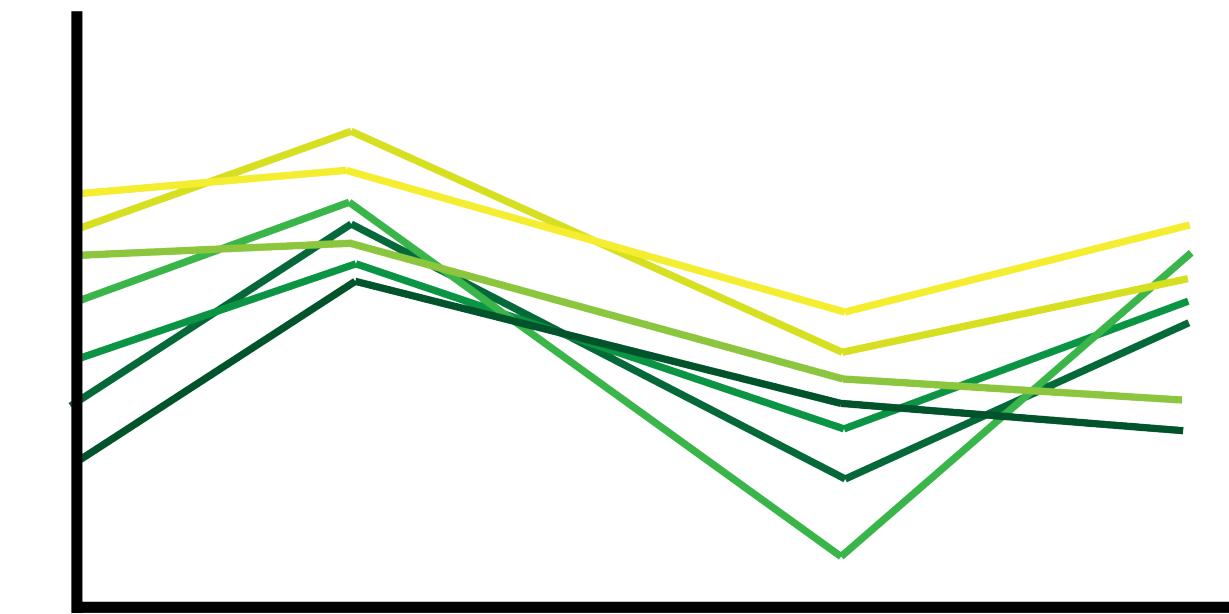
Used for reproduction number estimation

Also forecasting, elimination, effect of NPIs, etc

EpiEstim/EpiNow2/EpiFilter all use it

## Sequential Monte Carlo

A ***method*** for fitting hidden-state models



Can handle any ***sequential*** model

Also known as “particle filters”

Very flexible!

# Renewal Models

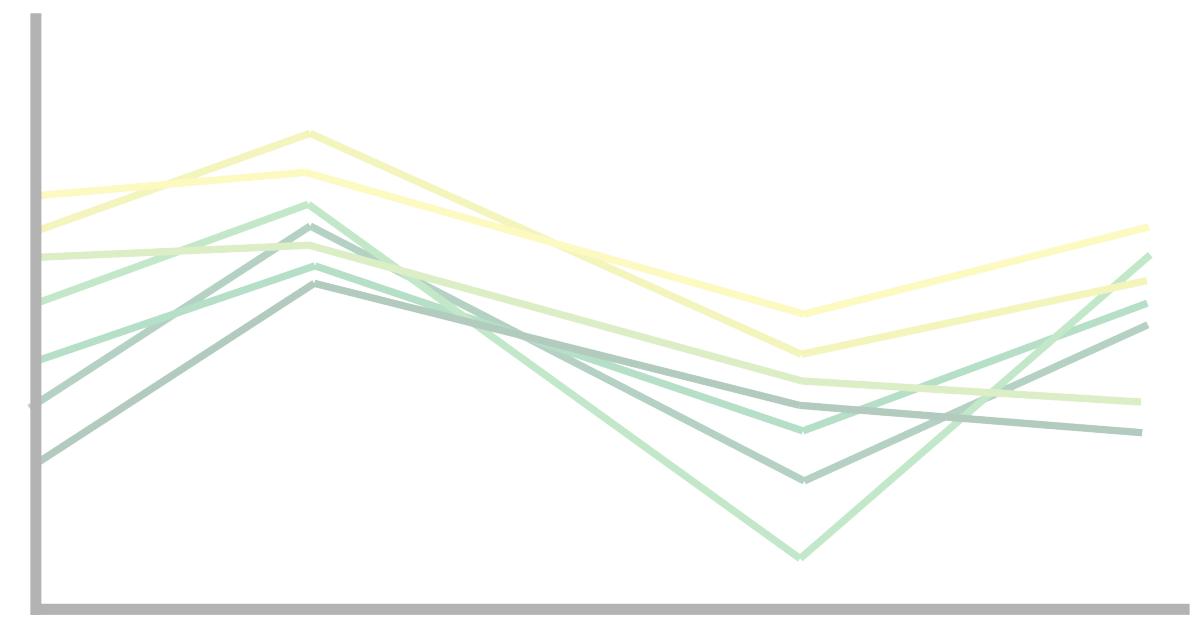
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# The renewal model

- Originally attributed to Euler in 1767, modelling population dynamics
- Lotka gave us the current (continuous time form) in 1907:

$$B(t) = \int_0^t B(t-a)p(a)v(a) da$$

births      P(survival)      fertility

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births      P(survival)      fertility

infections      P(infectious)      infectiousness

- Adapted to infections by Kermack and McKendrick (1927)\*

# The renewal model

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- Adapted to allow for time-varying average infectiousness, then
- (Fraser, 2007) separated infectiousness into the reproduction number and the generation time distribution\*:

$$I(t) = R(t) \int_0^t I(t - \tau) \omega(\tau) d\tau$$

# The renewal model

- A more familiar form was introduced in (Diekmann, 1977)
- Adapted to allow for time-varying average infectiousness, then
- (Fraser, 2007) separated infectiousness into the reproduction number and the generation time distribution
- and introduced the **discrete-time analogue**:

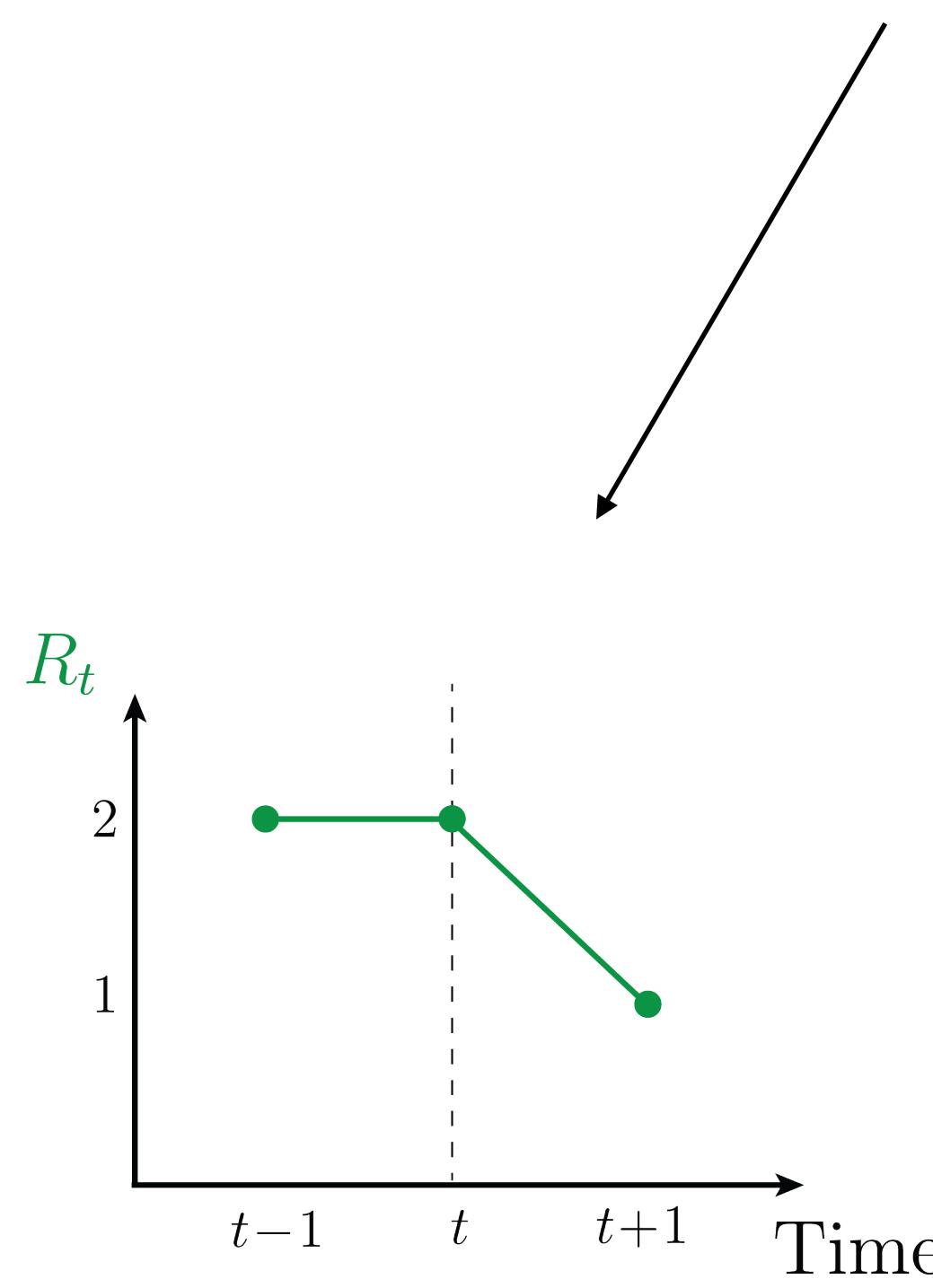
$$I_t = R_t \sum_{\tau=0}^t I_{t-\tau} \omega_\tau$$

- Which was then popularised by the EpiEstim software package (Cori et al, 2013)

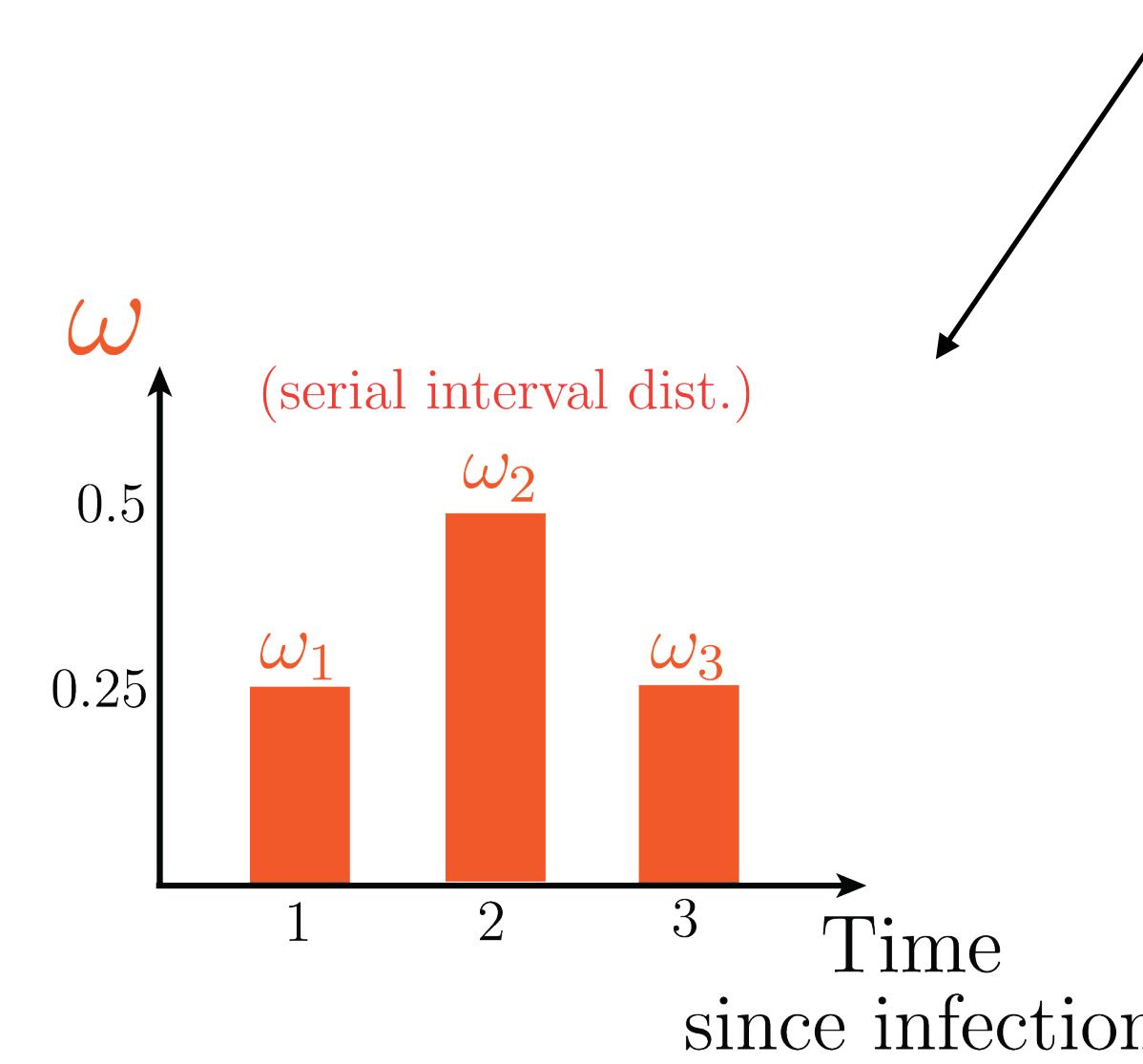
# The renewal model

## A toy example

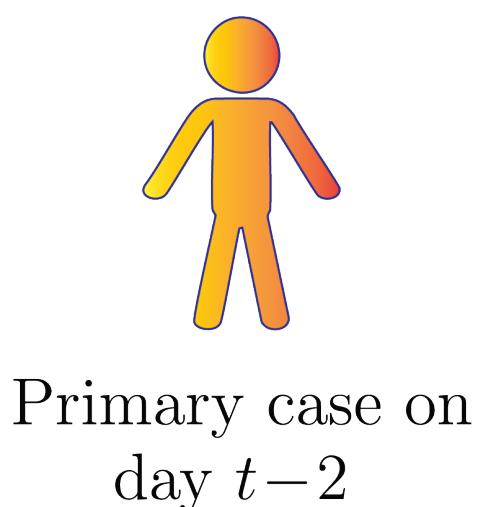
Reproduction number over time



A serial interval distribution

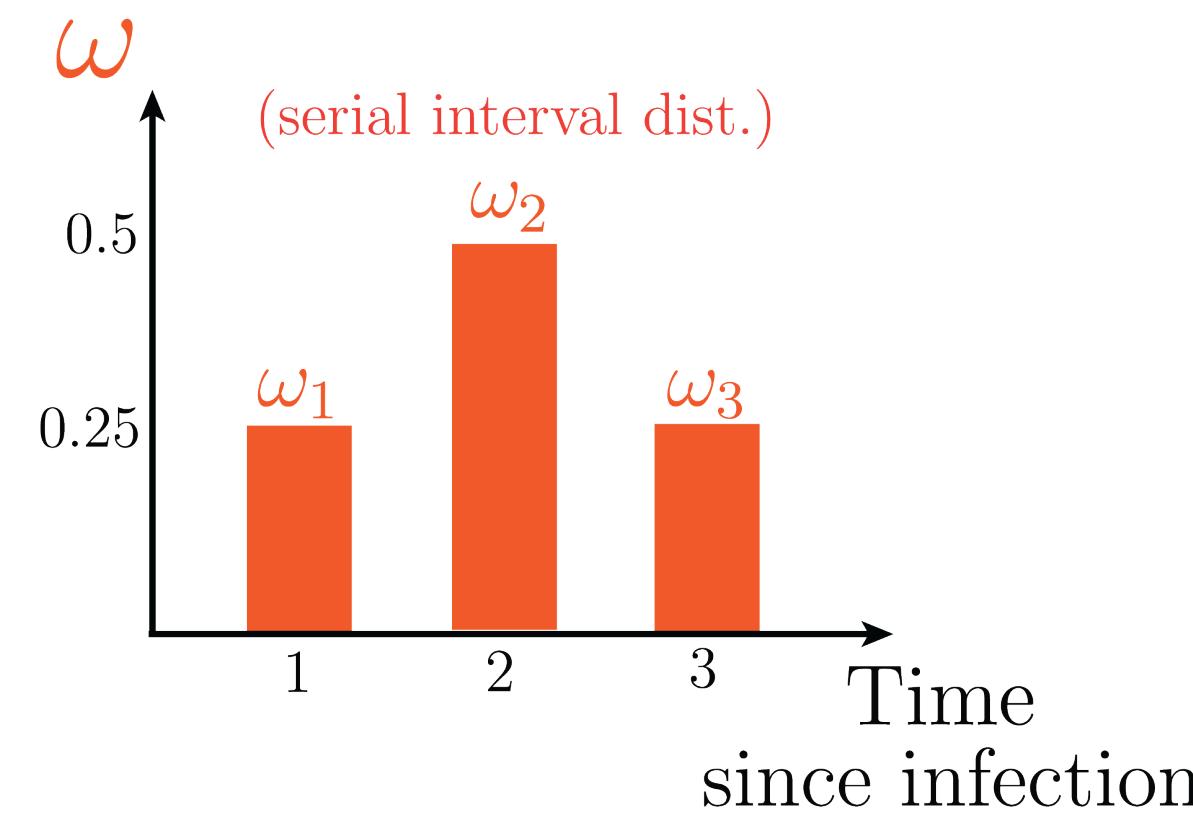
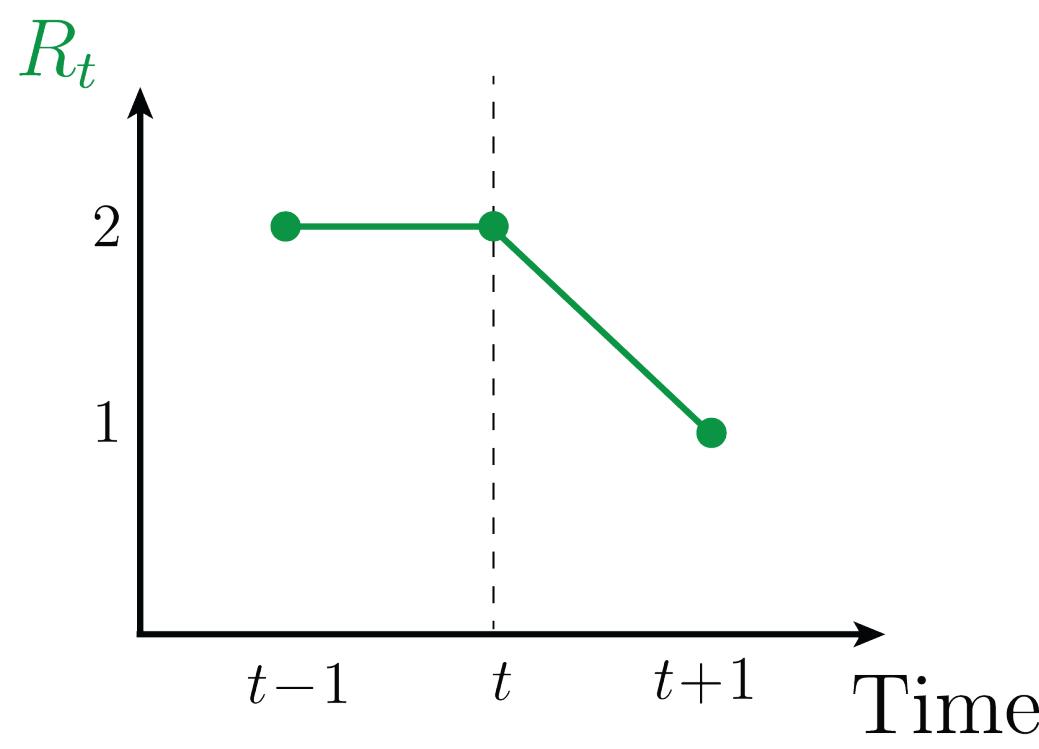


# The renewal model

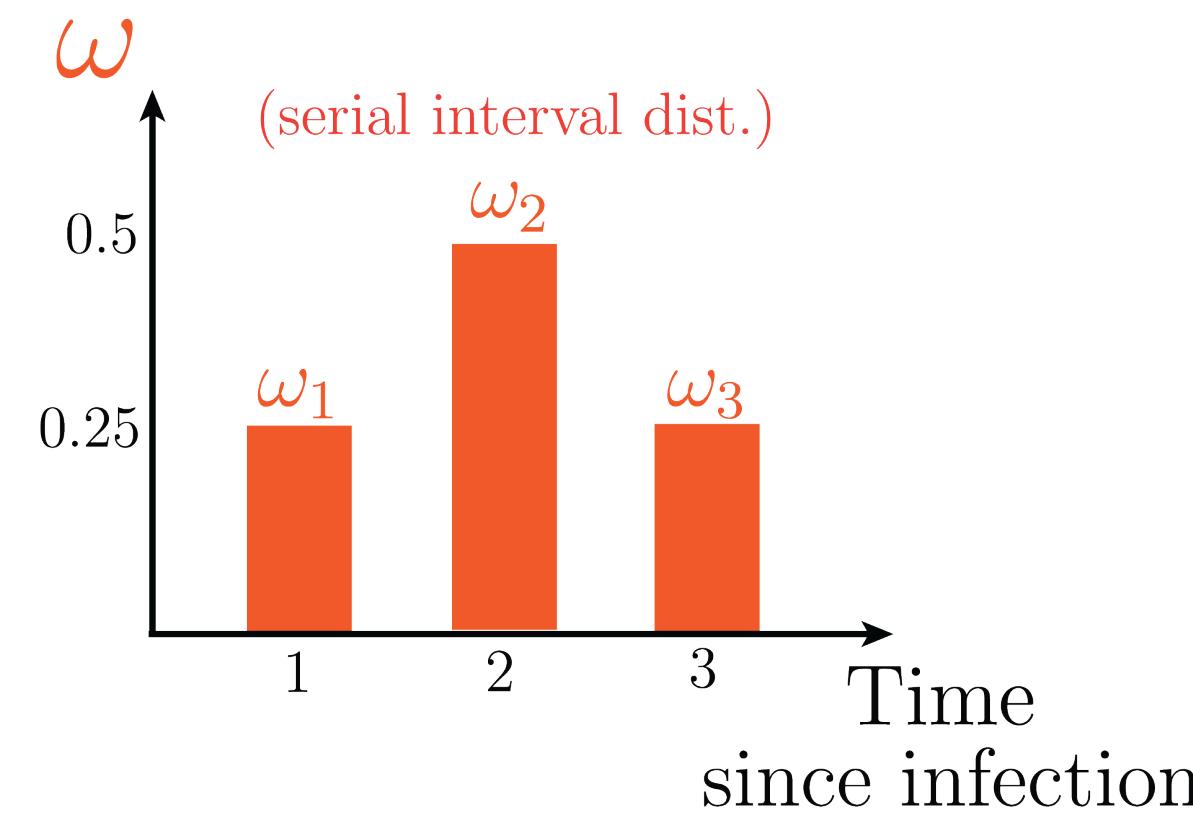
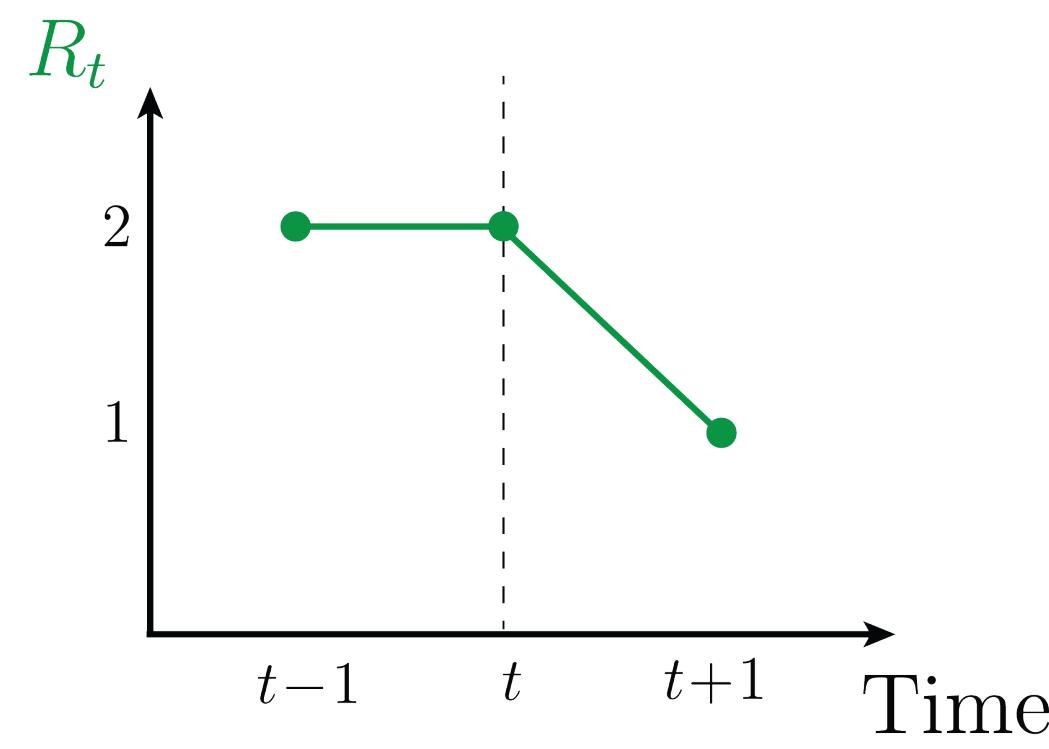
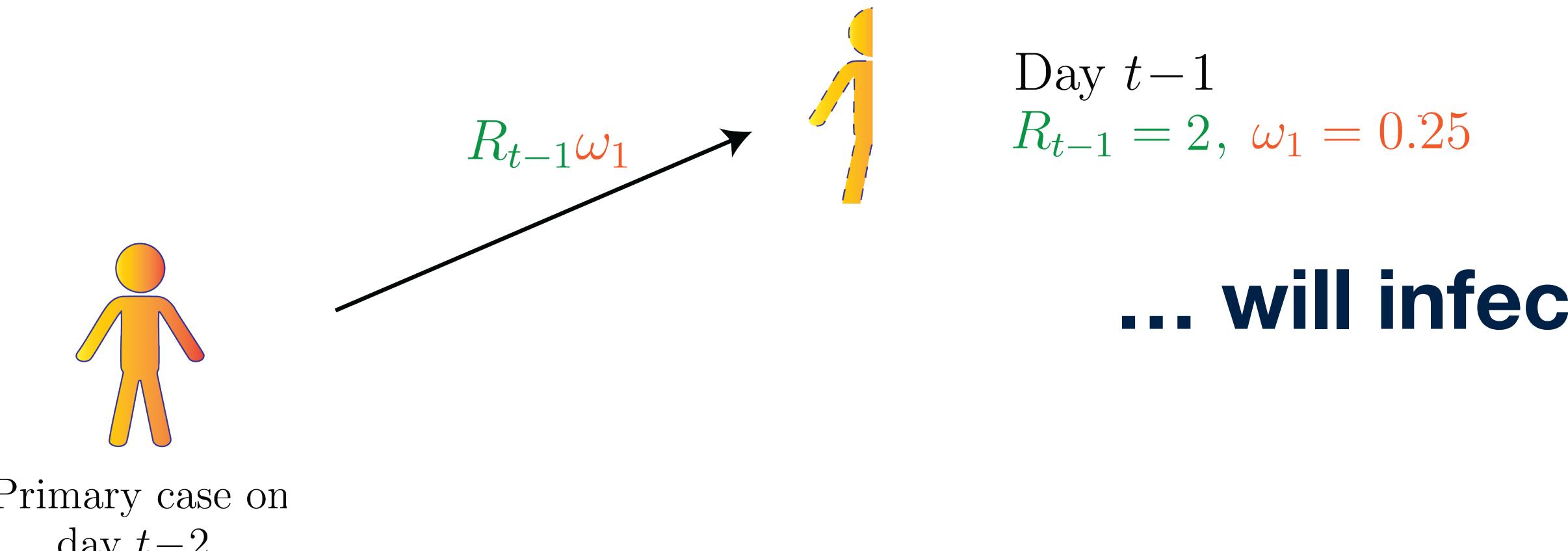


Primary case on  
day  $t-2$

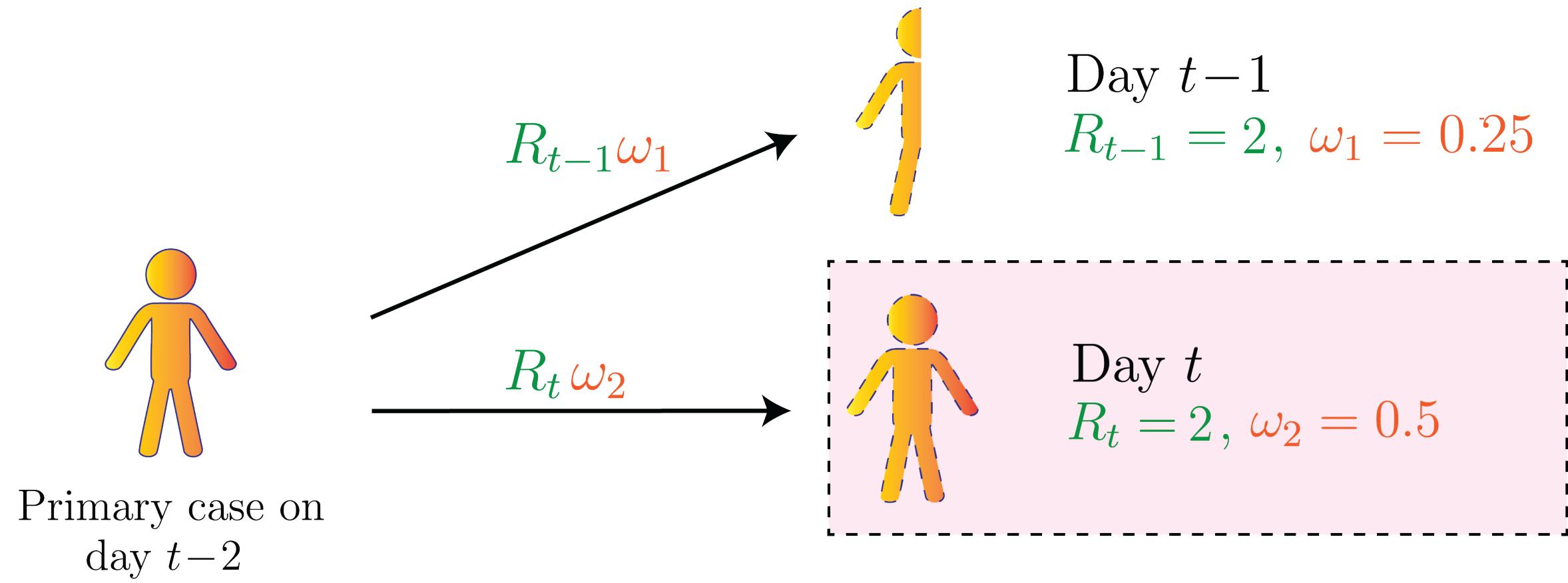
Someone infected two days ago...



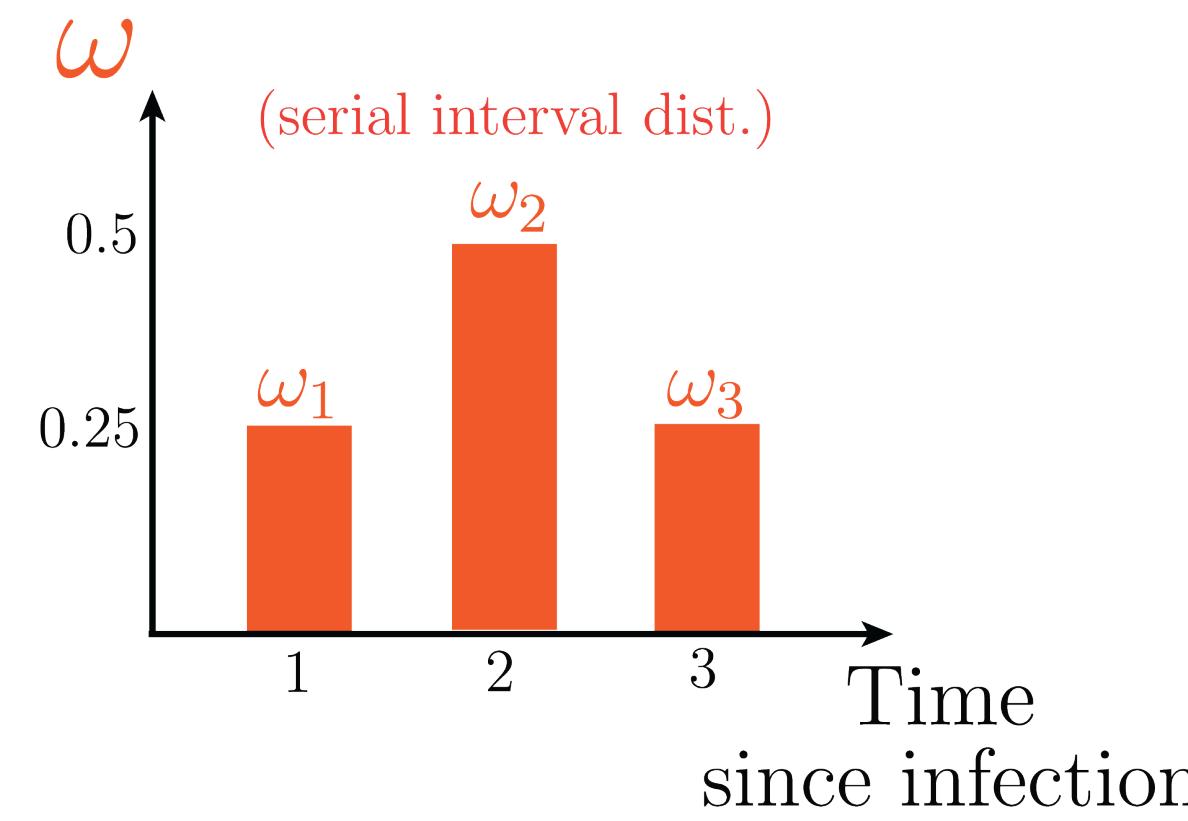
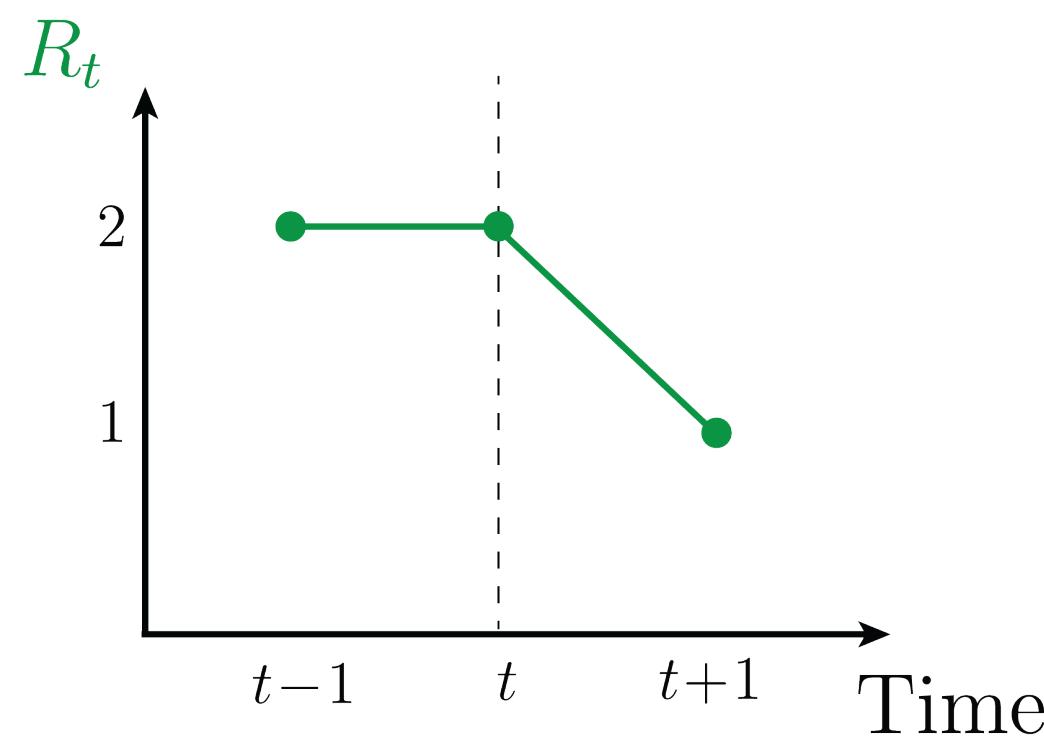
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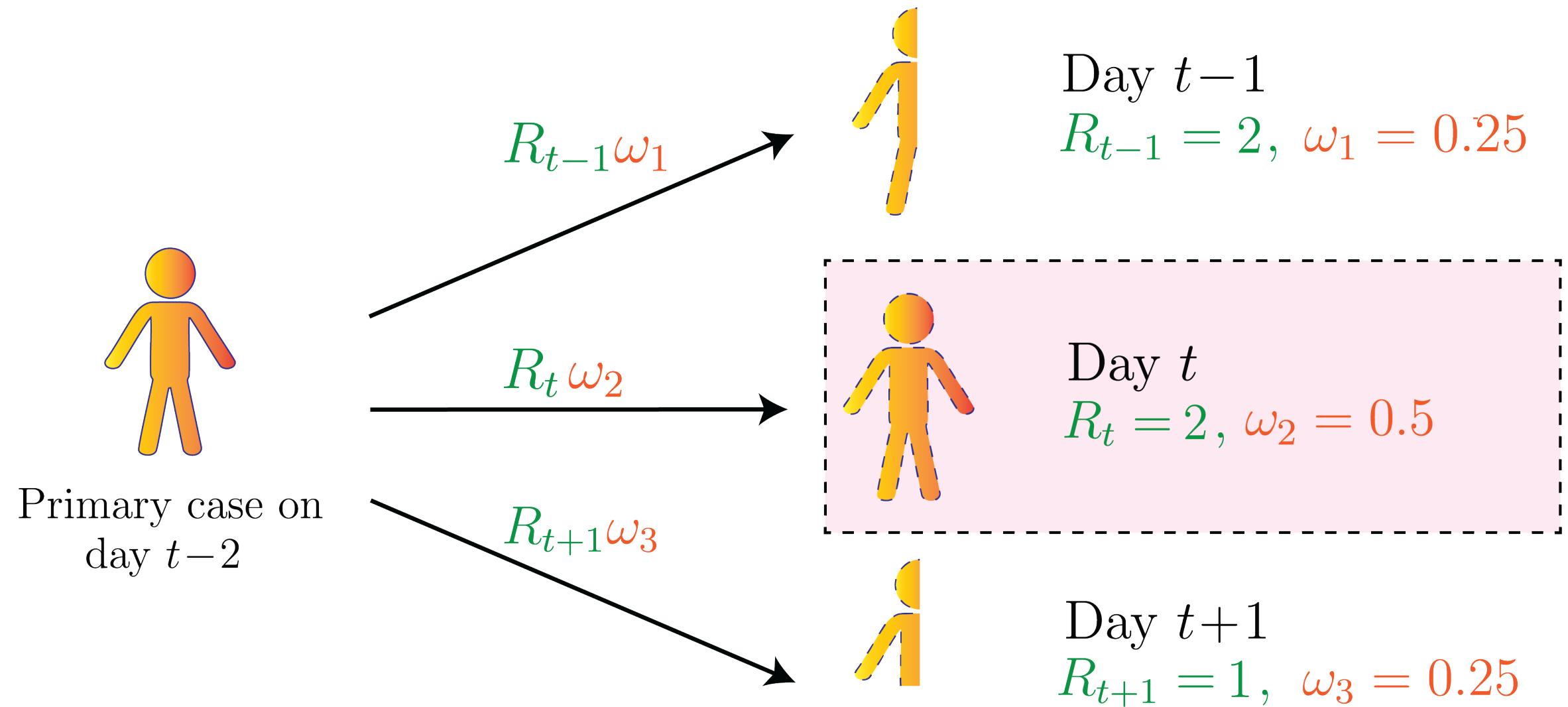
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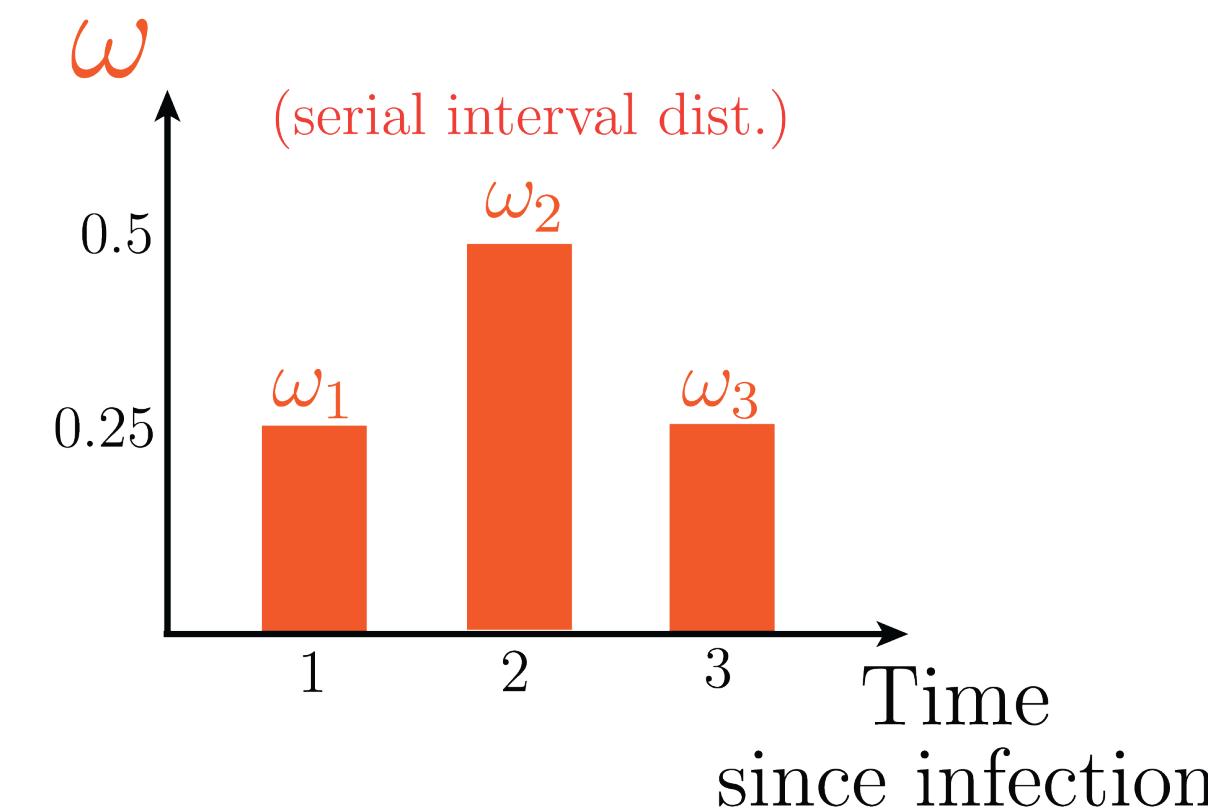
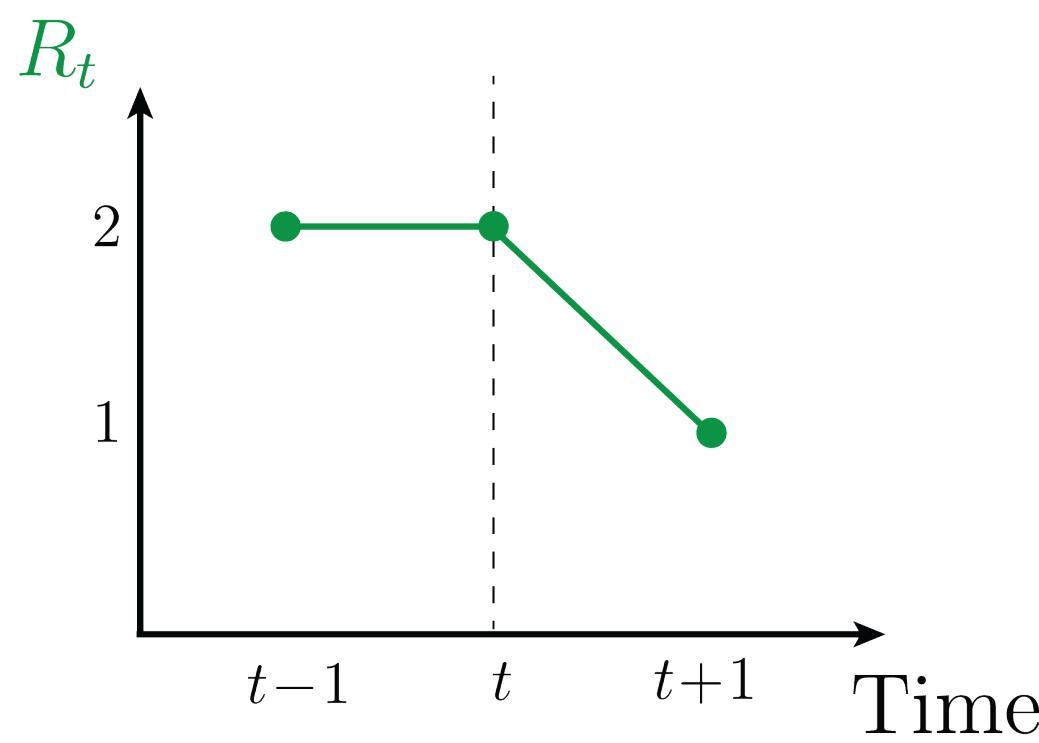
... an average of 1 person today, and...



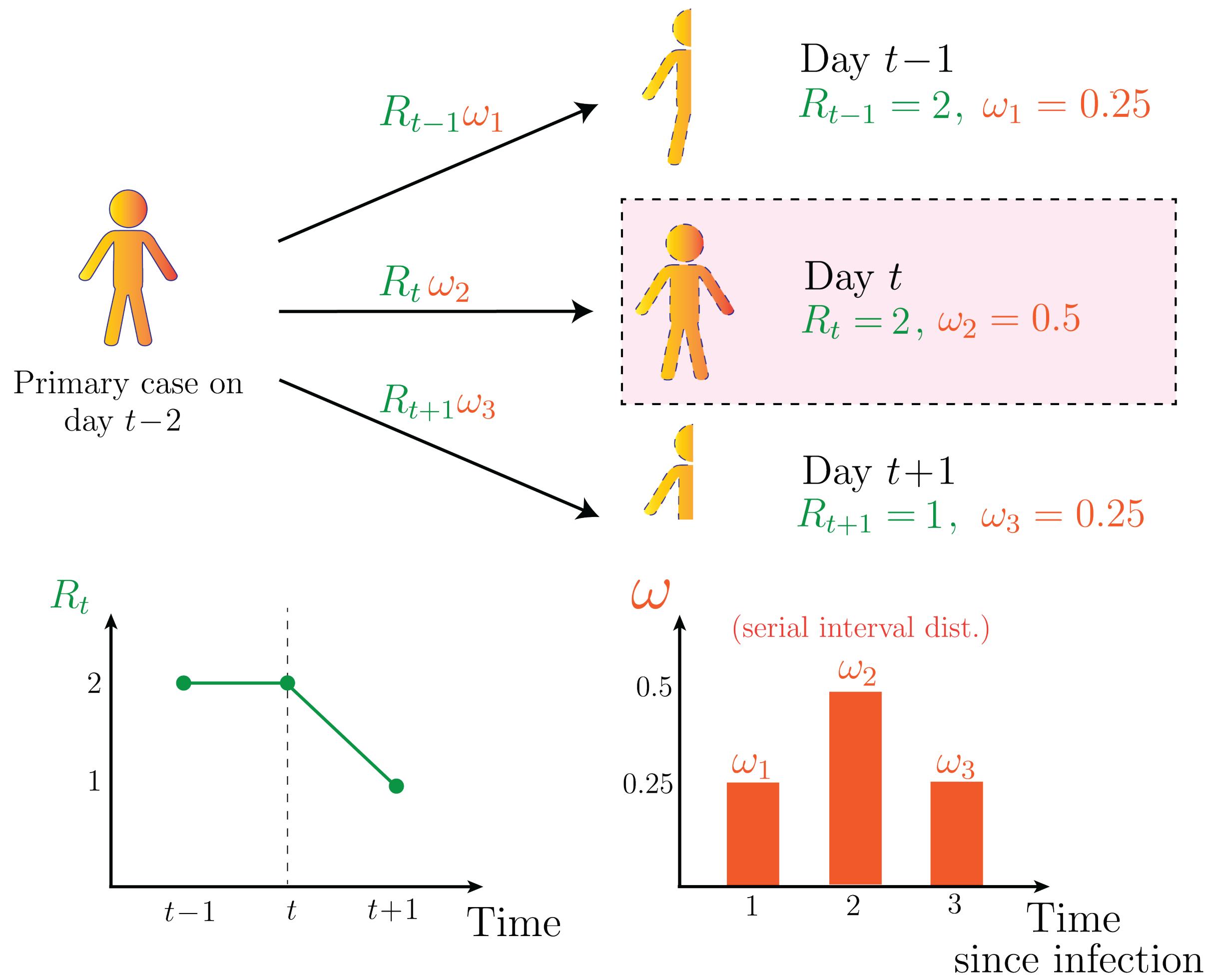
# The renewal model



... an average of 0.25 people tomorrow.



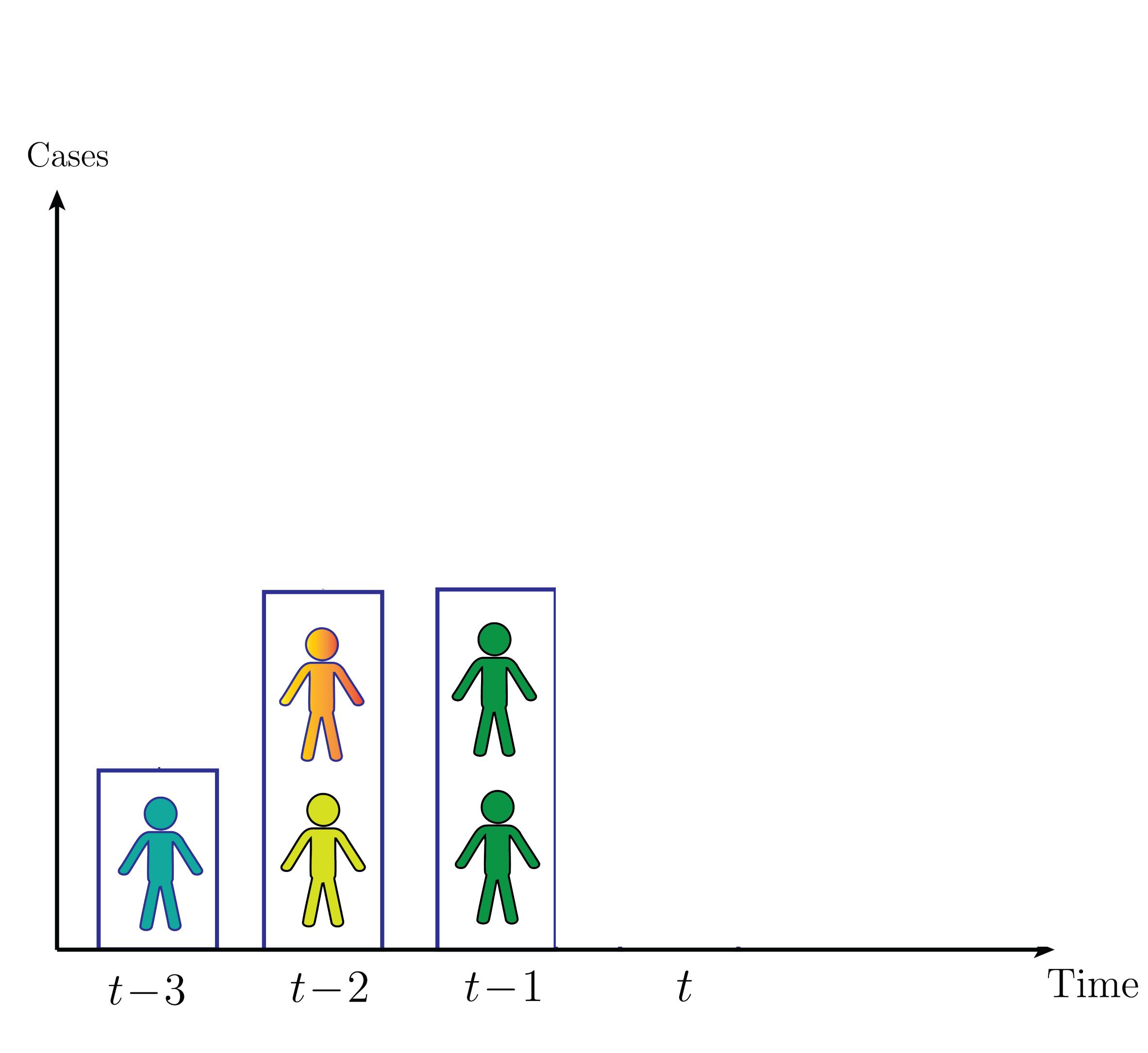
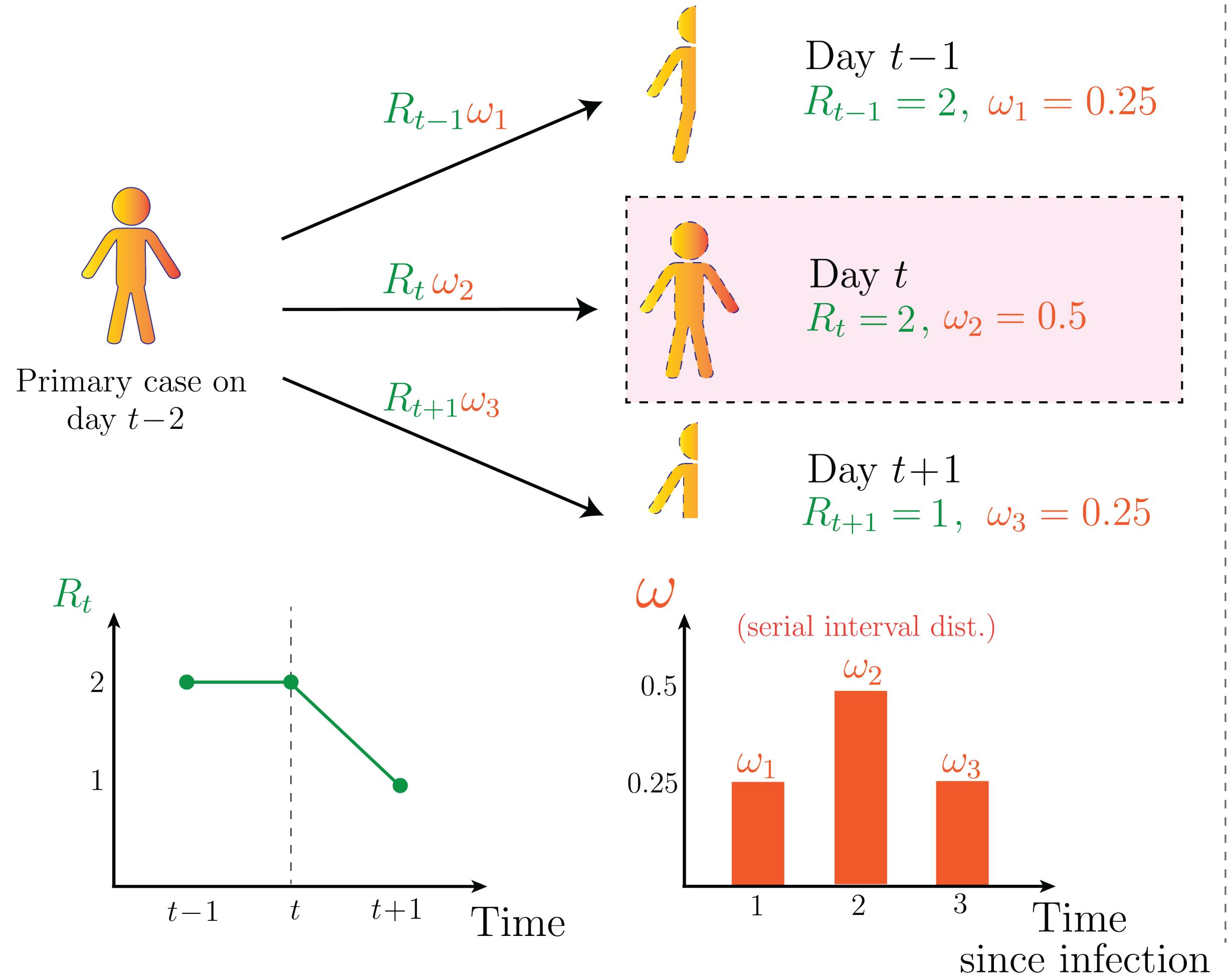
# The renewal model



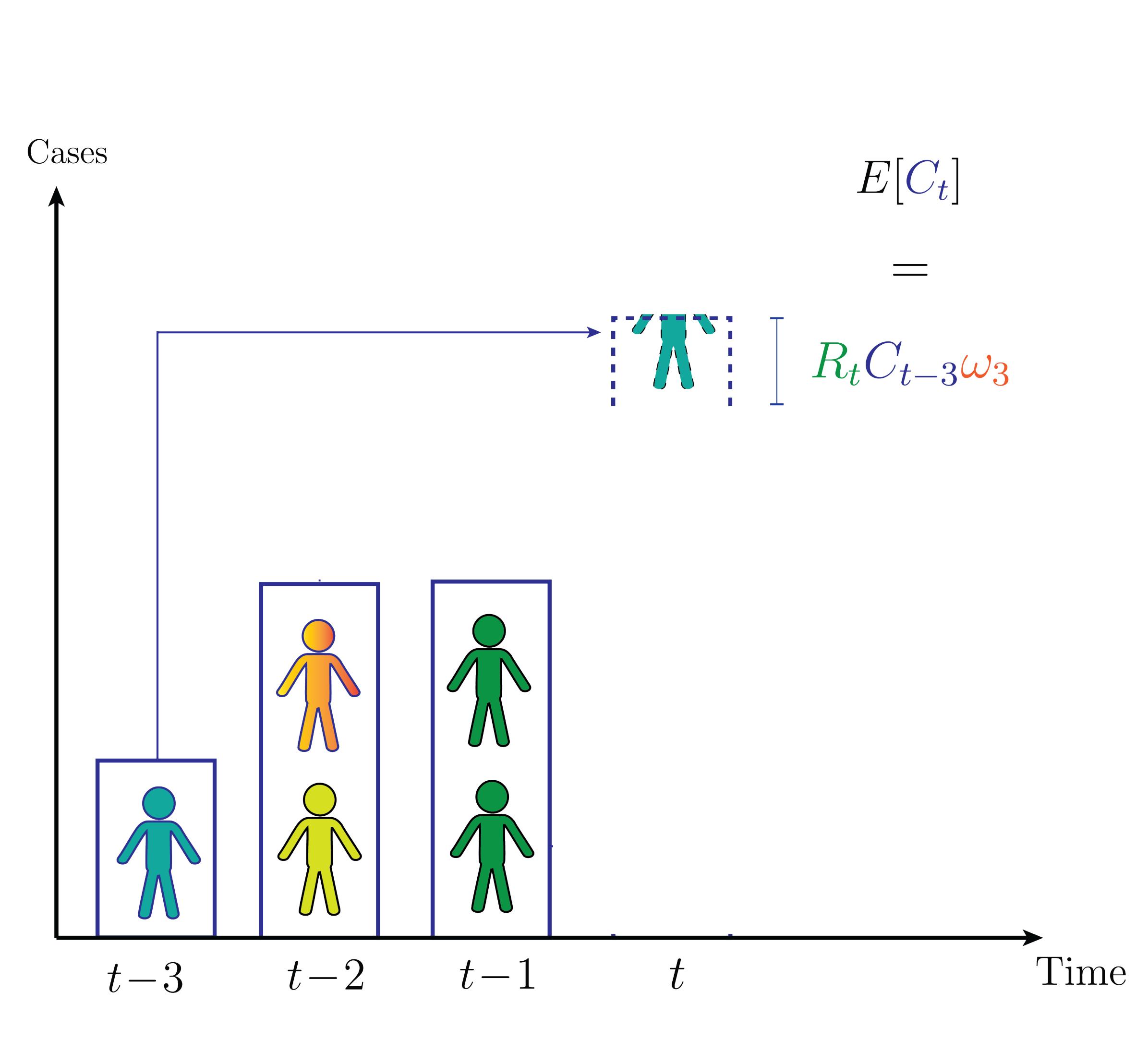
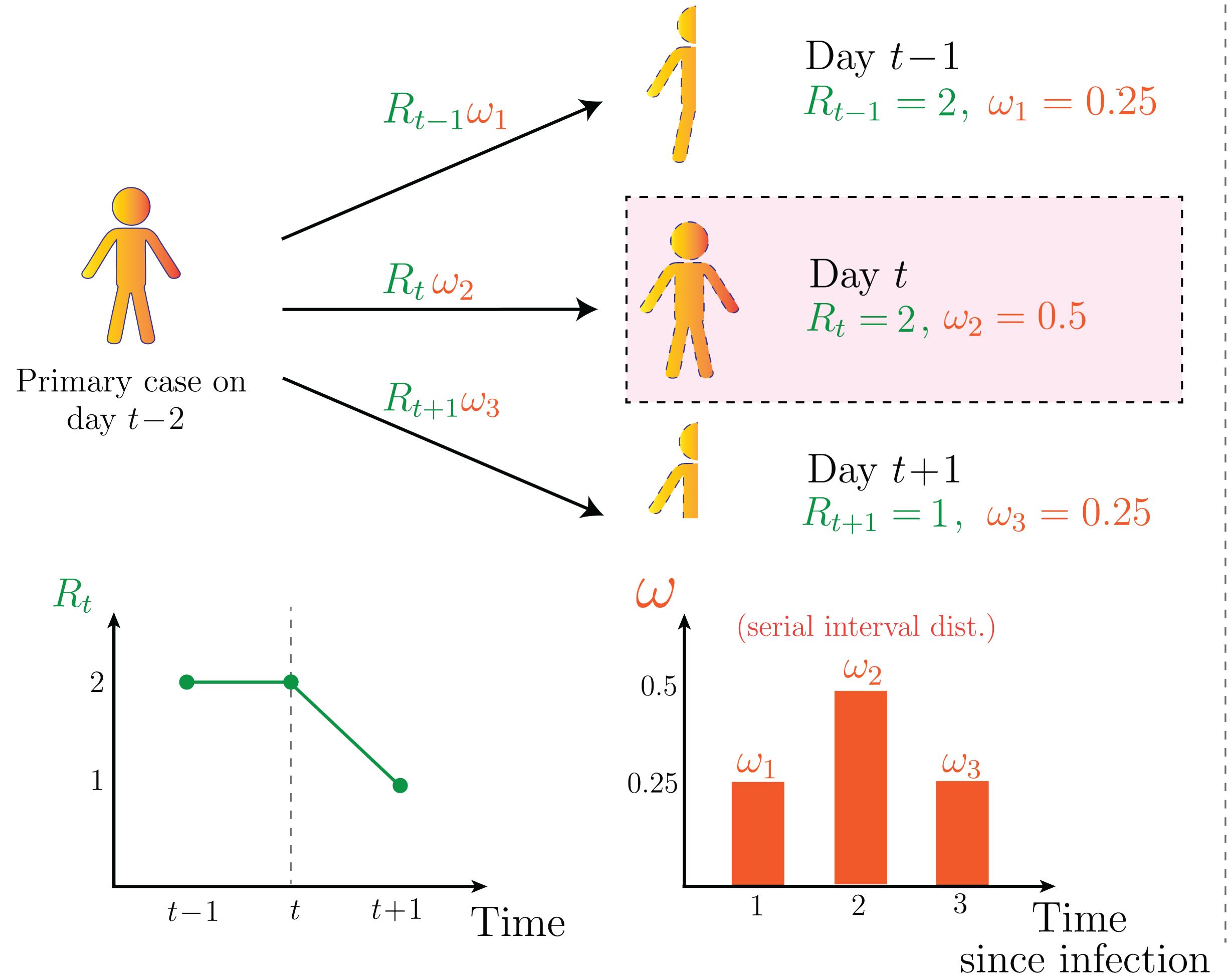
So...

**How many *total* cases  
do we expect today?**

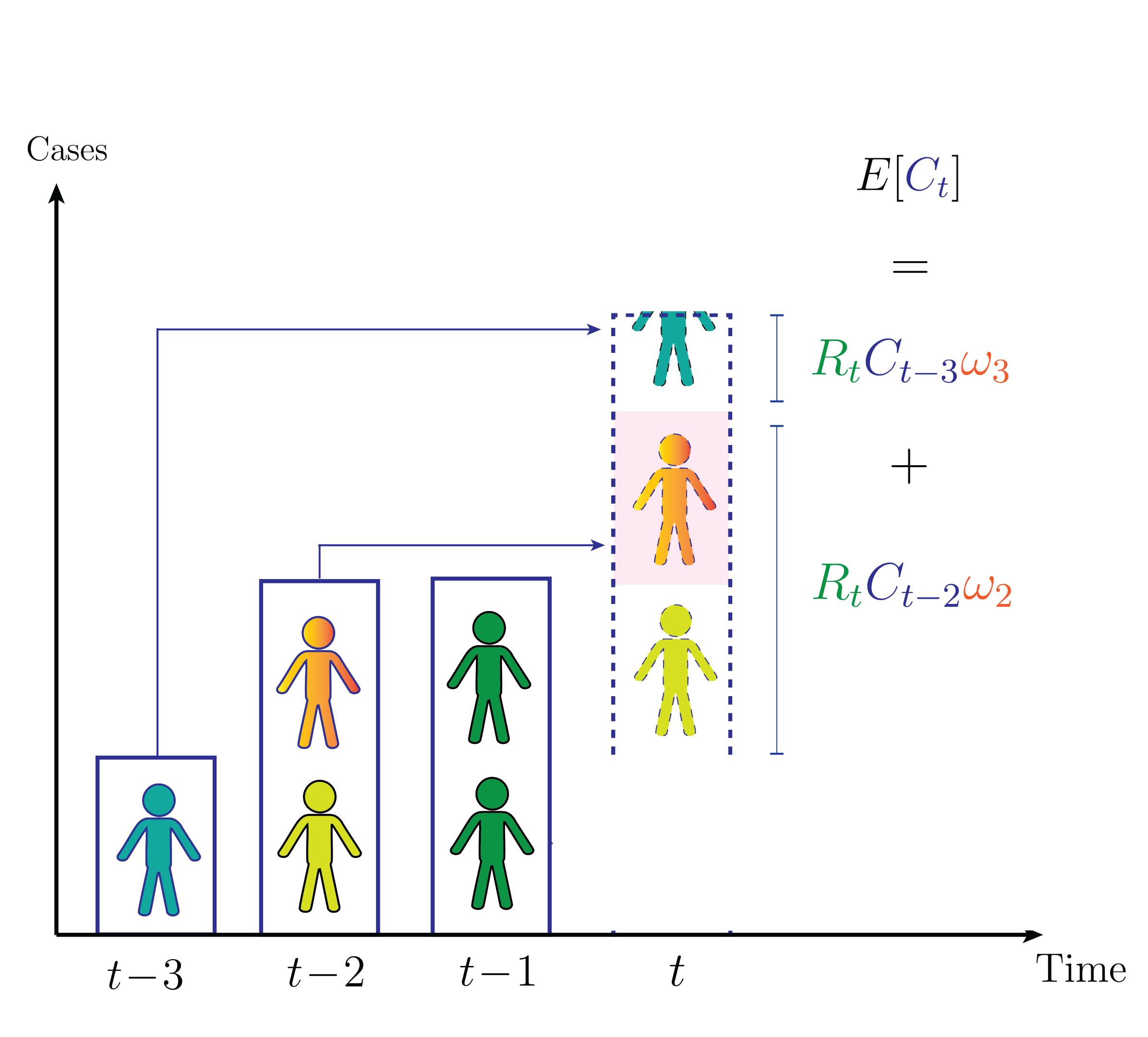
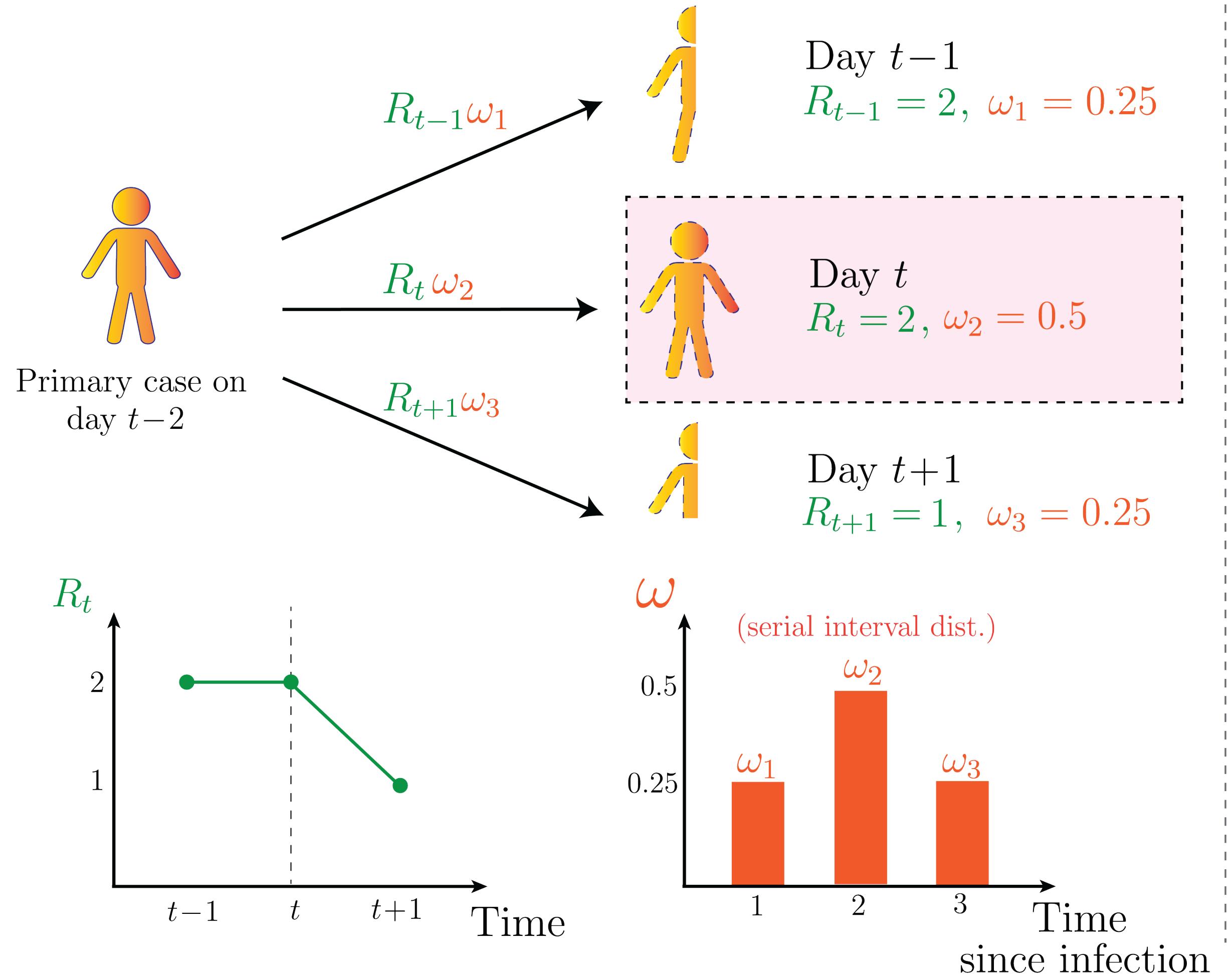
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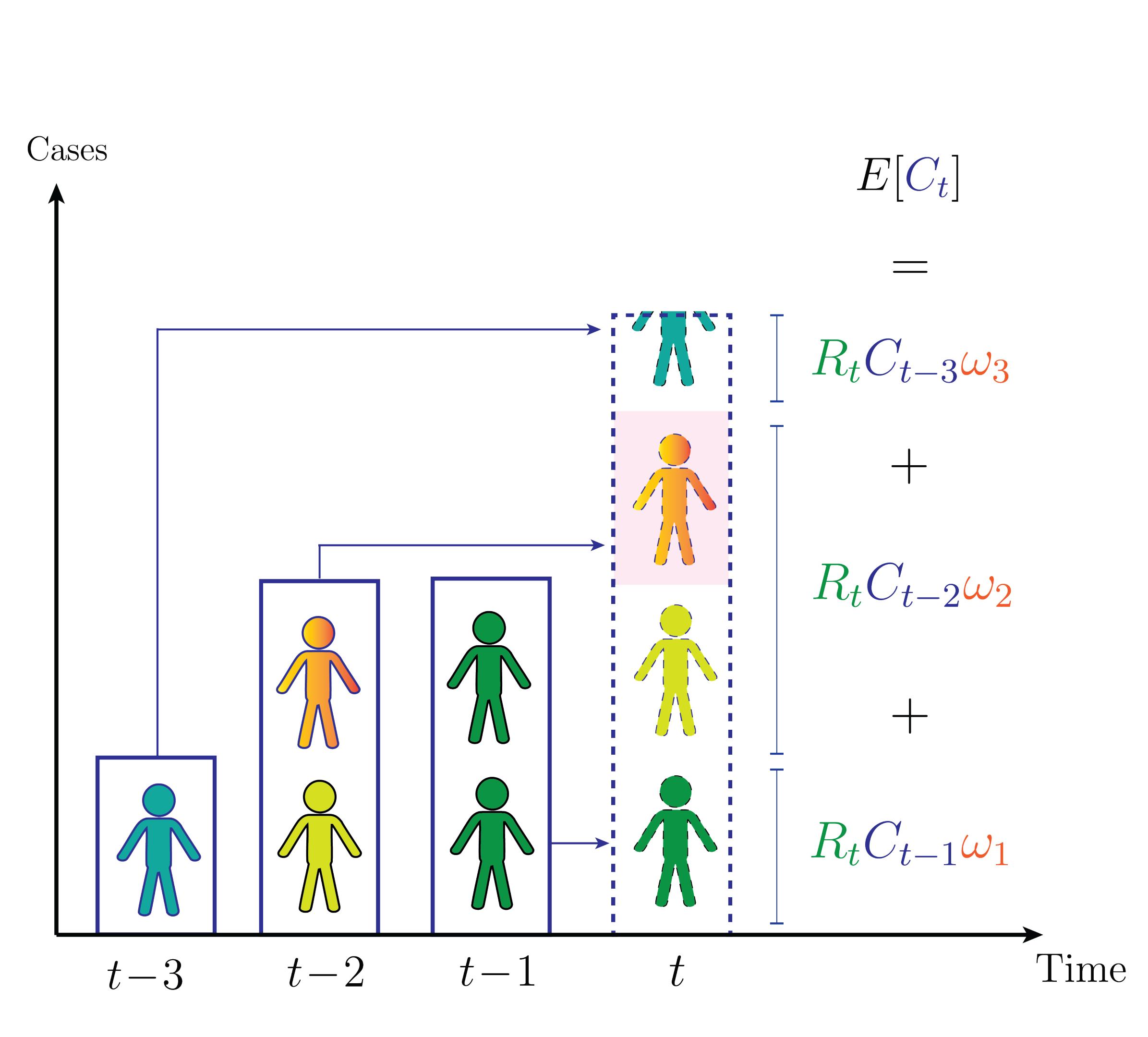
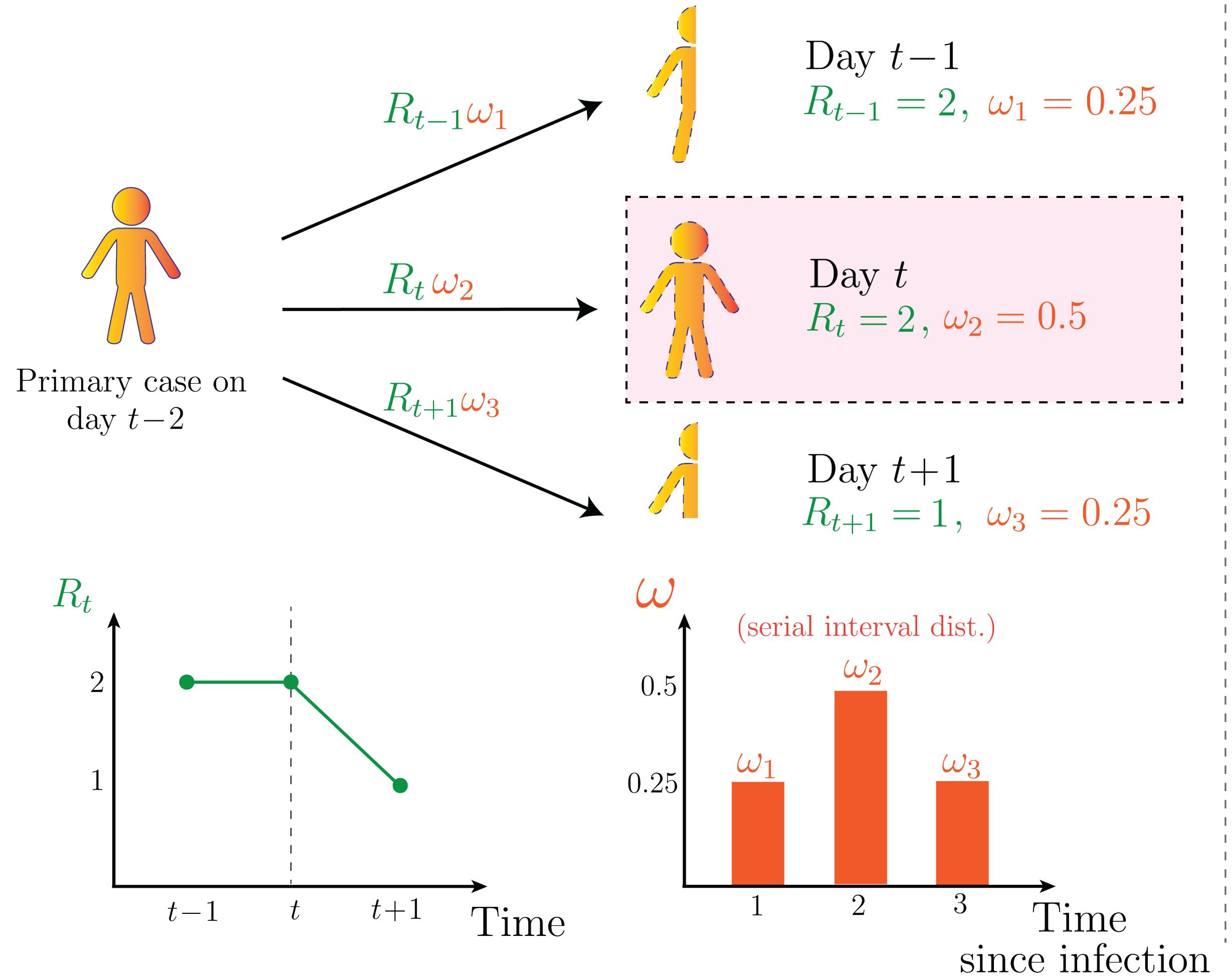
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# The renewal model



# The renewal model

- This is a **very** flexible model

*Prior for  $R_t$*

$$P(R_t)$$

×

*Likelihood*

$$P(C_t | R_t, C_{1:t-1}) = \text{Poiss} \left( R_t \sum C_{t-u} \omega_u \right)$$

=

*Posterior for  $R_t$*

$$P(R_t | C_{1:t})$$

**Pointwise estimates of  $R_t$**

*Dynamic model for  $R_t$*

$$P(R_t | R_{t-1}) = N(R_{t-1}, \sigma)$$

×

*Observation model*

$$P(C_t | R_t, C_{1:t-1}) = \text{Poiss} \left( R_t \sum C_{t-u} \omega_u \right)$$

=

*Posterior for  $R_t$*

$$P(R_t | C_{1:t})$$

**Smoothed estimates of  $R_t$**

*Dynamic model for  $R_t$*

$$P(R_t | R_{t-1}) = N(R_{t-1}, \sigma)$$

×

*Dynamic model for  $I_t$*

$$P(I_t | R_t, I_{1:t-1}) = \text{Poiss} \left( R_t \sum I_{t-u} \omega_u \right)$$

×

*Observation model*

$$P(C_t | I_t) = \text{Binomial}(I_t, \rho)$$

=

*Posterior for  $R_t$*

$$P(R_t | C_{1:t})$$

**+ observation noise**

# Renewal Models

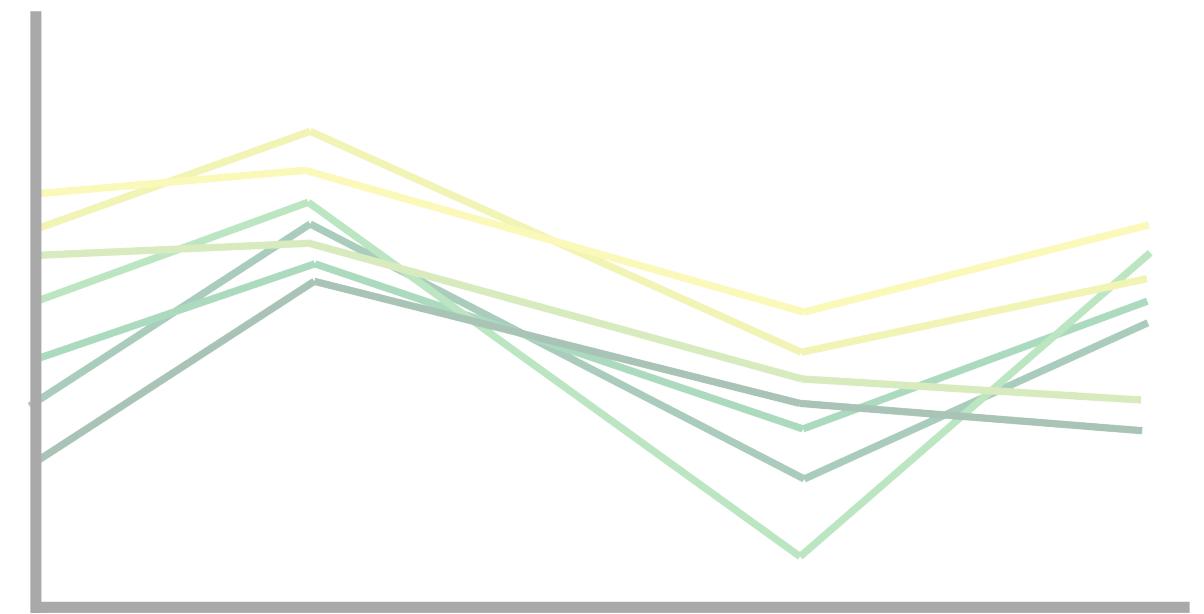
A simple ***model*** of an epidemic

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Used for reproduction number estimation

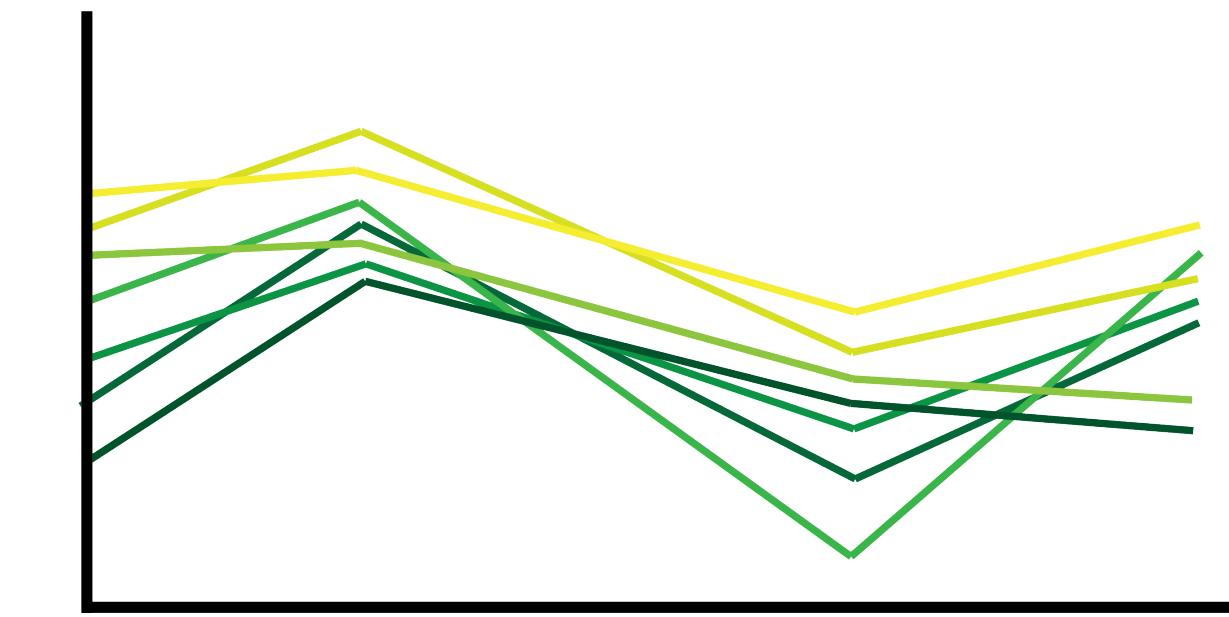
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# Sequential Monte Carlo

A **method** for fitting hidden-state models



Also known as “particle filters”

Very flexible!

Can account for many biases at once

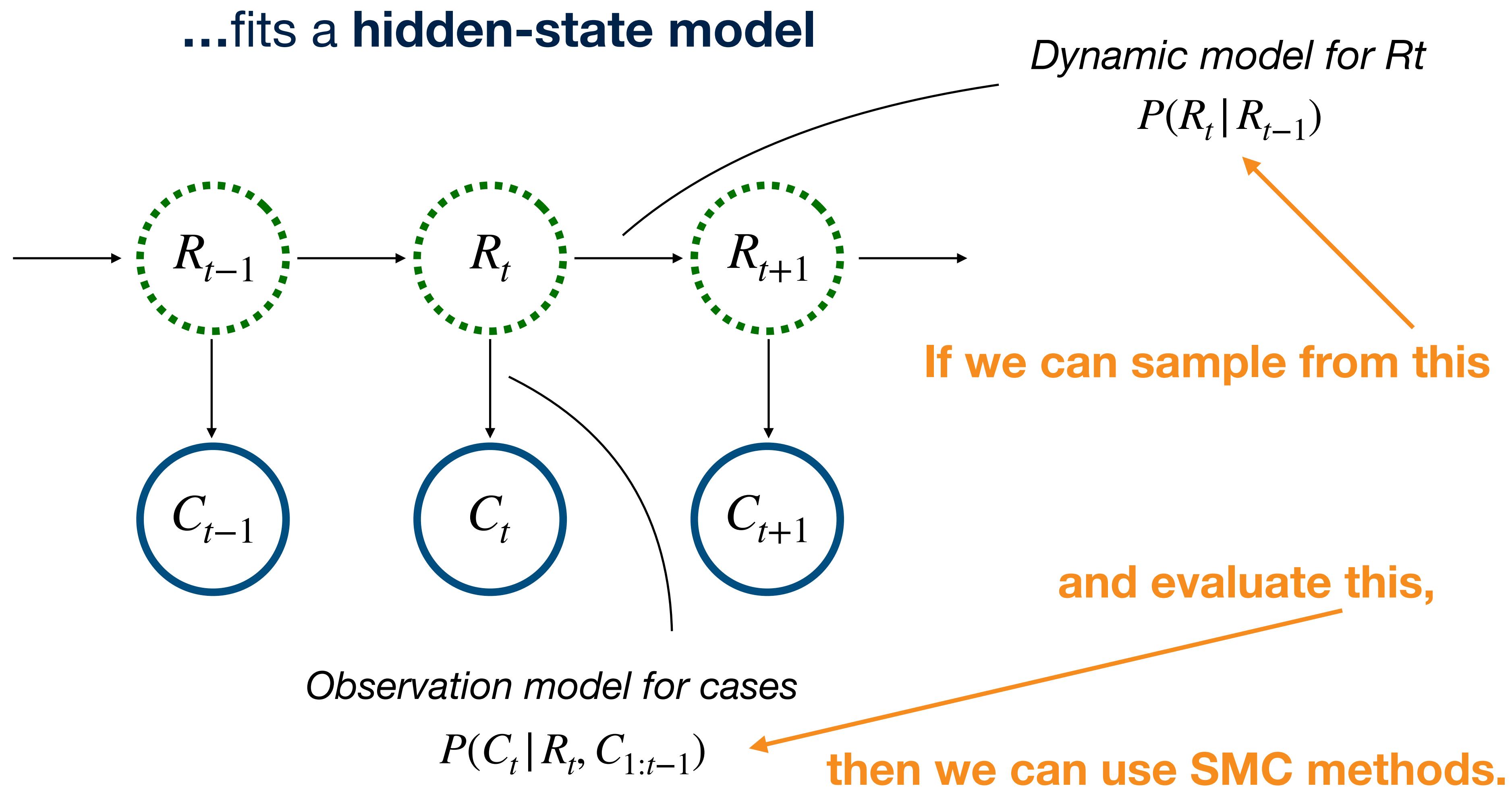
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# Sequential Monte Carlo

- Origins in **importance sampling** and **Bayesian filtering/smoothing** (e.g. Kalman filters)
- **Bootstrap filter (Gordon, 1993)**: the first method for non-linear non-Gaussian state-space models
- Then advanced by Doucet, Del Moral, Chopin, Kantas, Andrieu and many others\*
- We consider **hidden-state estimation** and **parameter estimation** separately

*(These names are listed as they are associated with helpful tutorial/summary/overview papers!)*

# Sequential Monte Carlo: the bootstrap filter...



# Sequential Monte Carlo: the bootstrap filter

Assume we have samples:

$$R_t^{(i)} \sim P(R_t | C_{1:t}), \quad i = 1, \dots, N$$

We update these samples at  $t+1$  by:

**1. Projecting** according to the dynamic model

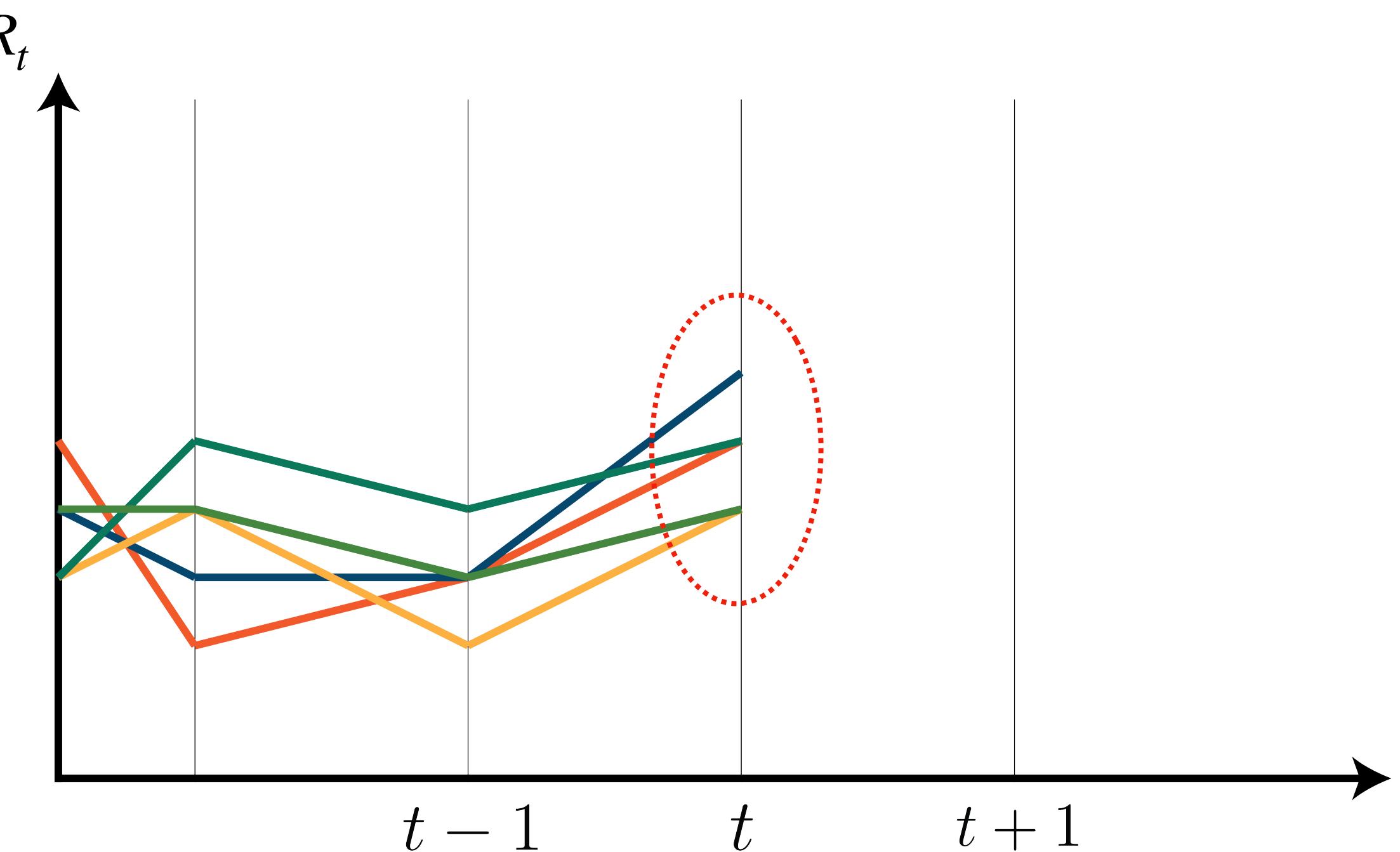
$$\tilde{R}_{t+1}^{(i)} \sim P(R_{t+1} | R_t^{(i)})$$

**2. Weighting** according to the observation model

$$W_{i,t+1} = P(C_{t+1} | \tilde{R}_{t+1}^{(i)}, C_{1:t})$$

**3. Resampling** projected “particles”, giving:

$$R_{t+1}^{(i)} \sim P(R_{t+1} | C_{1:t+1}), \quad i = 1, \dots, N$$



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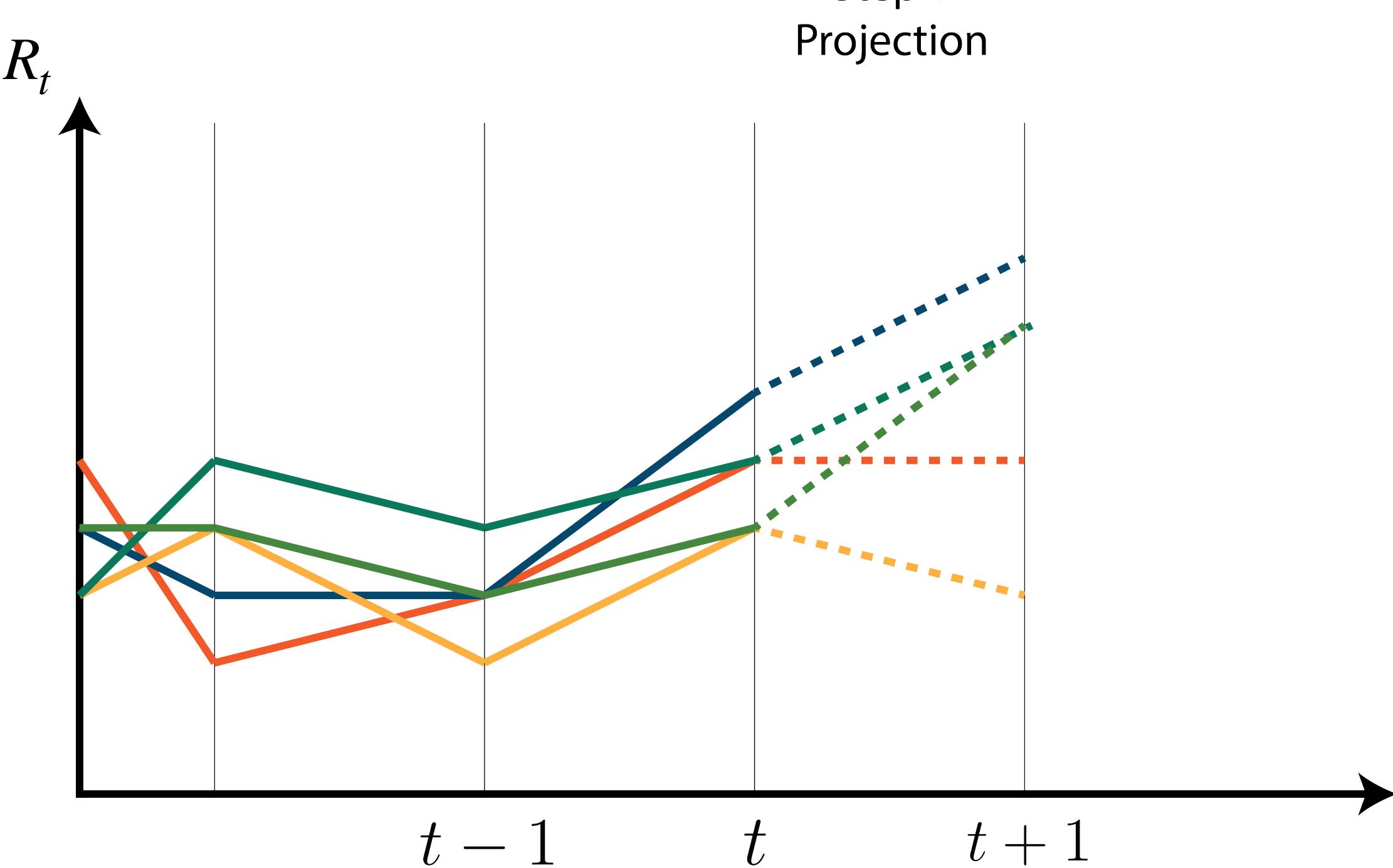
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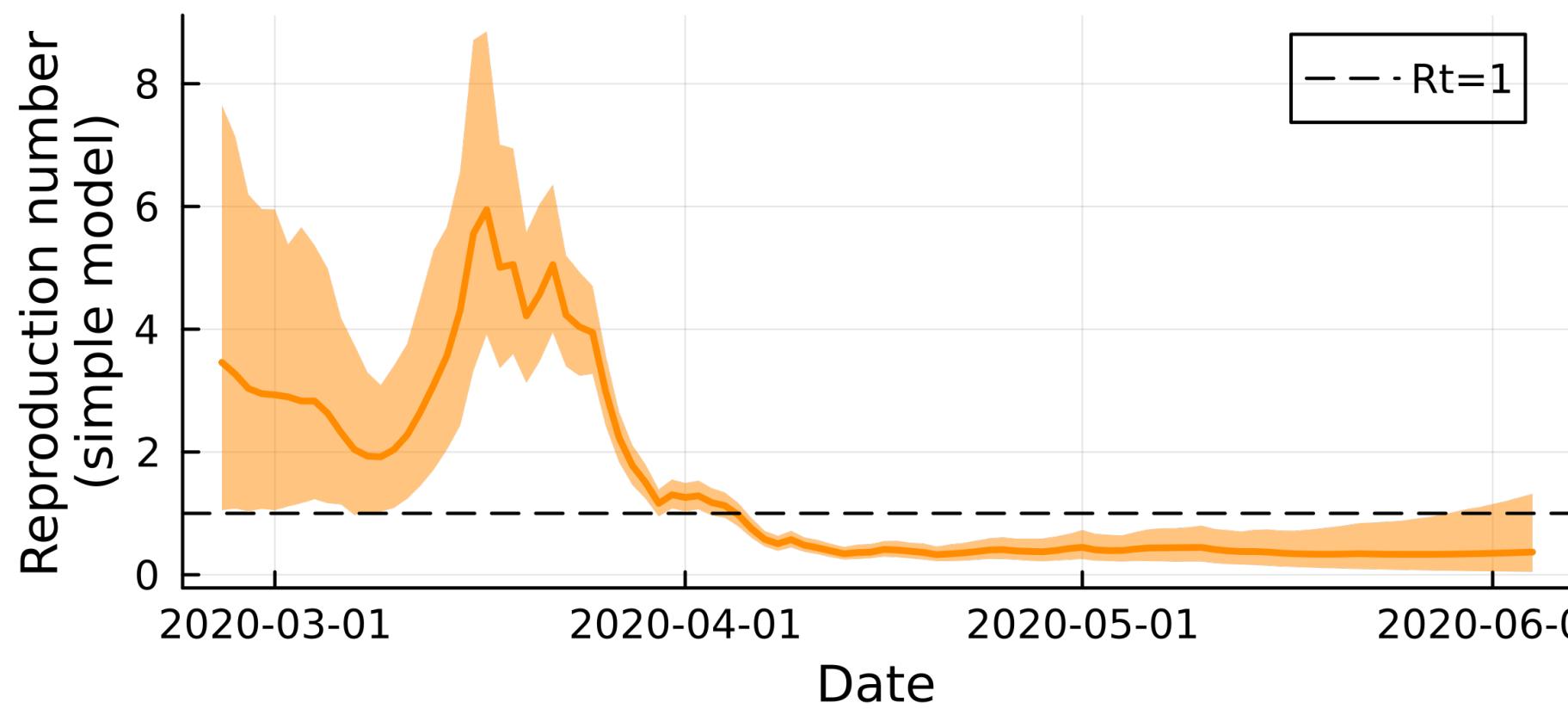
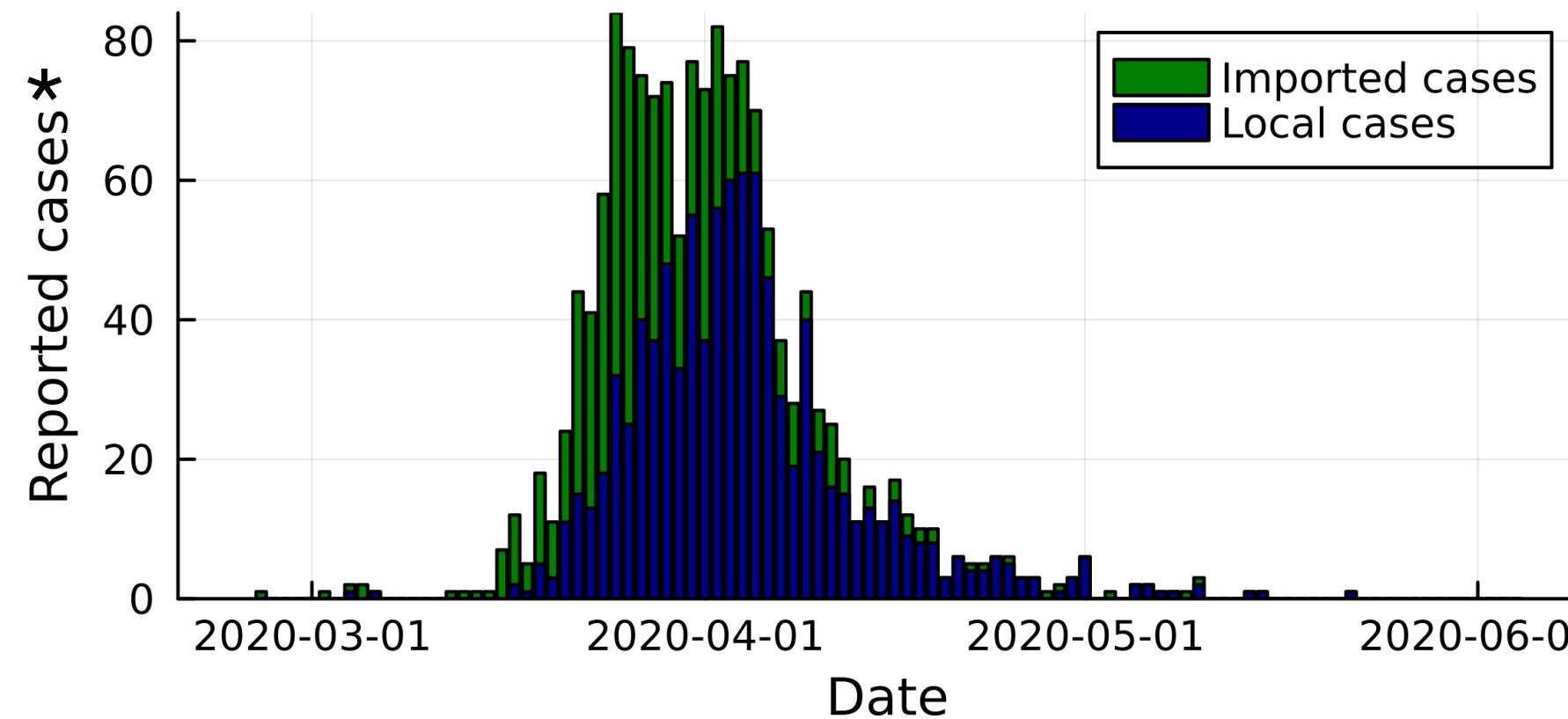
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# A simple example...



\* Reported case data are from the first 100 days of the COVID-19 outbreak in New Zealand

## Dynamic model for $R_t$

$$\log R_t | R_{t-1} \sim \text{Normal}(\log R_{t-1}, \sigma)$$

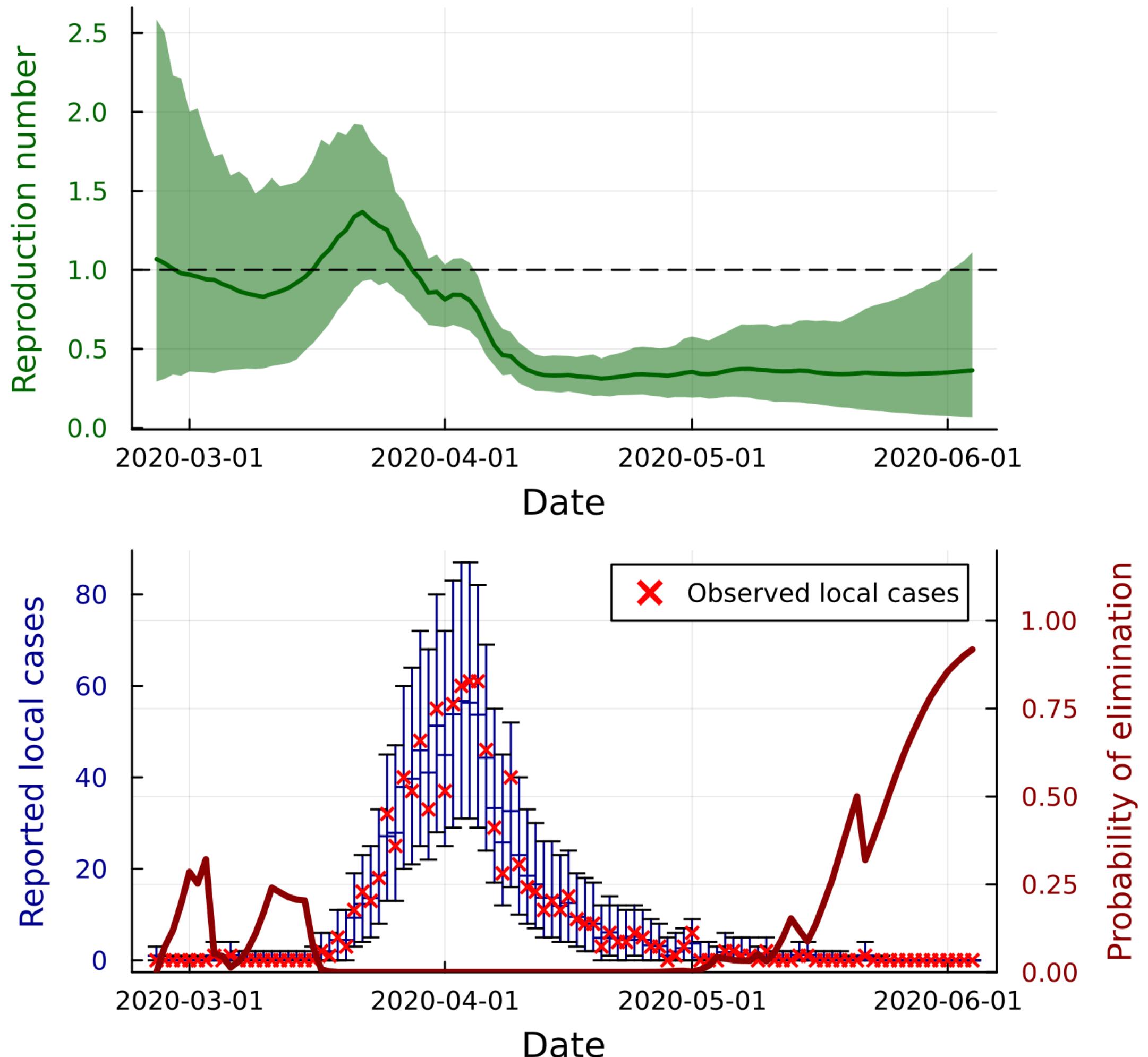
## Observation model

$$C_t \sim \text{Poisson}\left(R_t \sum_{u=1}^{\omega_{\max}} C_{t-u} \omega_u\right)$$

The entire algorithm requires only 11 lines of code:

```
# Setup:  
N = 100000 # Number of particles  
ω = pdf.Gamma(2.36, 2.74), 1:100 # Serial interval  
Y = loadData("NZCOVID") # Load data  
R = zeros(N, 100) # Matrix to store particle values (of log Rt)  
  
# Run the bootstrap filter:  
R[:,1] = log.(rand(Uniform(0, 10), N)) # Sample initial values  
for tt = 2:100  
    R[:,tt] = rand.Normal.(R[:,tt-1], 0.2) # Project (log) Rt  
    Λ = sum(Y.Ct[(tt-1):-1:1] .* ω[1:(tt-1)]) # Calculate the force of infection  
    W = pdf.Poisson.(Λ * exp.(R[:,tt])), Y.Ct[tt]) # Calculate weights  
    R[1:N, max(tt-40, 1):tt] = R[wsample(1:N, W, N), max(tt-40, 1):tt] # Resample  
end
```

# Imported cases, reporting noise, and elimination



**Dynamic model for  $R_t$**

$$\log R_t | R_{t-1} \sim \text{Normal}(\log R_{t-1}, \sigma)$$

$$I_t | R_t, I_{1:t-1} \sim \text{Poisson} \left( R_t \sum_{u=1}^{\omega_{\max}} (I_{t-u} + M_{t-u}) \omega_u \right)$$

**Observation model**

$$C_t \sim \text{Negative binomial}(\dots)$$

**Probability of elimination** estimated by simulating the model forward two weeks and checking whether any new infections occur.

# Sequential Monte Carlo: Parameter estimation

- A much harder and more expensive problem!
- We use Particle Marginal Metropolis Hastings (PMMH)

**Dynamic model for  $R_t$**

$$\log R_t | R_{t-1} \sim \text{Normal}(\log R_{t-1}, \sigma)$$

**Observation model**

$$C_t \sim \text{Poisson}\left(R_t \sum_{u=1}^{\omega_{\max}} C_{t-u} \omega_u\right)$$

# Sequential Monte Carlo: Parameter estimation

- Predictive decomposition of the likelihood:

$$L(\theta) = P(C_{1:T} | \theta) = P(C_1 | \theta) \prod_{t=2}^T \underbrace{P(C_t | C_{1:t-1}, \theta)}_{\text{one-step-ahead likelihood}}$$

- Where the one-step-ahead likelihood is just the average of the bootstrap filter weights:

$$\overbrace{P(C_t | C_{1:t-1}, \theta)}^{\text{one-step-ahead likelihood}} = E_{R_t | C_{1:t-1}}[P(C_t | R_t, C_{1:t-1}, \theta)] \approx \frac{1}{N} \sum_{i=1}^N W_{t,i}$$

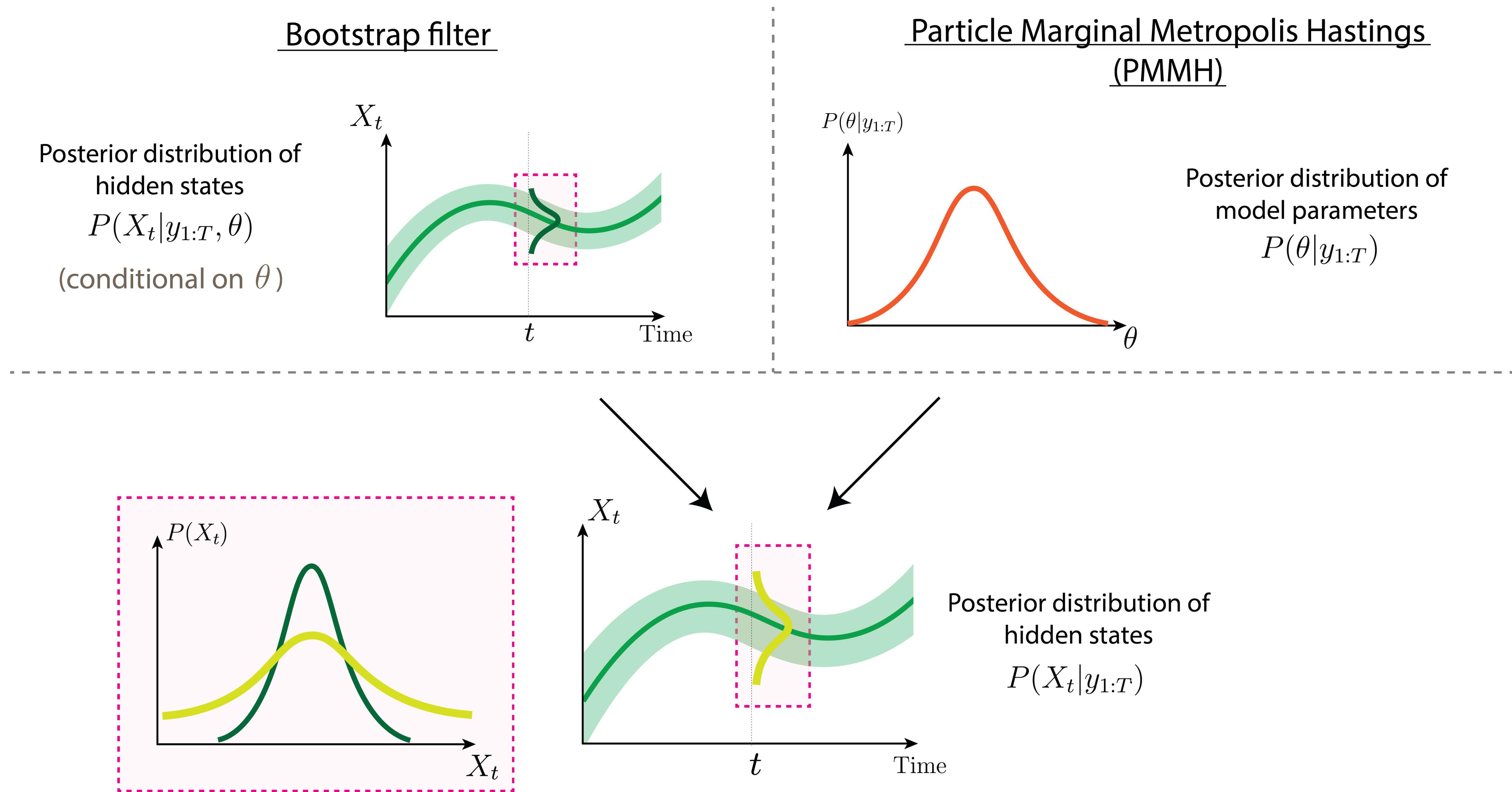
# Sequential Monte Carlo: Parameter estimation

- To estimate the log-likelihood, we run the bootstrap filter and calculate:

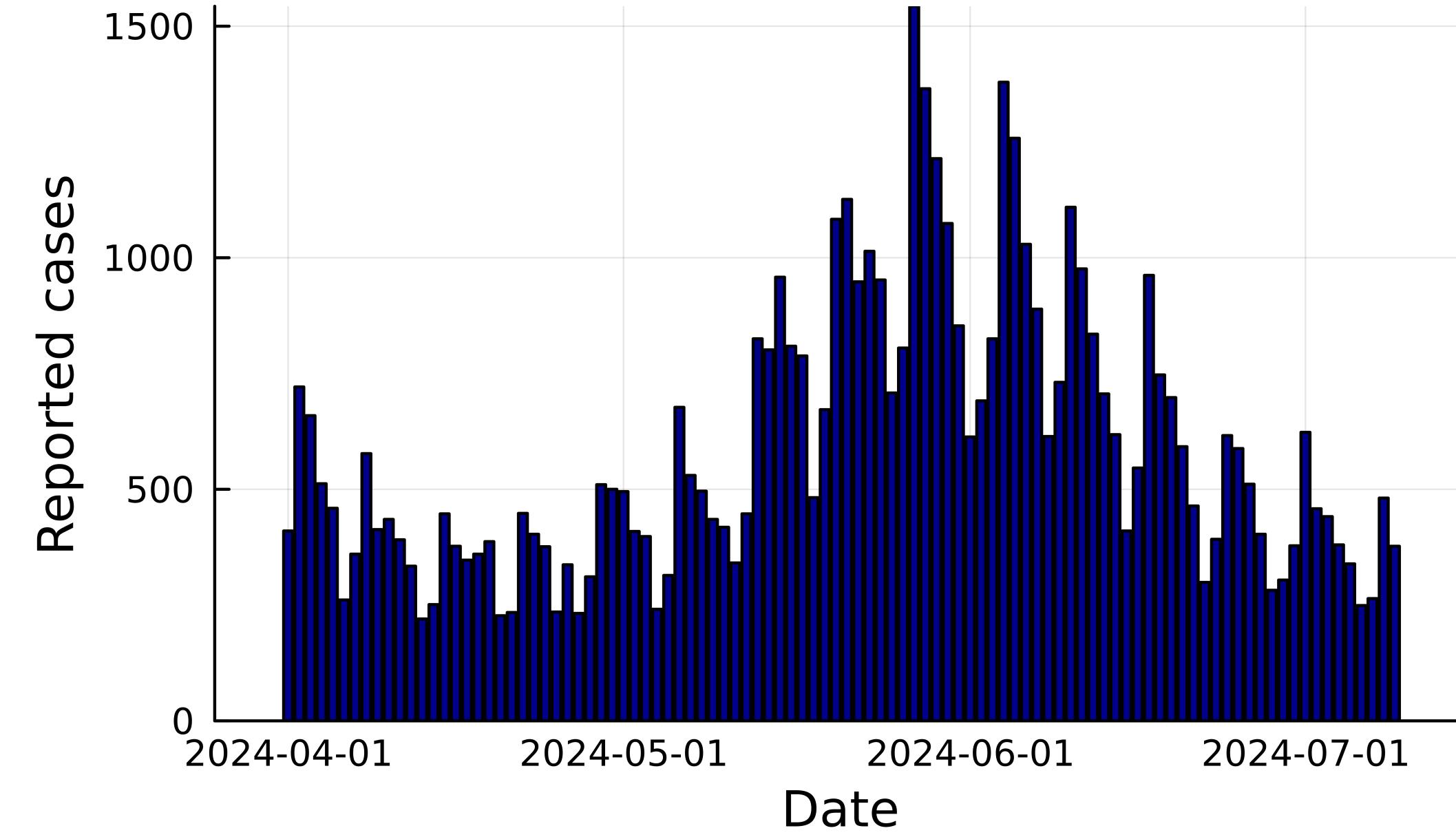
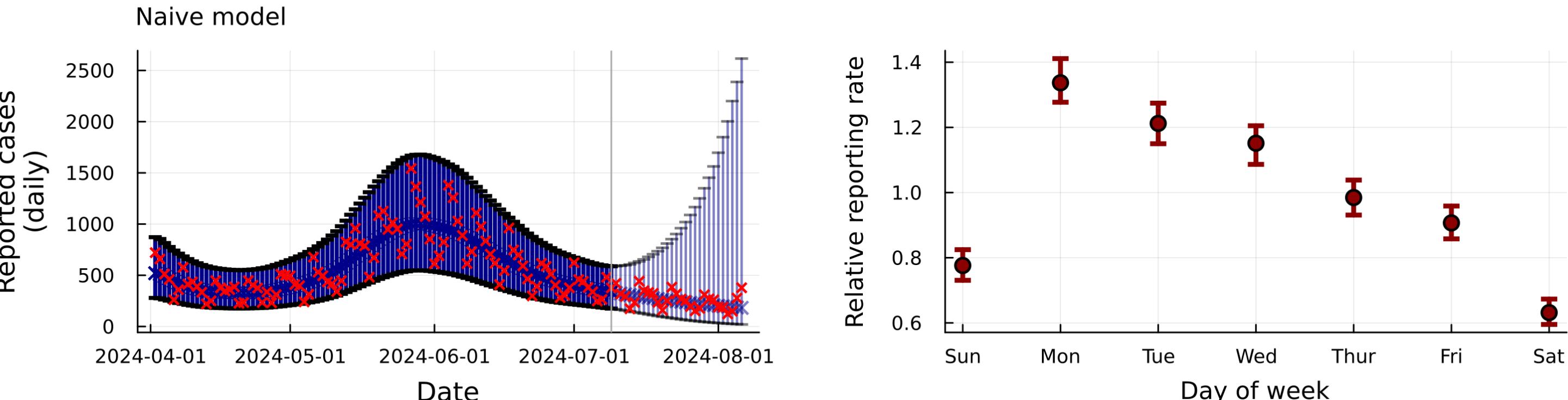
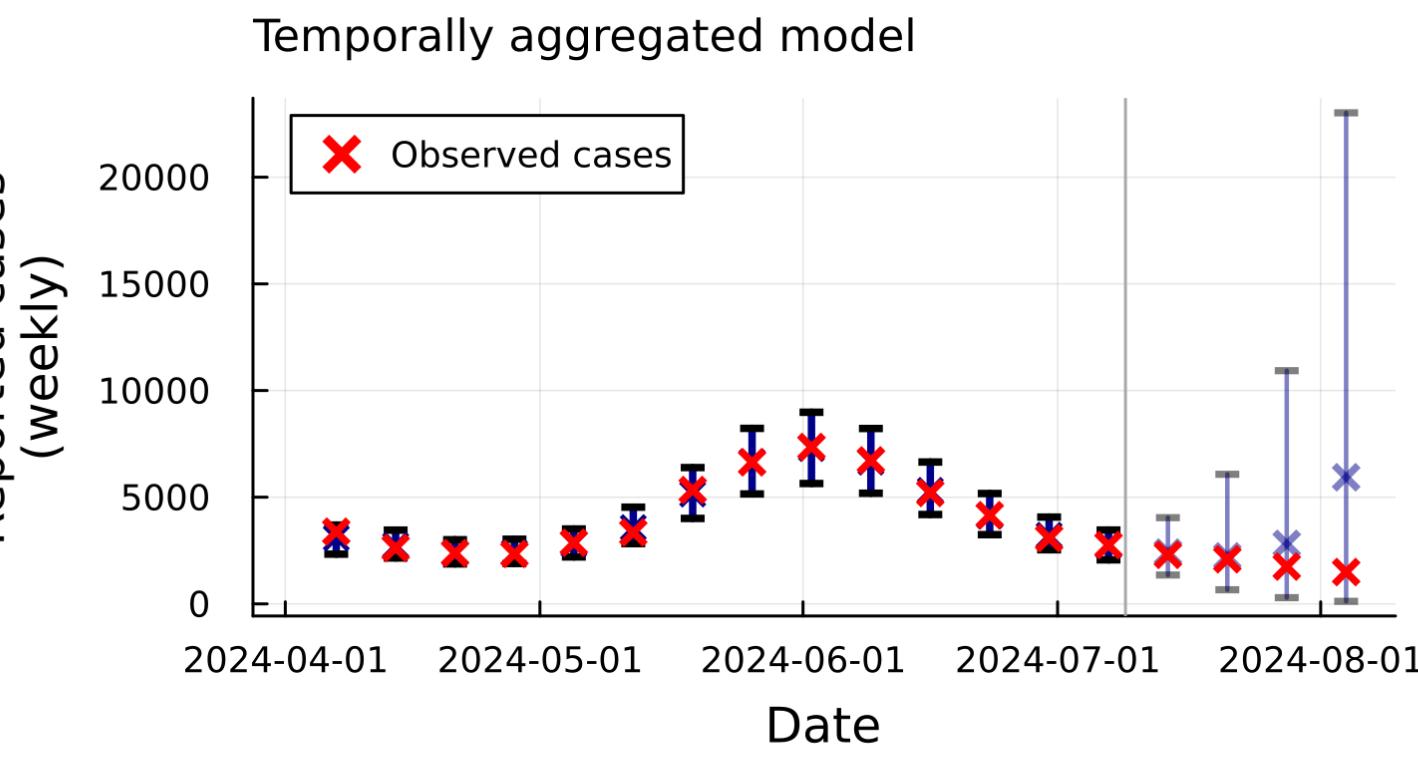
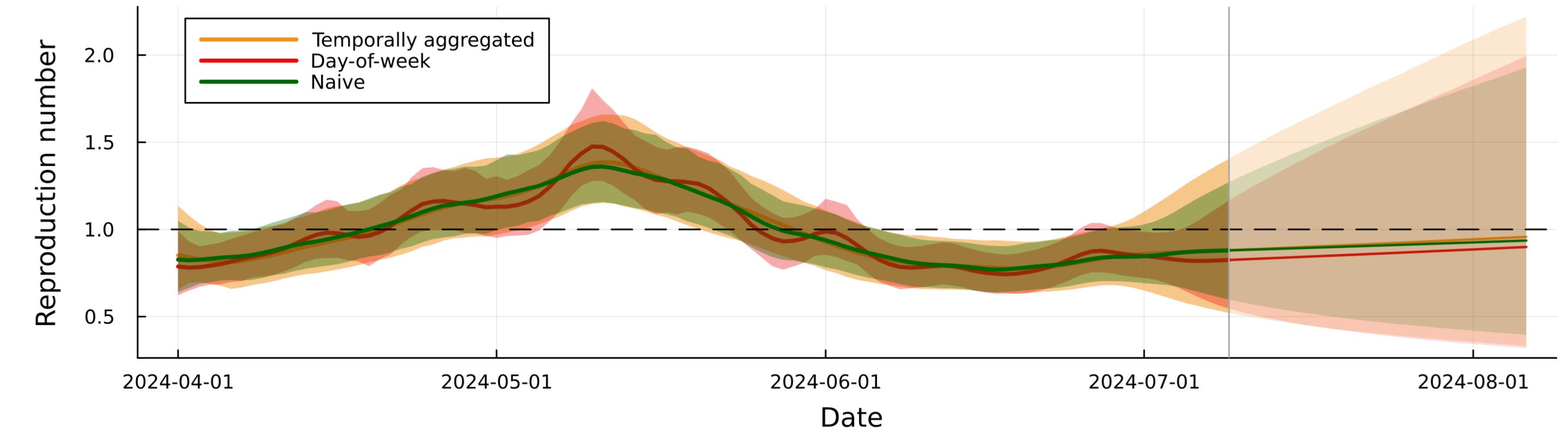
$$\hat{\ell}(\theta) = \sum_{t=1}^T \log \bar{W}_t$$

- Then just plug this into a standard Metropolis Hastings algorithm!
- Need to be careful about  $S.D.(\hat{\ell}(\theta))$ ....

# Sequential Monte Carlo: Overall approach



# Example: Temporal effects



## Three observation models:

- Naive
- Day-of-week effect
- Weekly aggregated

# Some final thoughts...

## Advantages of these methods and models

- ✓ Simple, intuitive, highly flexible
- ✓ Requires no external software
- ✓ No complicated mathematical approximations
- ✓ Produces well-calibrated estimates and predictions

## Disadvantages

- ✗ Far less established than existing methods in epi
- ✗ Can struggle with high-dimensional  $\theta$
- ✗ Sequential nature is still somewhat restrictive

Website



Preprint



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