

# Applications of Nowcasting Methods to Notifiable Disease Surveillance

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# Our team

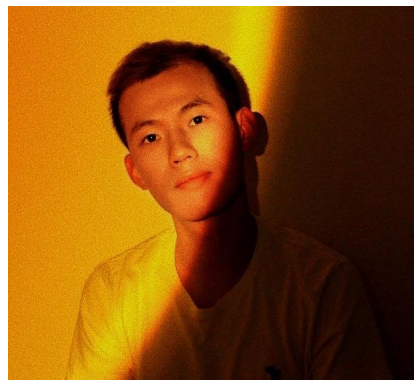
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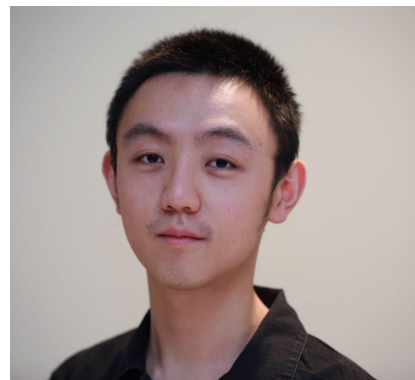
**Ian Painter**



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**Fred Yu**

# Scientific Background

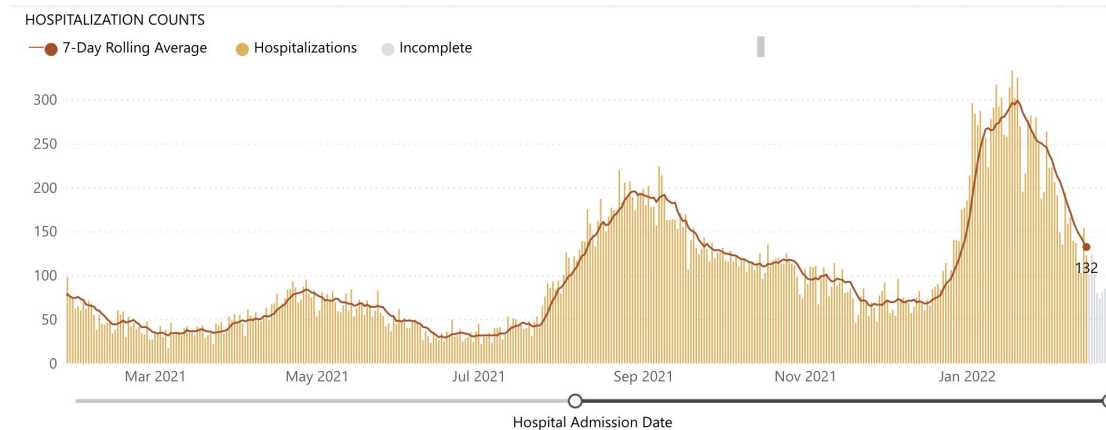


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# Problem overview

- > WA State DOH concerned with Covid-19 hospitalizations
  - Track by date of disease transmission to monitor disease spread
  - Positive test specimen collection date equated to disease transmission date for simplicity



# Delay of interest

- > Only concerned with cases that result in hospitalization
- > Variable delay between test collection and hospitalization (1-14 days)



- > Examples
  - Subject 1: Positive test on Day 1 → Hospital report on Day 10 → Delay = 9
  - Subject 2: Positive test on Day 9 → Hospital report on Day 11 → Delay = 2

# Objectives

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- > Understand the true nature of the data and delays
- > Develop nowcasting models to mitigate impact of delays
- > Evaluate the performance and robustness of our models, and determine the highest-performing option
- > Establish the contexts under which our model may be used by the DOH

# Nowcasting Background

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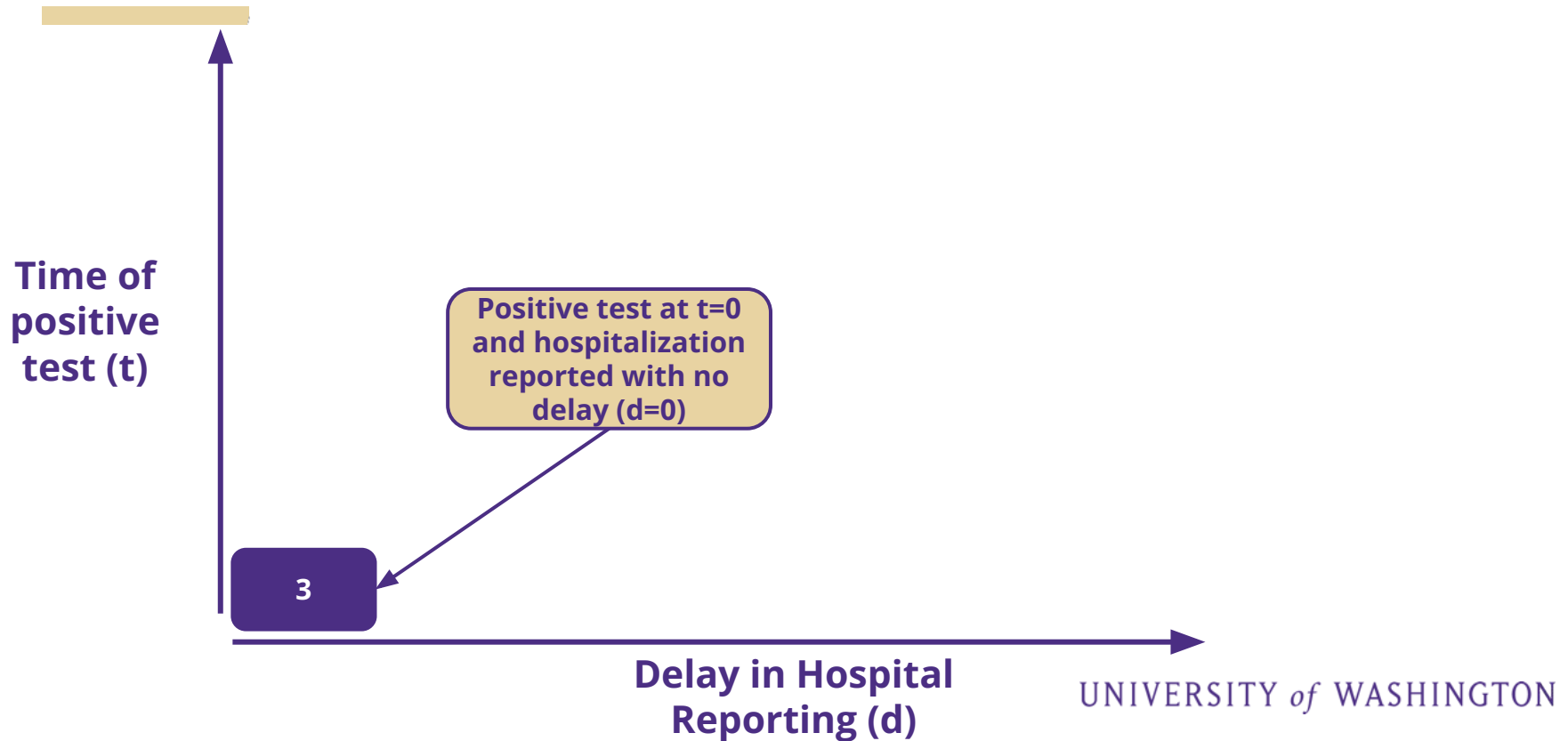
# Nowcasting - Predicting the present

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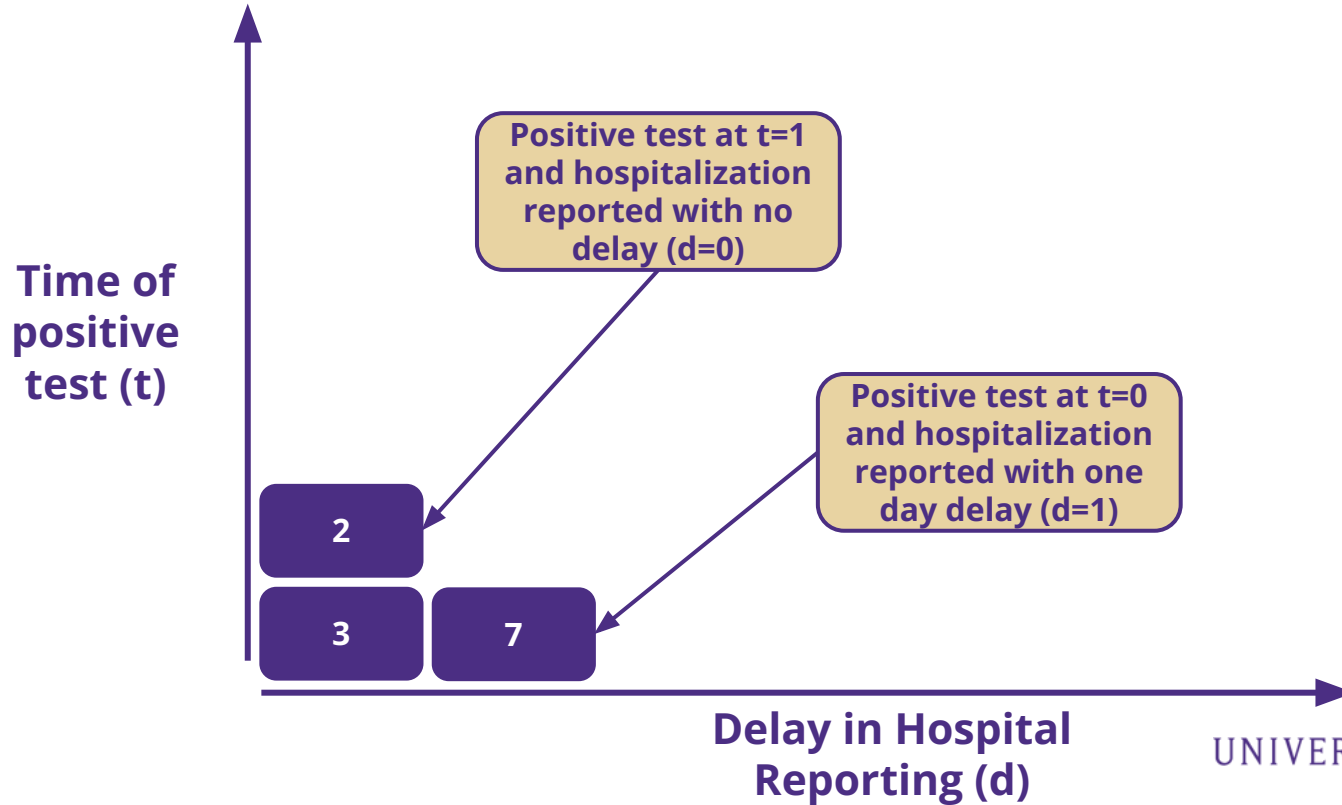
- > Estimate the **occurred-but-not-yet reported** events
  - e.g. How many hospitalizations will result from the positive tests collected yesterday?
- > Originated in actuarial literatures, found application when analysing AIDS/HIV epidemic
- > Modeling the **delay distribution** is a common feature



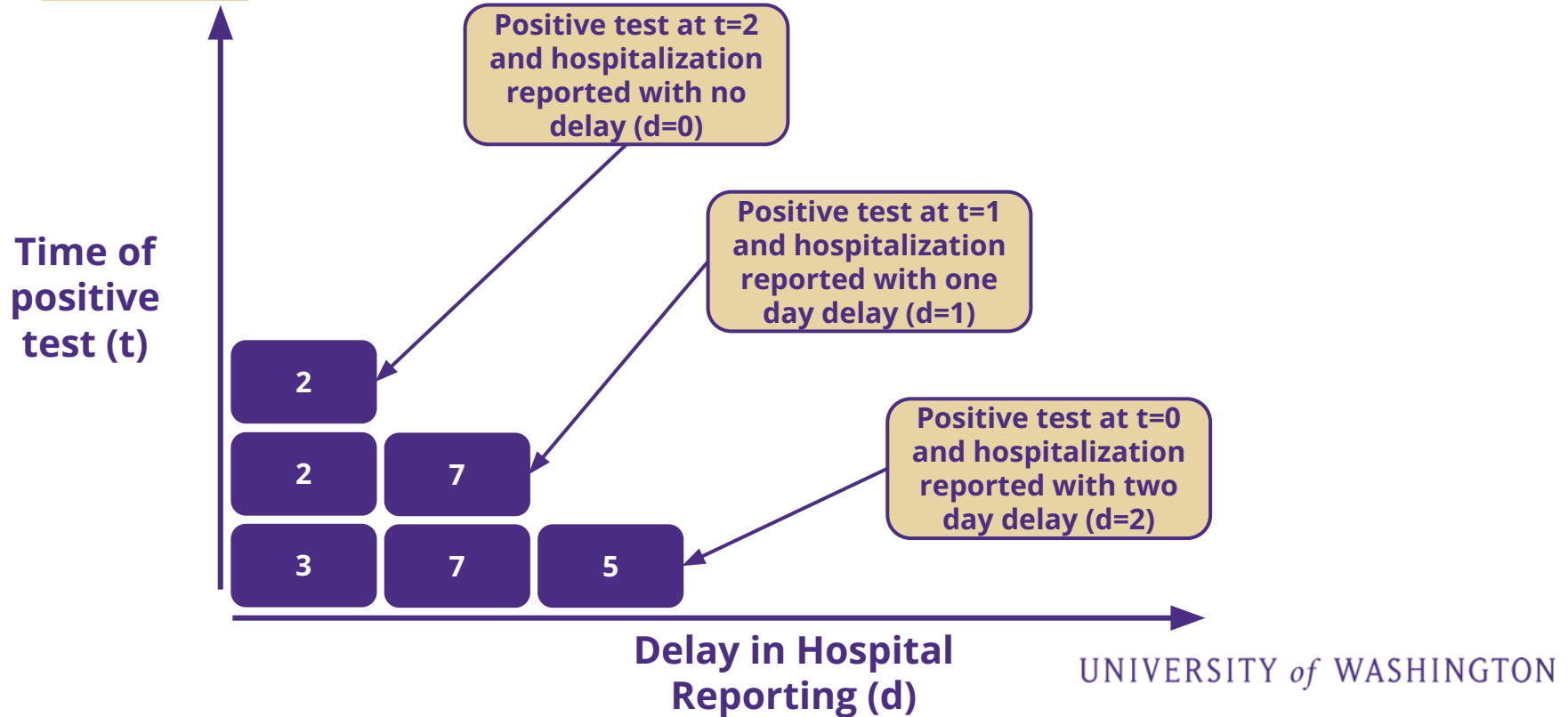
# Nowcasting example - Day 1



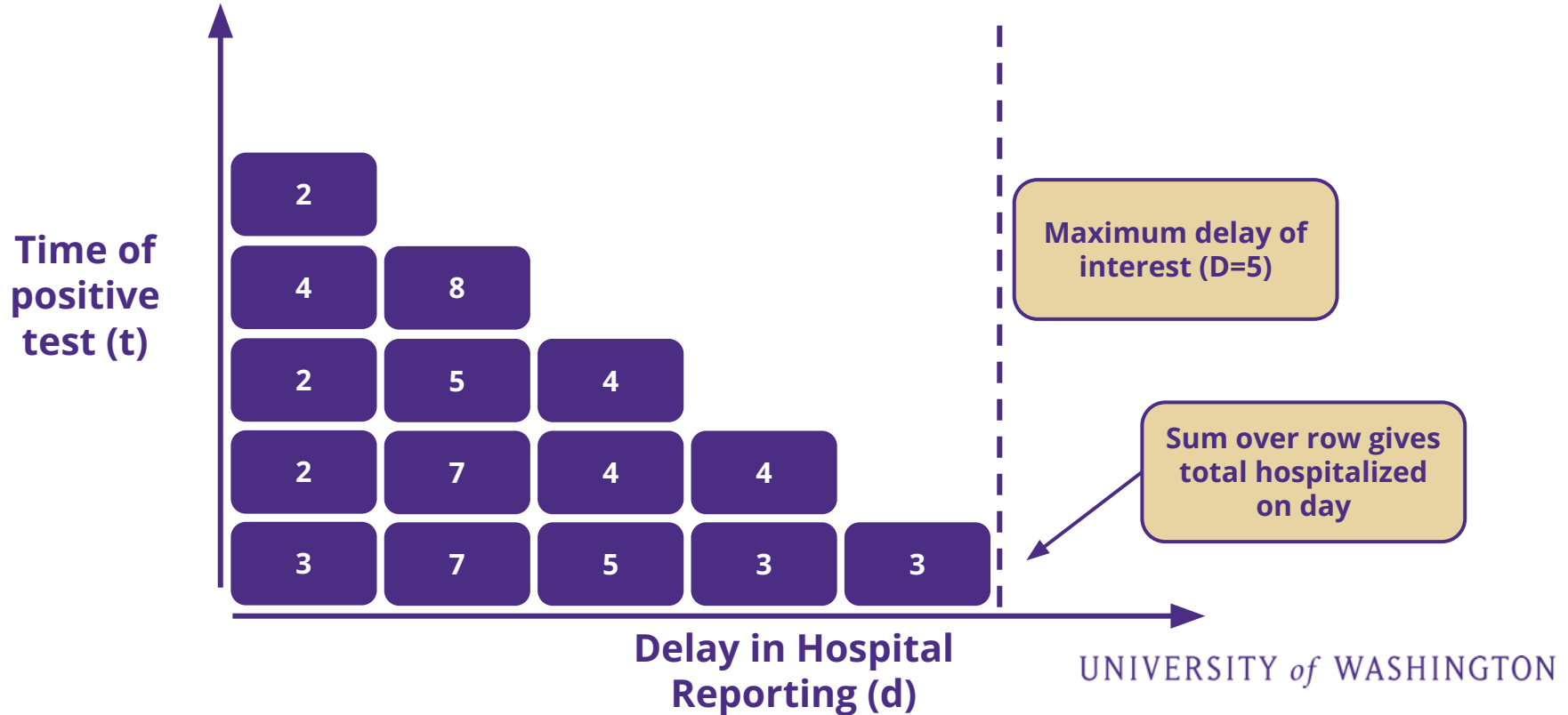
# Nowcasting example - Day 2



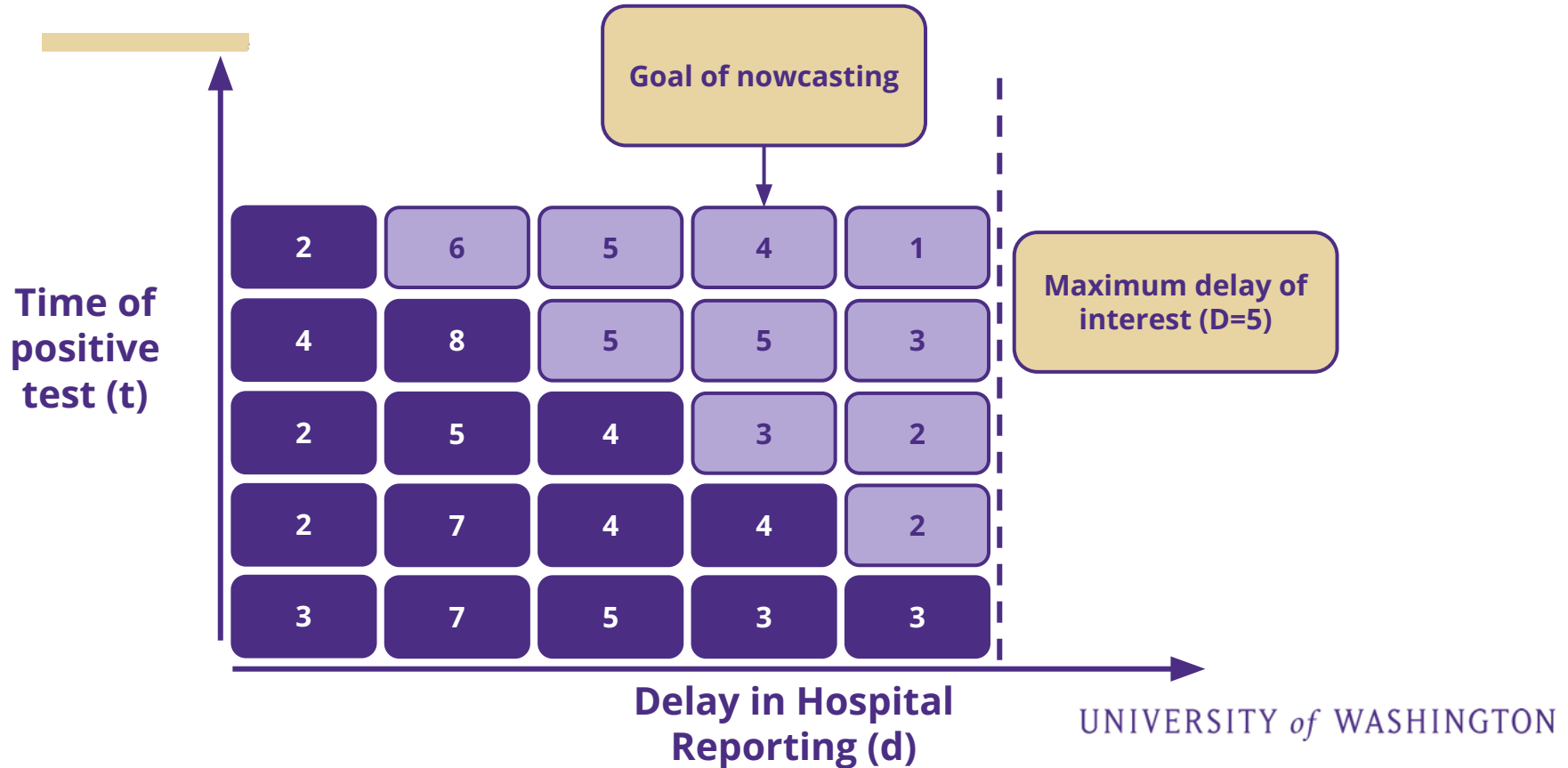
# Nowcasting example - Day 3



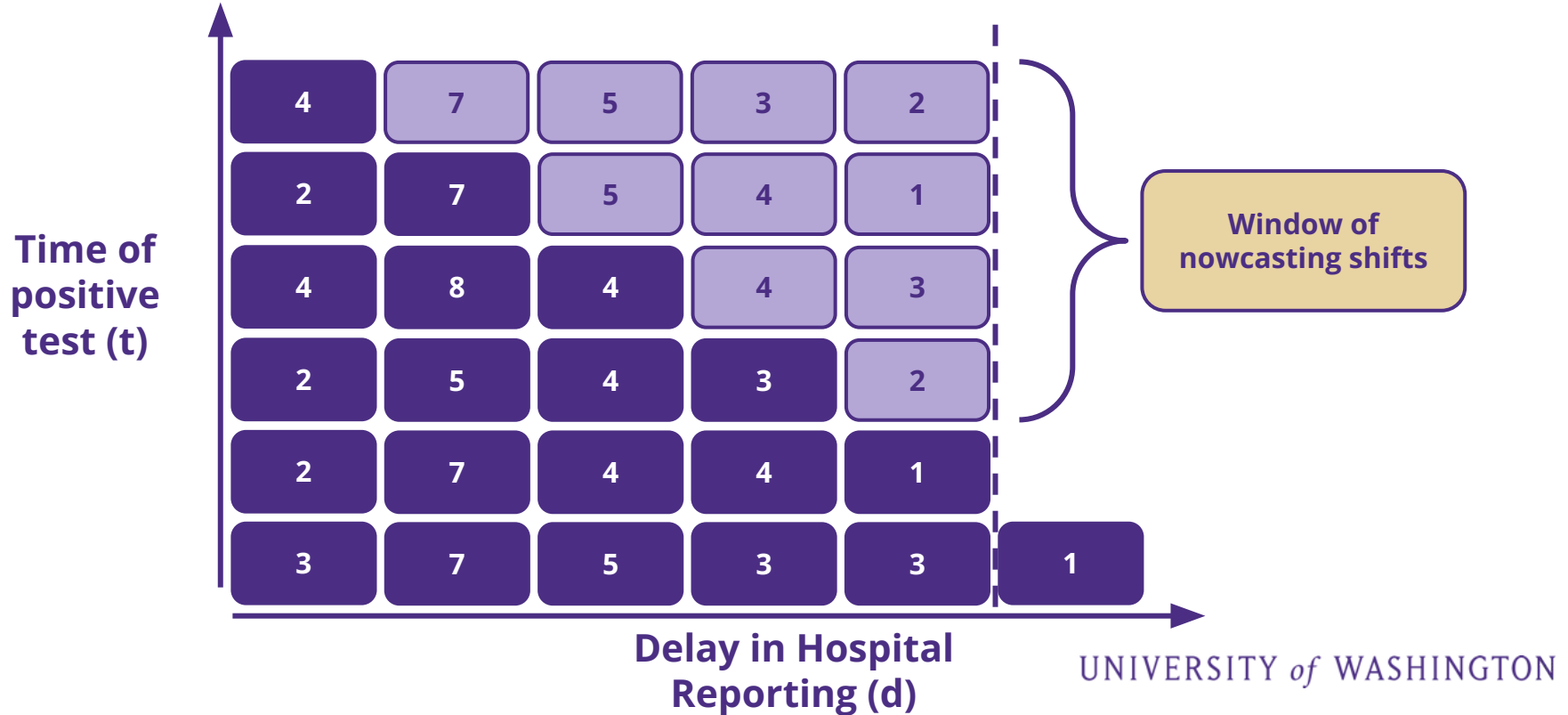
# Nowcasting example - Day 5



# Nowcasting example - Day 5



# Nowcasting example - Day 6



# Nowcasting Methods



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# General nowcasting framework

Assumption:  $n(t, d)$ , i.e. the case counts at day  $t$ , reported with  $d$  days of delay, follows an underlying in-homogeneous poisson process with the mean

The diagram illustrates the components of the mean  $\mu_{t,d}$ . It shows two yellow boxes: 'Epidemic curve' on the left and 'Delay distribution' on the right. Arrows point from each box to its respective term in the equation  $\mu_{t,d} = \lambda_t \times p_{t,d}$ . The epidemic curve is represented by  $\lambda_t$  and the delay distribution by  $p_{t,d}$ , which are multiplied together to give the mean  $\mu_{t,d}$ .

$$\mu_{t,d} = \lambda_t \times p_{t,d}$$

Nowcasting can thus be done through the following steps:

**Step 1: Estimate the rate of occurrence of cases at day  $t$ , i.e. the epidemic curve**

**Step 2: Estimate the proportions of cases at day  $t$ , delayed for  $d$  days, i.e. the delay distribution**

**Step 3: Predict the unobserved  $n(t, d)$ 's in order to compute the total  $N(t, T+D)$**



# Bayesian nowcasting models

Model	Epidemic curve	Delay distribution
<b>HH</b> ( <i>Hohle &amp; Heiden</i> )	Assume a gamma mean $\lambda_t \sim \text{Gamma}(a_\lambda, b_\lambda)$  Then the true count given mean is Poisson.	<ul style="list-style-type: none"><li>- Assume <b>time-invariant</b> delay</li><li>- Model using GD-Multinomial conjugacy framework</li></ul>
<b>NobBS</b> ( <b>Nowcasting by Bayesian Smoothing</b> )	Model as a first-order random walk	<ul style="list-style-type: none"><li>- Assume <b>time-invariant</b> delay</li><li>- Model using GD-Multinomial conjugacy framework</li></ul>

# Results

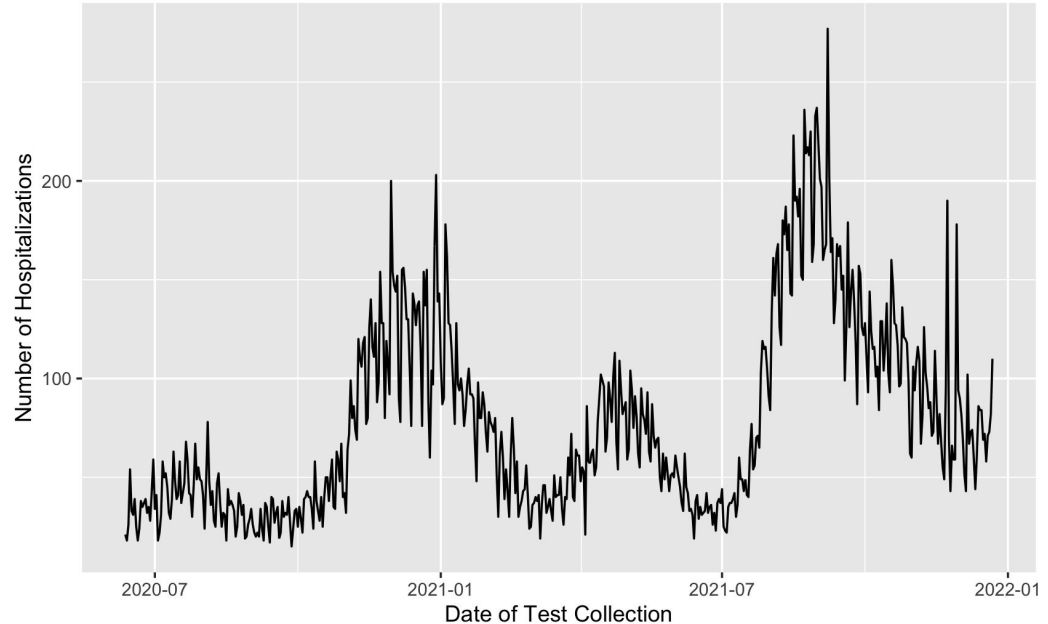


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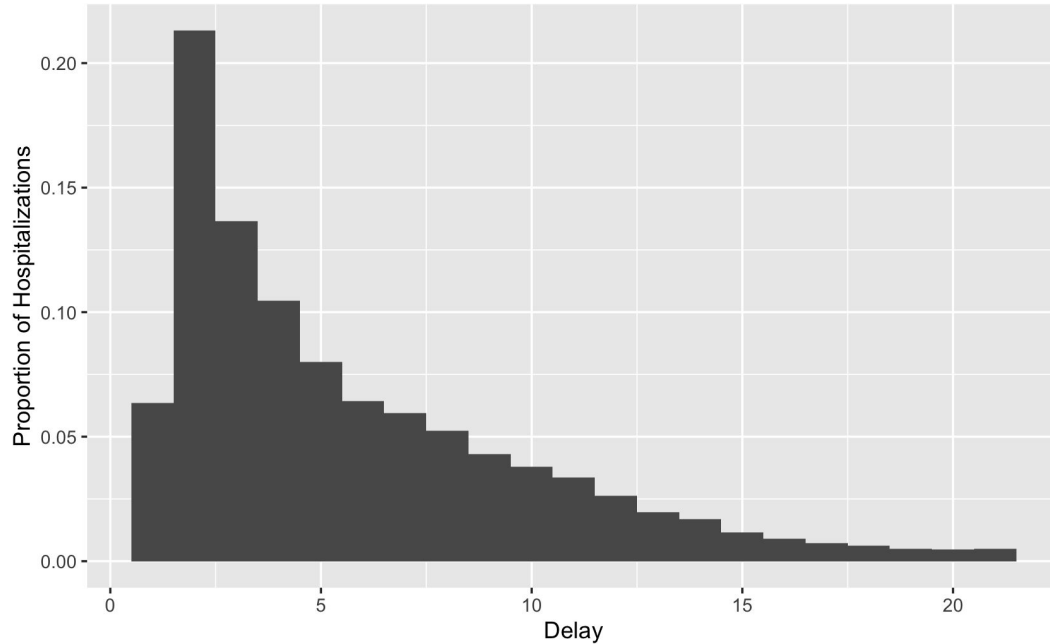


# Data structure and exploratory analysis

- > Schema: One row per hospitalization
- > Time of interest: Test collection to hospitalization reporting
- > Date range of data:  
6/11/20 to 1/12/22
- > Max delay of interest: **21 days**
  - Analysis start: 7/2/20
  - Analysis end: 12/21/21

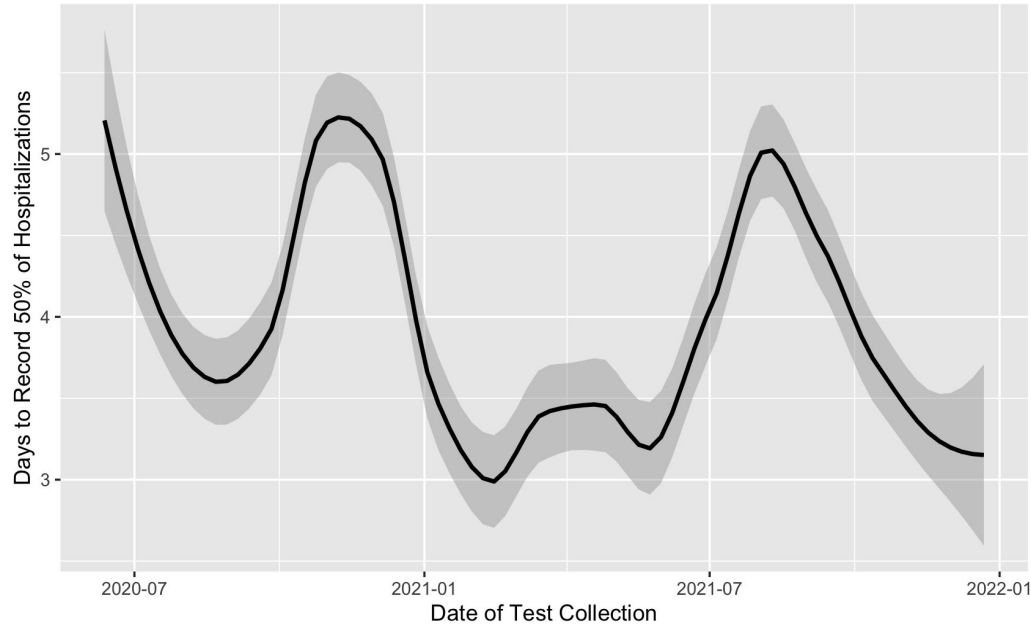


# Exploratory analysis



- > No hospitalizations confirmed on date of test (delay = 0)
- > Plurality of cases (20%+) reported with two days delay

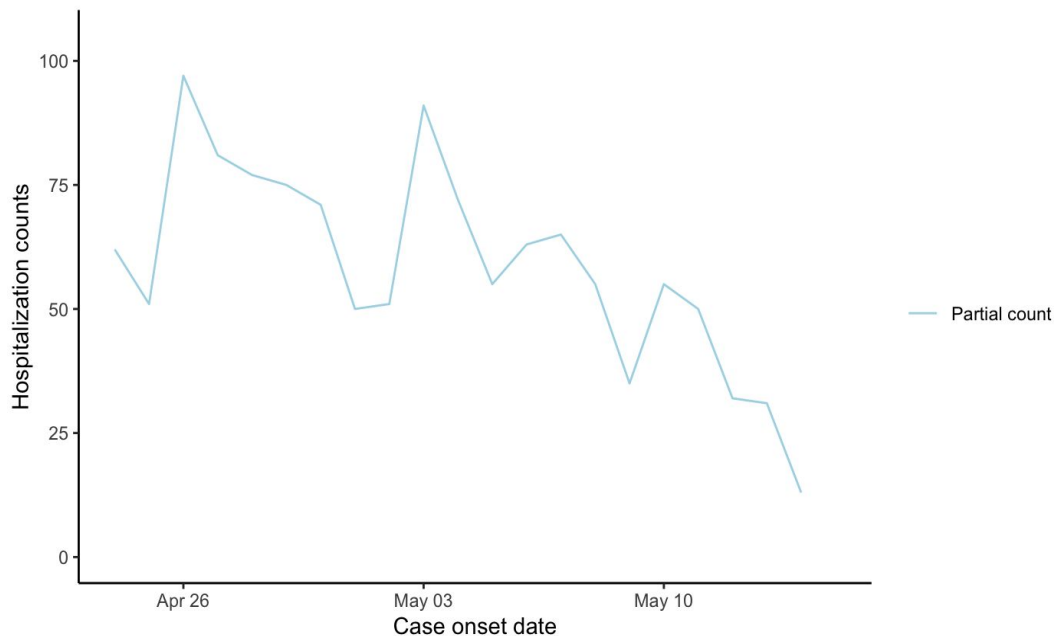
# Exploratory analysis



- > Days needed to observe 50% of total hospitalizations fluctuates over pandemic
- > Times associated with higher volumes require more days to reach 50% of hospitalizations

# Example nowcast at May 15, 2021

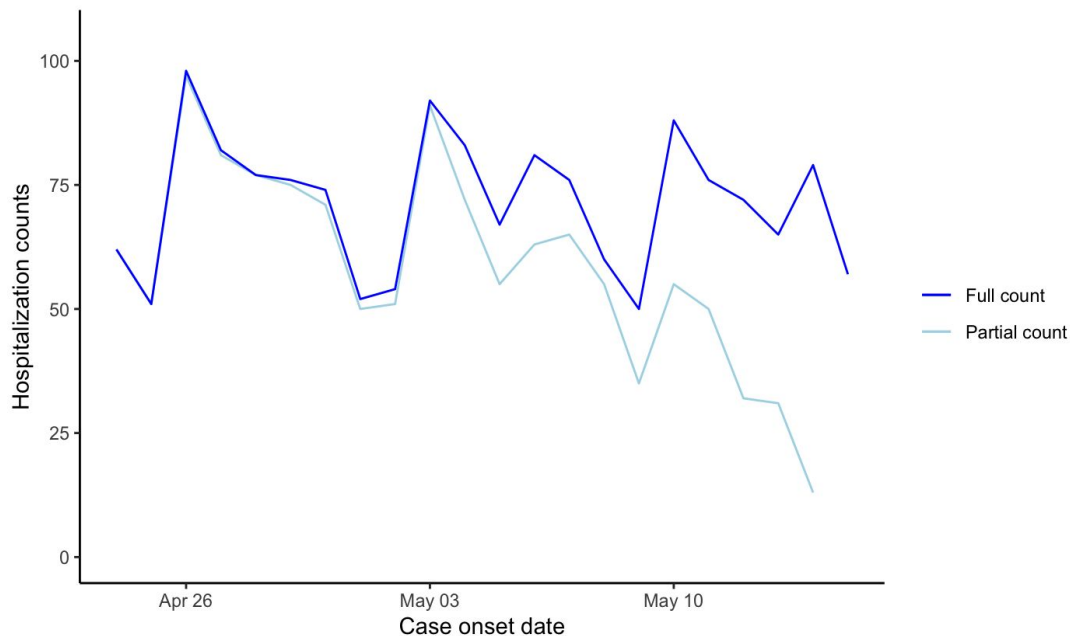
Nowcasting estimates at 2021-05-15 (T) and 21 days prior



- > Information available to us at time of nowcasting
- > Decreases in most recent days

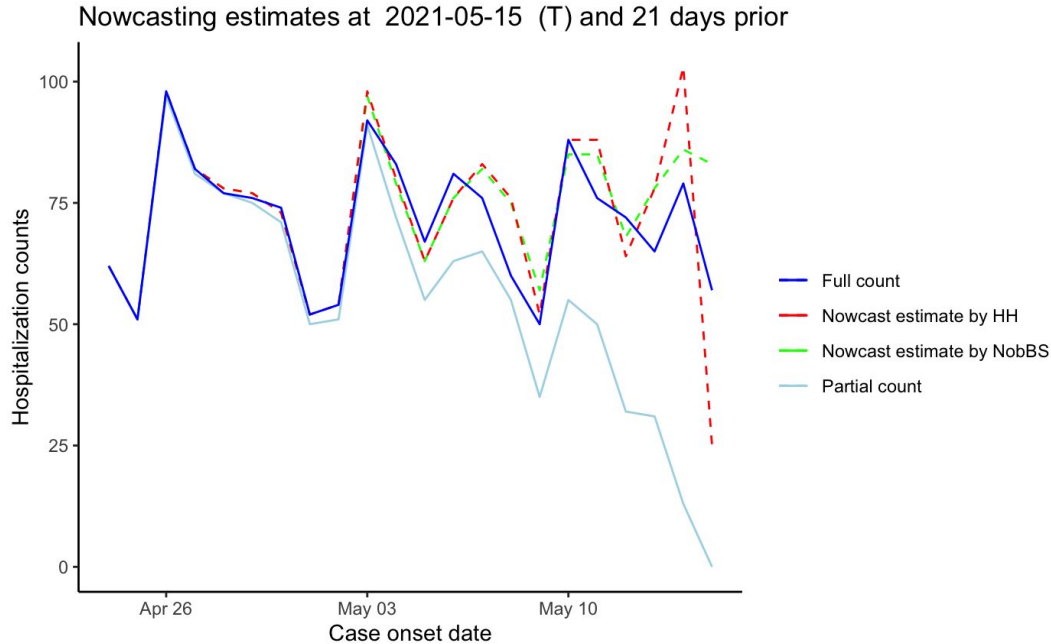
# Example nowcast at May 15, 2021

Nowcasting estimates at 2021-05-15 (T) and 21 days prior



- > Lookback time: Days between date of nowcasting (e.g. 5/15) and date of prediction (ranges from 0 to 21)
- > Gap between partial and full counts narrows as lookback time increases

# Example nowcast at May 15, 2021



- > Accuracy increases with increasing lookback time
- > Direction of error (overpredict v underpredict) varies by date

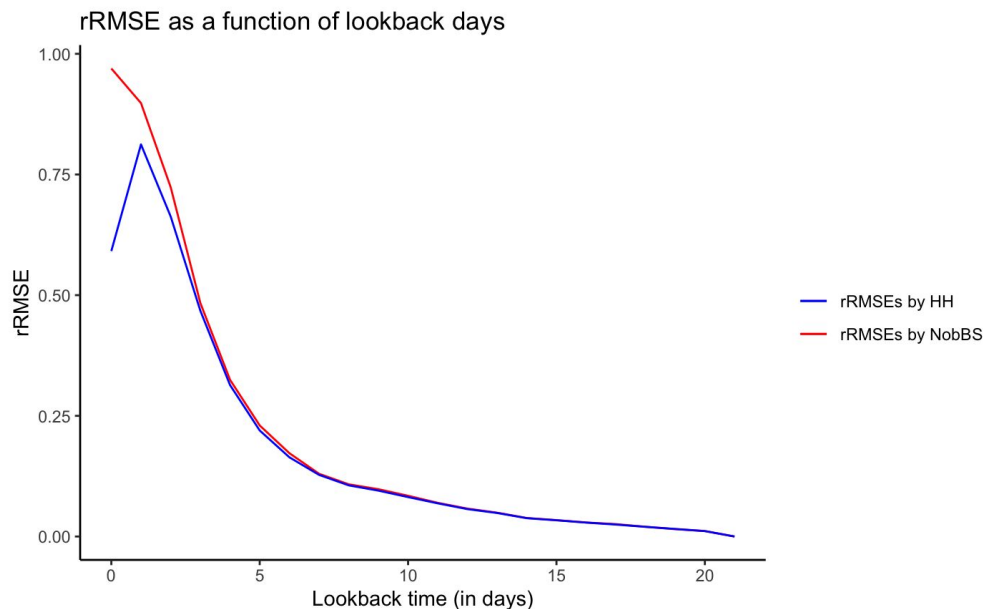


# Evaluation of nowcasts

- Given the max delay, **22** estimates generated for each date of nowcasting
- We use *relative root-mean-square error* (***rRMSE***) as the metric to evaluate the quality of different nowcasting models
  - Think of this as average relative error with penalty for large deviations
- The medians of *rRMSE* for all **539** times of nowcast

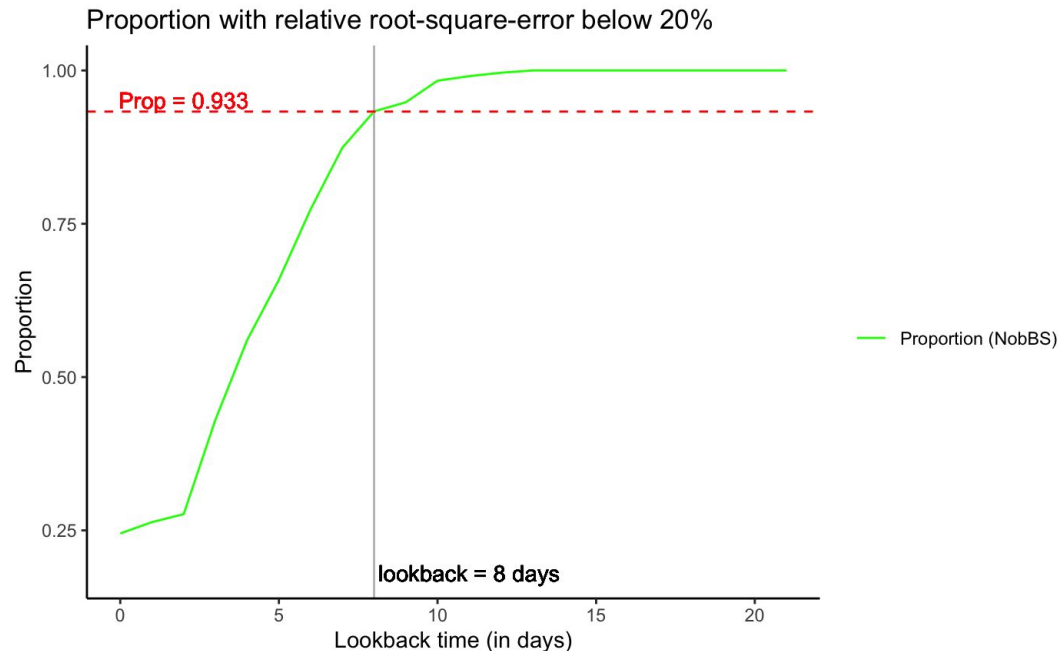
	HH	NobBS
rRMSE Median	0.261	0.197
rRMSE Median (lookback >= 5)	0.089	0.092

# rRMSE by lookback time (in days)



- > Both models see decrease in rRMSE as lookback time increases
- > HH method appears more performant at shorter lookback times

# Nowcasts with errors below threshold



- > Collaborated with DOH to identify **20%** as tolerable error, as daily fluctuations reach this level
- > For **lookback times of 8+**, we have high certainty of estimating within 20%
- > Current DOH practice excludes most recent 16 days; **NobBS** model expedites decision-making by **8 days**

# Next steps

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- > Handoff to WA State Department of Health
  - Determine parameters of use
  - Handoff code and guidelines
  - Productionize model and monitor
- > Improve model
  - Incorporate reporting hospital information
  - Perform HH method with epidemic curve modeled by spline

# Acknowledgement

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**Minh Vo**

**And our Capstone program cohort!**

# Questions?



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# Appendix

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