



BAYESIAN CHANGE-POINT DETECTION FOR PROCESS MONITORING WITH FAULT DETECTION

KERN CENTER FOR THE SCIENCE OF HEALTH CARE DELIVERY

Alexander C. Murph

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CONTROL TOWER INTERFACE

The screenshot displays the Mayo Clinic Control Tower Interface. At the top, the Mayo Clinic logo is on the left, and navigation links for Rochester, Saint Marys, Methodist, Arizona, Florida, and Mayo Health Systems are in the center. On the right, there's a search bar labeled 'find a patient', a notification bell with 374 alerts, and a user profile for Ann Haun. Below the navigation, there are 'List View' and 'Geo View' buttons. A sorting dropdown is set to 'Palliative Score', and pagination shows page 1 of 89. A 'Demo Mode' button is also present.

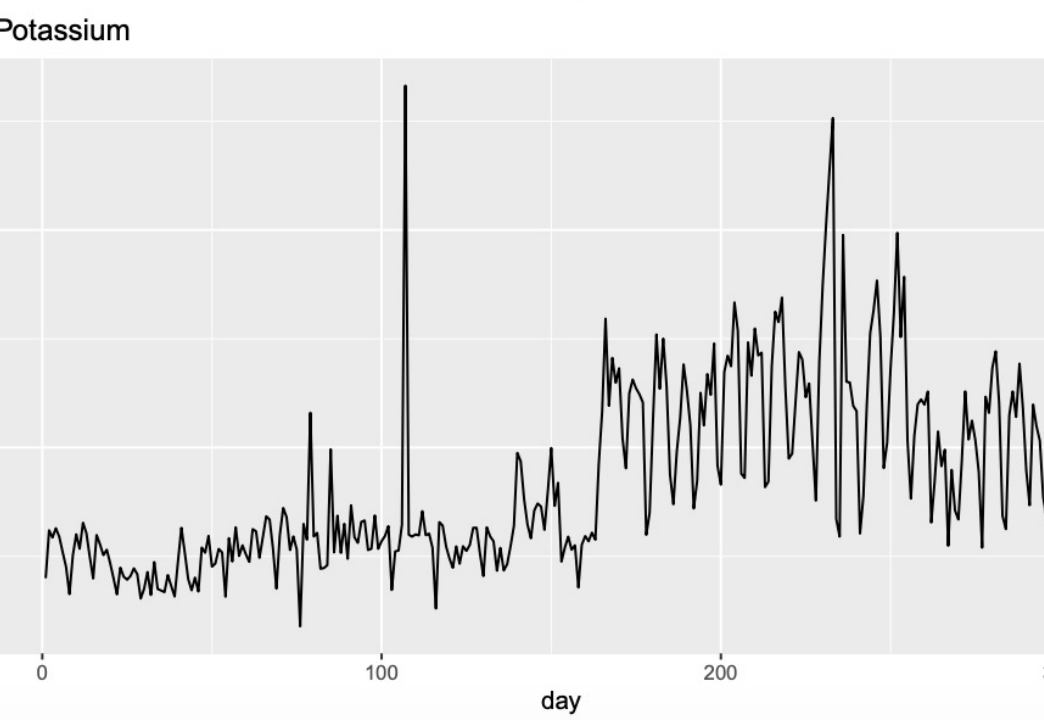
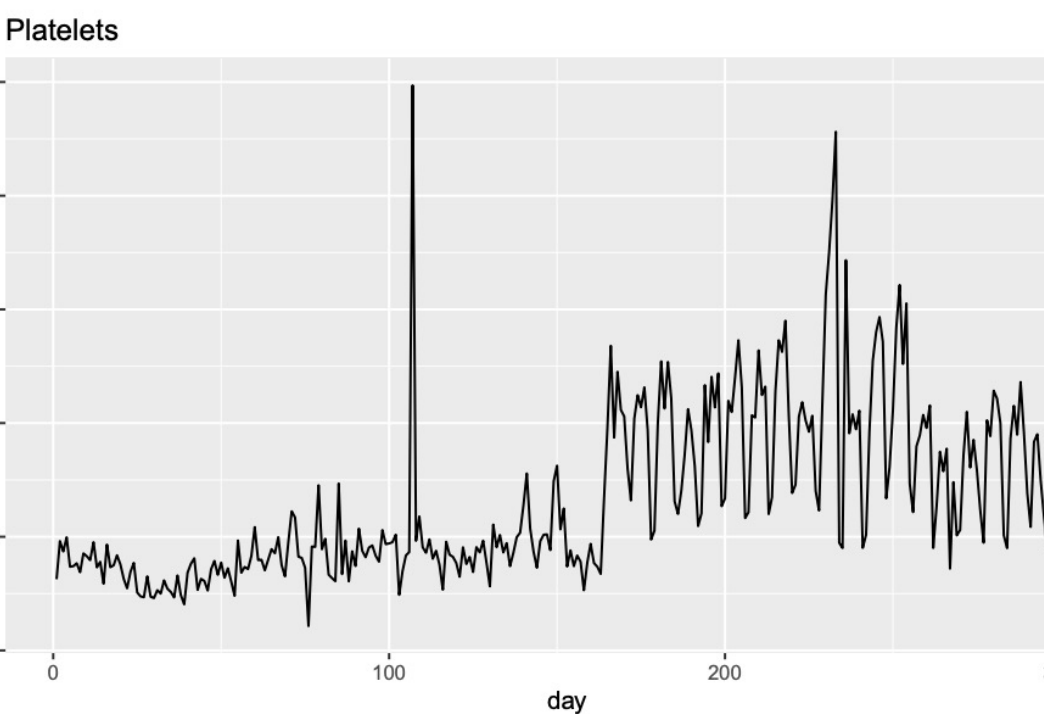
The main content area lists three patients, each with a 'Patient, Test' summary, a 'Problems List', and '24-hour Events'. Each patient entry includes a palliative score in a colored circle (red for 100, green for 99) and a bell icon with a number in a circle (2, 1, 2 respectively).

Patient, Test	Problems List	24-hour Events
Patient, Test 0-000-000 58 years Male 1 day in the hospital Facility: ROSMC Dept: RODO6D Room: RDO6307 307-P 100 Palliative	Problems List 44 total 1. Abdominal Pain 2. Cellulitis Groin 3. Infection Urinary Tract Acute 4. History Of Falling	24-hour Events 19 total 1. CM Care Management PROGRESS 2. DX ABDOMEN PORTABLE ANTERIOR POSTERIOR 1 VIEW 3. DX CHEST PORTABLE 1 VIEW 4. Pharmacy PROGRESS
Patient, Test 0-000-000 79 years Female 2 days in the hospital Facility: ROSMC Dept: ROAL7E Room: RAL7421 421-P 100 Palliative	Problems List 11 total 1. Hypoxia 2. Acute And Chronic Respiratory Failure With Hypoxia (HCC) 3. Chronic Respiratory Failure (HCC) 4. Acute Respiratory Failure With Hypoxia (HCC)	24-hour Events 11 total 1. Critical Care Medicine PROGRESS 2. Critical Care Medicine PROGRESS 3. Critical Care Medicine PROGRESS 4. Respiratory Therapy PROGRESS
Patient, Test 0-000-000 22 years Male 3 days in the hospital Facility: ROSMC Dept: RODO2D Room: RDO2308 308-P 99 Palliative	Problems List 34 total 1. Compression Of Brain (HCC) 2. Hemorrhage Subarachnoid Personal History 3. Encephalopathy 4. Diabetes Mellitus Drug Or Chemical Induced With Hyperglycemia (HCC)	24-hour Events 11 total 1. Oncology PROGRESS 2. Neurology PROGRESS 3. Nursing Services CARE PLAN 4. GIM Integrative Medicine and Health PROGRESS

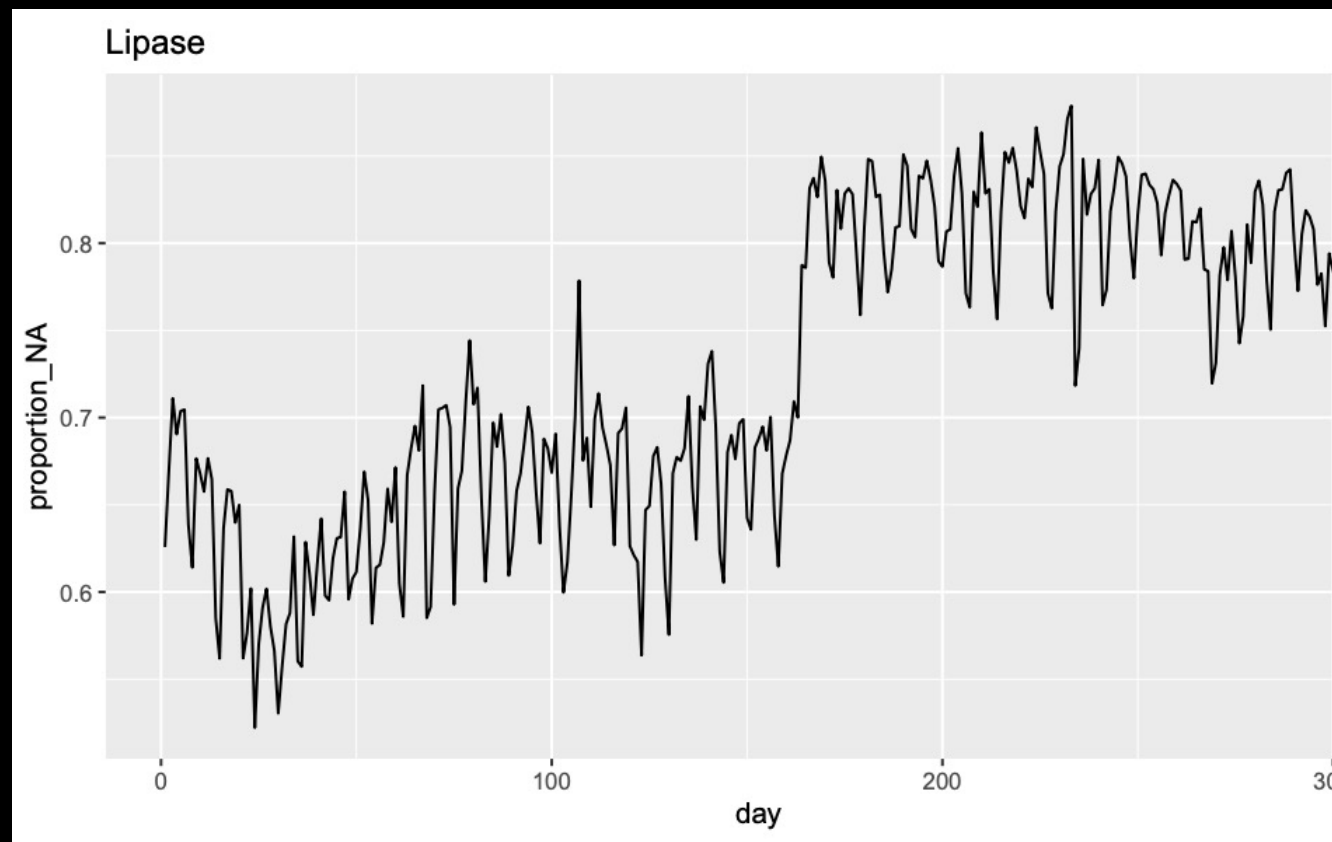
Number in the circle:

“The probability of getting an inpatient palliative care consultation in the next 7 days.”

Thank you to Patrick Wilson for sharing this slide!



WELL, WHAT COULD GO WRONG?



A (NOT SO) ELEGANT PROBLEM

(Sub) problems:

- Missing Data
- Mixed Data (discrete, binary, continuous)
- Censored Values
- HUGE data (n in millions, $p \approx 250$)

(Sub-Sub) problems:

- Lack of information for some priors
- Learning parameters for a GGM (especially the conditional independence graph structure)

(Sub) solutions:

- Bayesian Latent Variables
- Gaussian Graphical Models (GGMs)

(Sub-Sub) solutions:

- Bayesian Hierarchical Model
- Double Reversible Jump Metropolis Hastings, Conjugate priors.

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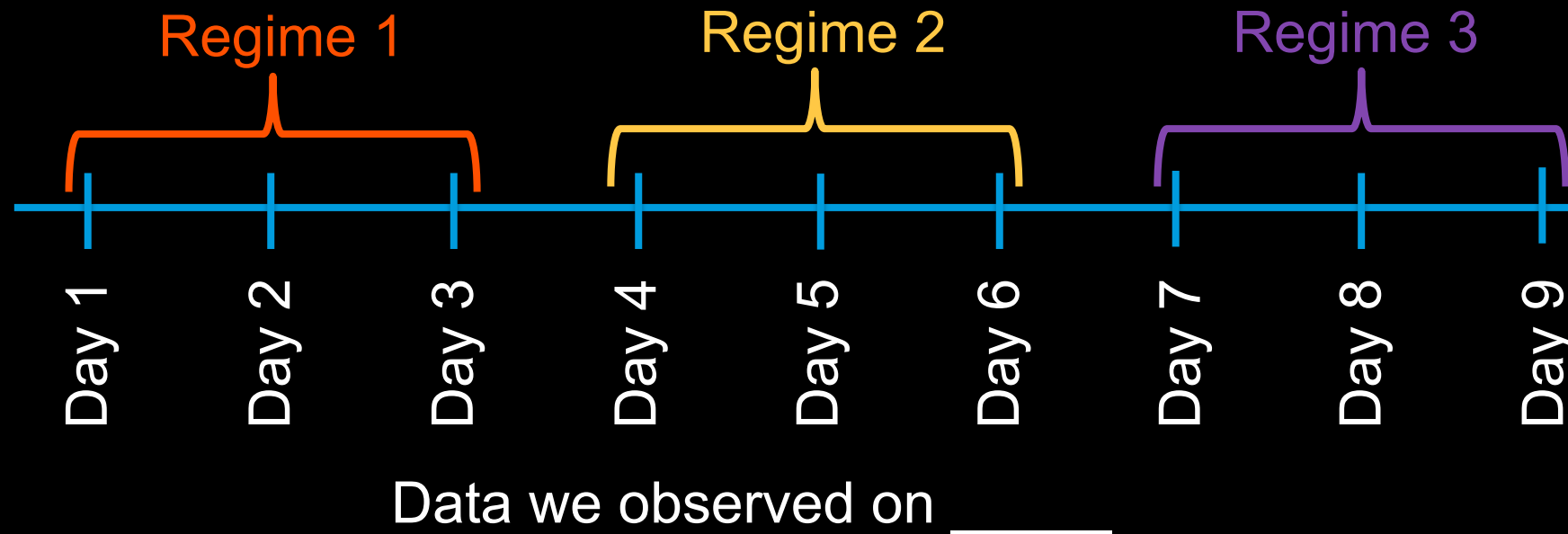
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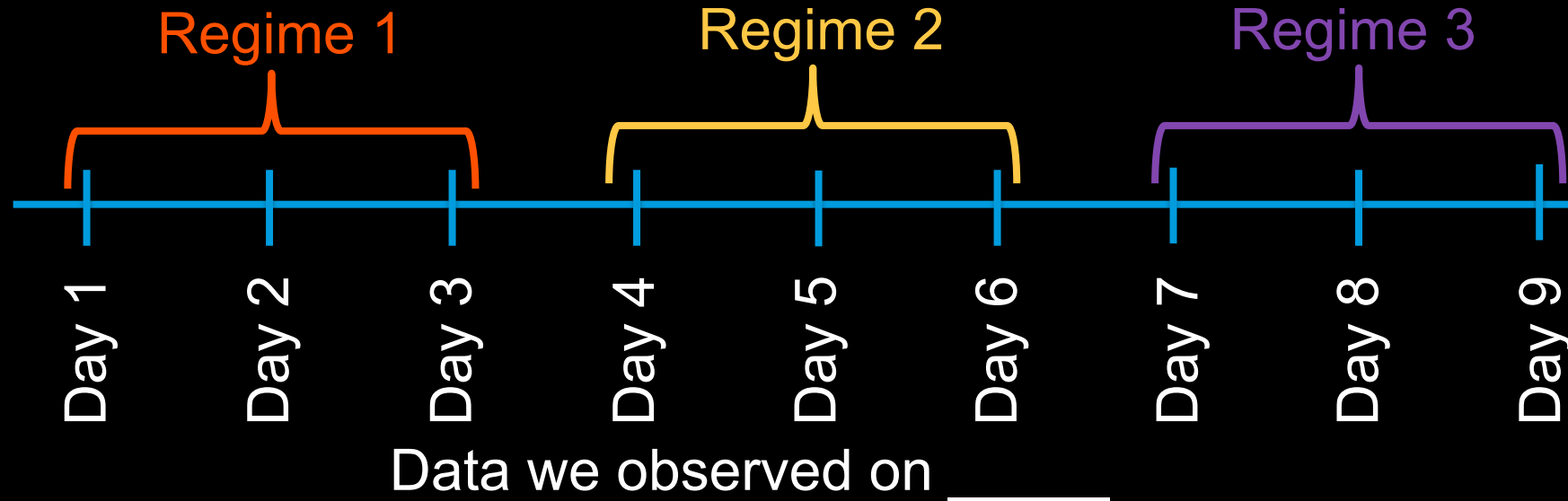
HOW DO WE FIND A CHANGE POINT?

LET'S CLASSIFY OBSERVATIONS OVER TIME INTO "REGIMES"



SUPPOSE WE OBSERVED 9 DAYS OF DATA...

We can *encode* each day's regime assignment as a vector of length 9:

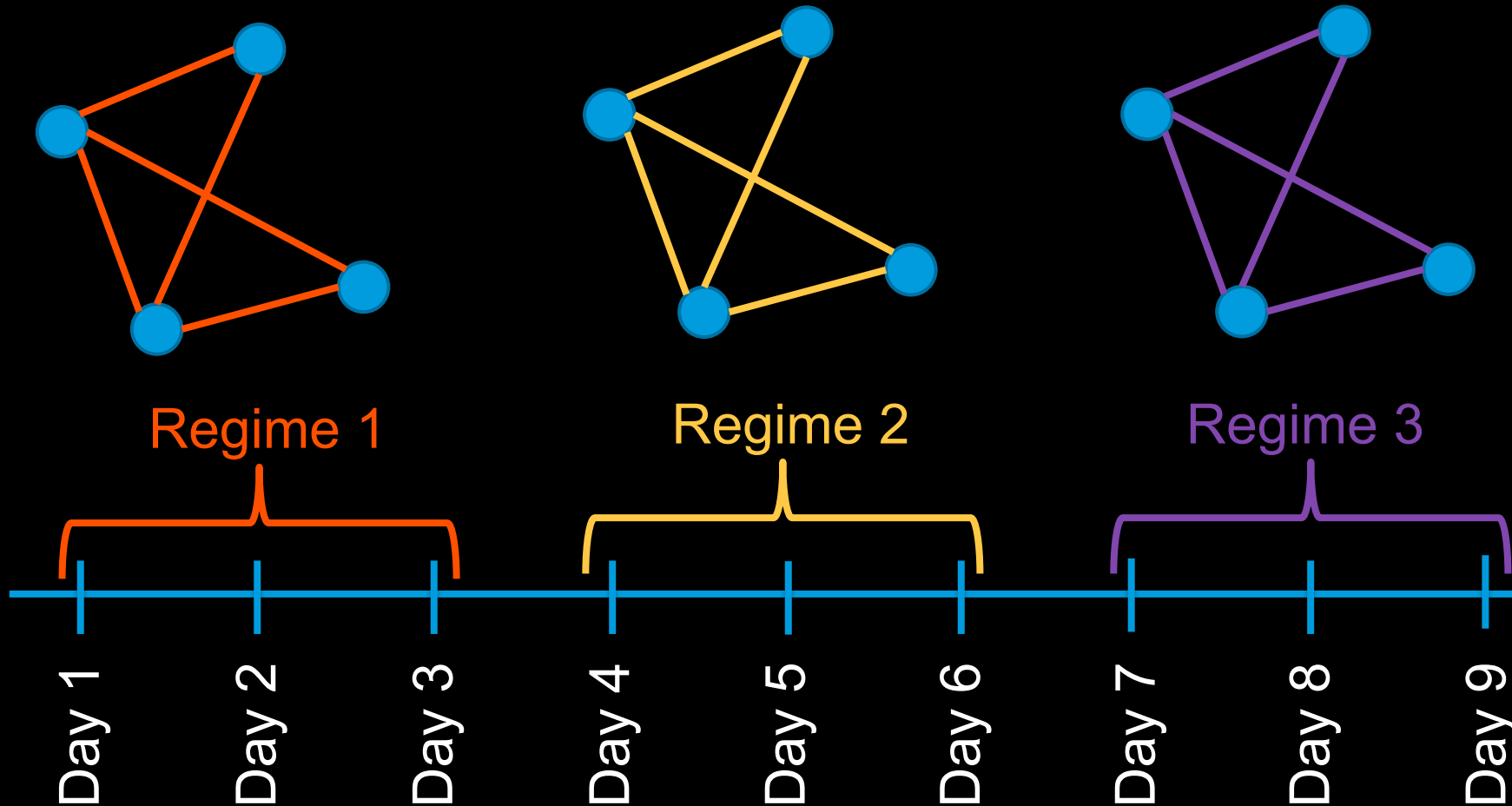


becomes

$$\phi = (1, 1, 1, 2, 2, 2, 3, 3, 3)$$

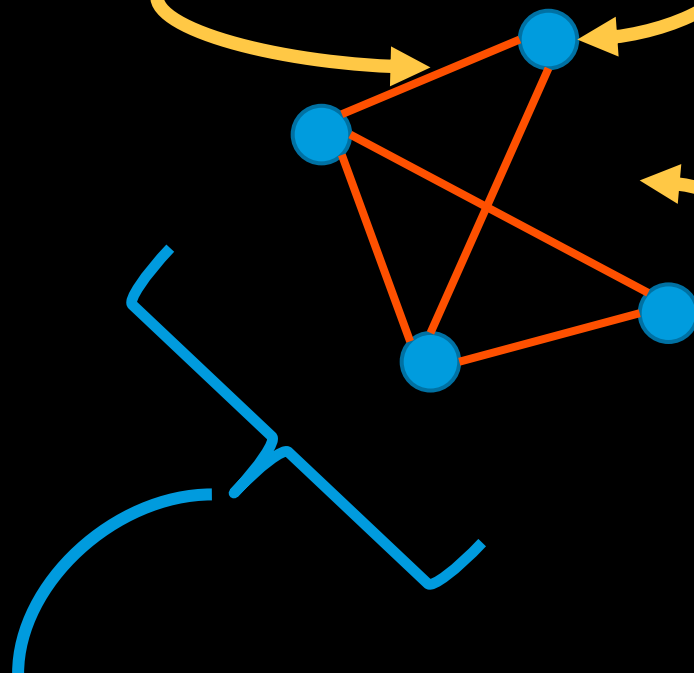
WHAT'S THE LIKELIHOOD?

WE MUST DEFINE THE DISTRIBUTION OF THE DATA *GIVEN* A REGIME VECTOR



Weighted edges
correspond to values in
the *precision matrix* of
a multivariate normal
model.

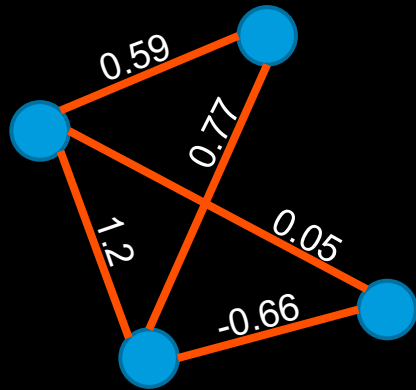
The nodes correspond to the
variables in our dataset.



The *absence* of an edge
corresponds to a zero in
the precision matrix

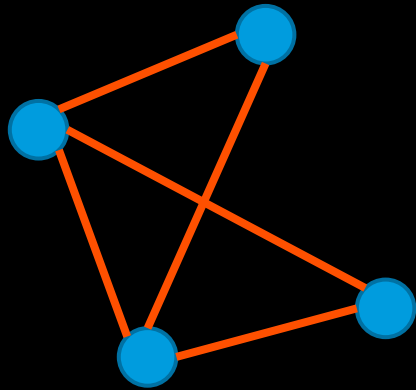
Λ , where $Y \sim \mathcal{N}_4(\mu, \Lambda^{-1})$

FOR INSTANCE,



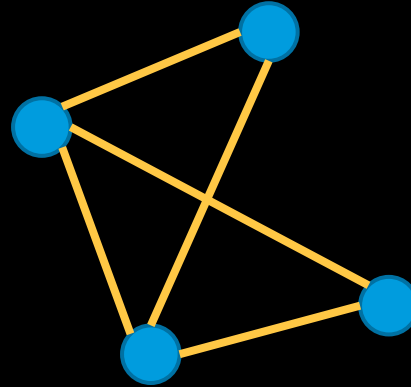
$$\Lambda = \begin{bmatrix} 1 & 0.59 & 0.05 & 1.2 \\ 0.59 & 1 & 0 & 0.77 \\ 0.05 & 0 & 1 & -0.66 \\ 1.2 & 0.77 & -0.66 & 1 \end{bmatrix}$$

$$\sim \mathcal{N}(\mu_1, \Lambda^{-1}_1)$$



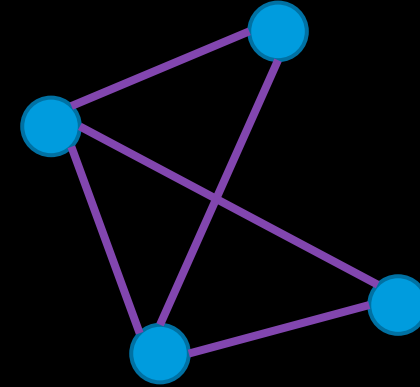
Regime 1

$$\sim \mathcal{N}(\mu_2, \Lambda^{-1}_2)$$

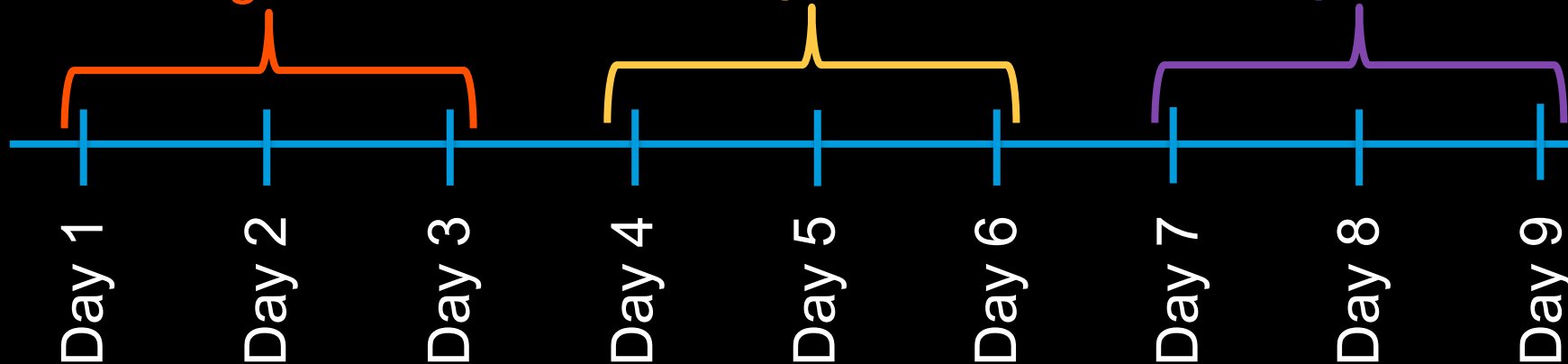


Regime 2

$$\sim \mathcal{N}(\mu_3, \Lambda^{-1}_3)$$



Regime 3



$$\phi = (1, 1, 1, 2, 2, 2, 3, 3, 3)$$

INFERENCE ON *THREE* PARAMETERS OF INTEREST

- The set-up on the previous slides admits a likelihood for our data (a product of normal mixture models).
- To get good posterior samples on ϕ , μ , Λ , G , we require:
 - Priors on each parameter;
 - Sampling procedures for proposal values.

SPARSITY ENCODED USING THE G-WISHART


Prior Distributions on the Parameters

$$\Lambda \sim \text{G-Wishart}(\delta, I)$$

(δ , degrees of freedom)

$$\mu|\Lambda \sim \mathcal{N}(\mu_0, \Lambda^{-1})$$

$$(\mu, \Lambda) \sim f_{\mu|\Lambda} f_{\Lambda}$$

$$p(\Lambda := \Sigma^{-1} | G) = \frac{1}{I_G(\delta, D)} (\det \Lambda)^{(\delta-2)/2} \exp \left\{ -\frac{1}{2} \langle \Lambda, D \rangle \right\}$$


★ is a normalizing constant gotten by integrating over a space of precision matrices with fixed zero structure

TO EVALUATE

Laplace Approximation: (Dobra and Lenkowski 2011)

$$h_{\delta,D}(\Lambda) = -\frac{1}{2}[\text{tr}(\Lambda^T D) - (\delta - 2) \log(\det K)]$$

$$I_G(\hat{\delta}, D) = \exp \left\{ h_{\delta,D}(\hat{\Lambda}) \right\} (2\pi)^{|V|/2} [\det H_{\delta,D}(\hat{\Lambda})]^{-1/2}$$

(V is the set of free elements in Λ , H is the hessian of h)

When we must calculate the normalizing constant, there exists a fast approximation!

TO SAMPLE

Algorithm 1 Exact sampling from the precision matrix.

Input: A graph $G = (V, E)$ with precision matrix K and $\Sigma = K^{-1}$

Output: An exact sample from the precision matrix.

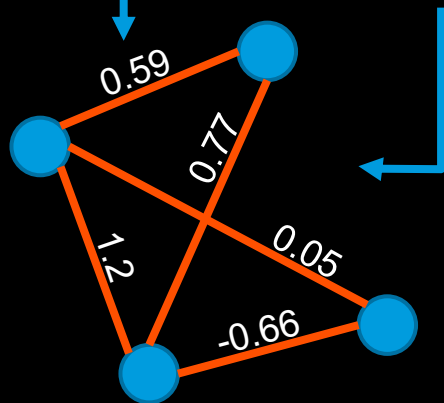
- 1: Set $\Omega = \Sigma$
- 2: **repeat**
- 3: **for** $i = 1, \dots, p$ **do**
- 4: Let $N_i \subset V$ be the neighbor set of node i in G . Form Ω_{N_i} and $\Sigma_{N_i,i}$ and solve $\hat{\beta}_i^* = \Omega_{N_i}^{-1} \Sigma_{N_i,i}$.
- 5: Form $\hat{\beta}_i \in R^{p-1}$ by padding the elements of $\hat{\beta}_i^*$ to the appropriate locations and zeros in those locations not connected to i in G .
- 6: Update $\Omega_{i,-i}$ and $\Omega_{-i,i}$ with $\Omega_{-i,-i} \hat{\beta}_i$.
- 7: **end for**
- 8: **until** convergence
- 9: **return** $K = \Omega^{-1}$

(algorithm Mohammadi 2019)

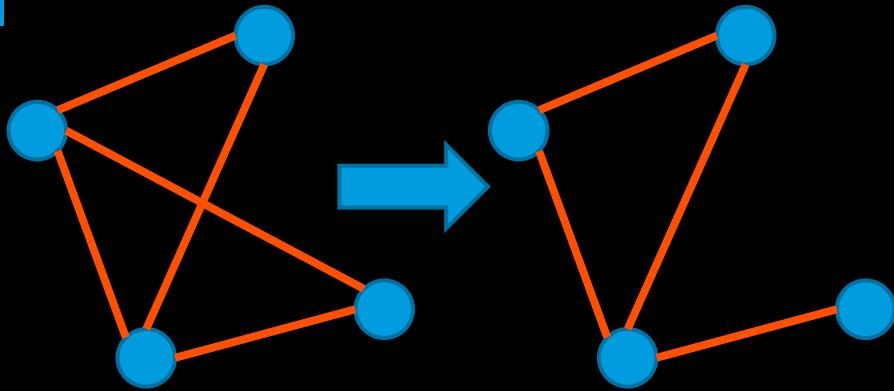
There exists algorithms to sample from a G-Wishart *without* having to calculate the normalizing constant!

Learning these edge weights is a relatively easy problem

Learning when edges are present/absent is **not** easy.



We would like to be able to...



DOUBLE REVERSIBLE JUMP METROPOLIS-HASTINGS MONTE CARLO

LEARNING THE *NUMBER* OF PARAMETERS, WHILE ALSO LEARNING THE PARAMETERS THEMSELVES.

This process combines two neat ideas:

1. Put a sampling distribution on the *Cholesky decomposition* $\Lambda = \Psi^T \Psi$ of the precision matrix Λ ;
2. Learn an ancillary grid of values along with the observed graph and precision matrix. Then, propose a new graph and matrix by *swapping*.

$$(\Lambda^d, G^d, \hat{\Lambda}^d, \hat{G}^d)$$

$$(\Lambda^{d+1}, G^{d+1}, \hat{\Lambda}^{d+1}, \hat{G}^{d+1})$$

SO WHERE ARE THE CHANGEPOINTS?

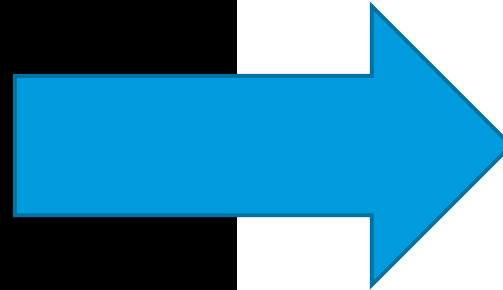
- (1, 1, 1, 2, 2, 2, 3, 3, 3)
- (1, 1, 2, 2, 2, 2, 2, 3, 3)
- (1, 1, 2, 2, 2, 2, 2, 2, 2)
- (1, 1, 2, 2, 3, 3, 3, 3, 3)
- (1, 1, 1, 1, 2, 2, 2, 2, 2)
- (1, 1, 1, 1, 2, 2, 2, 2, 3)
- (1, 1, 1, 1, 2, 2, 2, 2, 2)
- (1, 1, 1, 1, 1, 1, 1, 1, 1)

⋮

SO WHERE ARE THE CHANGEPOINTS?

- (1, 1, 1, 2, 2, 2, 3, 3, 3)
- (1, 1, 2, 2, 2, 2, 2, 3, 3)
- (1, 1, 2, 2, 2, 2, 2, 2, 2)
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- (1, 1, 1, 1, 2, 2, 2, 2, 2)
- (1, 1, 1, 1, 2, 2, 2, 2, 3)
- (1, 1, 1, 1, 2, 2, 2, 2, 2)
- (1, 1, 1, 1, 1, 1, 1, 1, 1)

⋮

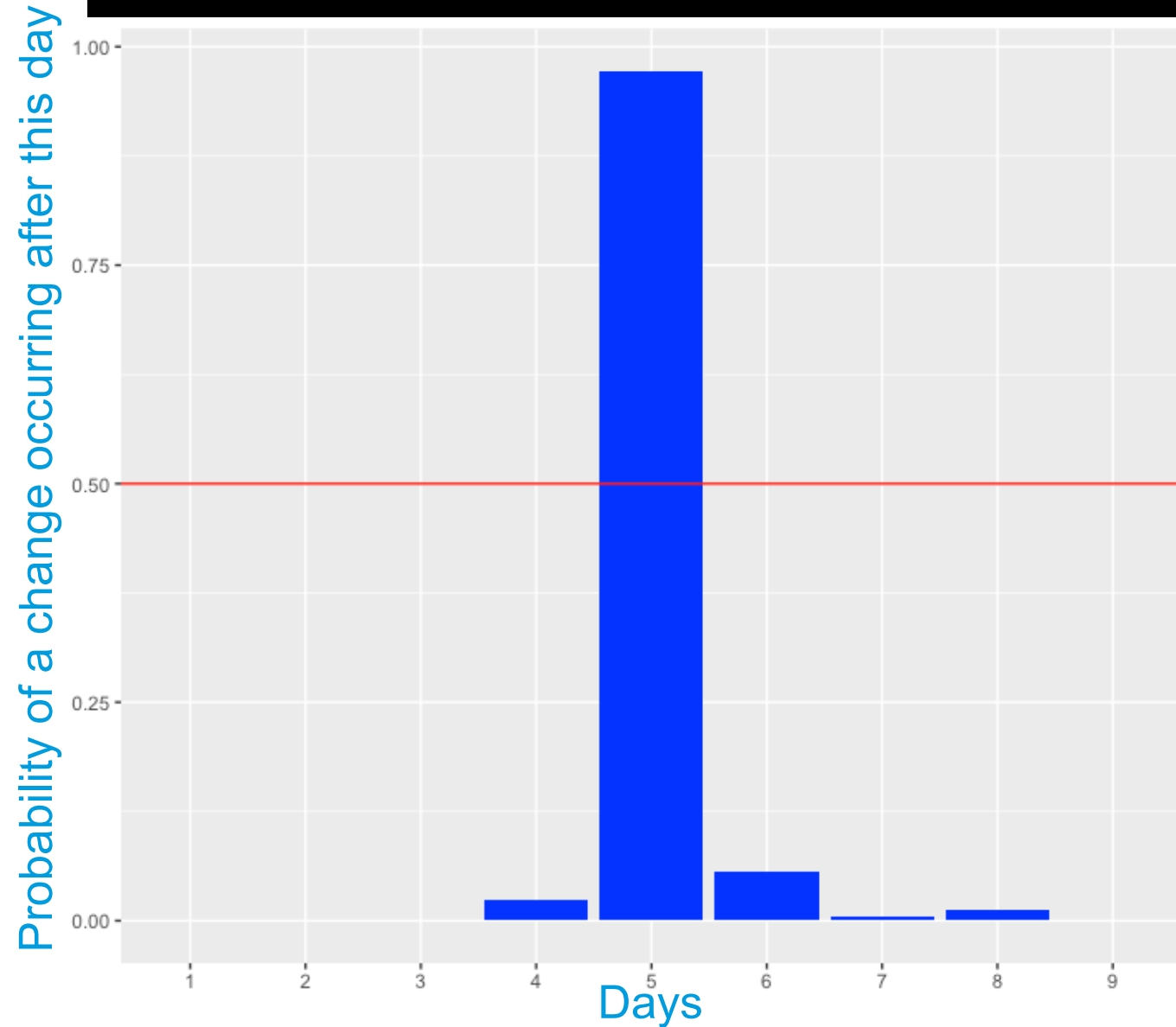
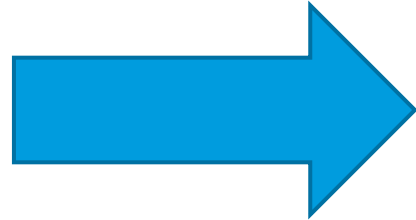


- (0, 0, 1, 0, 0, 1, 0, 0)
- (0, 1, 0, 0, 0, 0, 1, 0)
- (0, 1, 0, 0, 0, 0, 0, 0)
- (0, 1, 0, 1, 0, 0, 0, 0)
- (0, 0, 0, 1, 0, 0, 0, 0)
- (0, 0, 0, 1, 0, 0, 0, 1)
- (0, 0, 0, 1, 0, 0, 0, 0)
- (0, 0, 0, 0, 0, 0, 0, 0)

⋮

- (0, 0, 1, 0, 0, 1, 0, 0)
- (0, 1, 0, 0, 0, 0, 1, 0)
- (0, 1, 0, 0, 0, 0, 0, 0)
- (0, 1, 0, 1, 0, 0, 0, 0)
- (0, 0, 0, 1, 0, 0, 0, 0)
- (0, 0, 0, 1, 0, 0, 0, 1)
- (0, 0, 0, 1, 0, 0, 0, 0)
- (0, 0, 0, 0, 0, 0, 0, 0)

⋮



COMPETING METHODS

Hotelling T² (HT2) Sliding Window

Given four days of data, days 1,2, & 3 are compared against day 4.

$$T^* 2 = (\bar{y}_1 - \bar{y}_2) \left(\frac{S_1}{n_1} + \frac{S_2}{n_2} \right)^{-1} (\bar{y}_1 - \bar{y}_2) \sim T_{p,v}^2$$

\bar{y}_1 Mean value for days 1, 2, & 3

\bar{y}_2 Mean value for day 4

S_1 Sample variance for days 1, 2, & 3

S_2 Sample variance for day 4

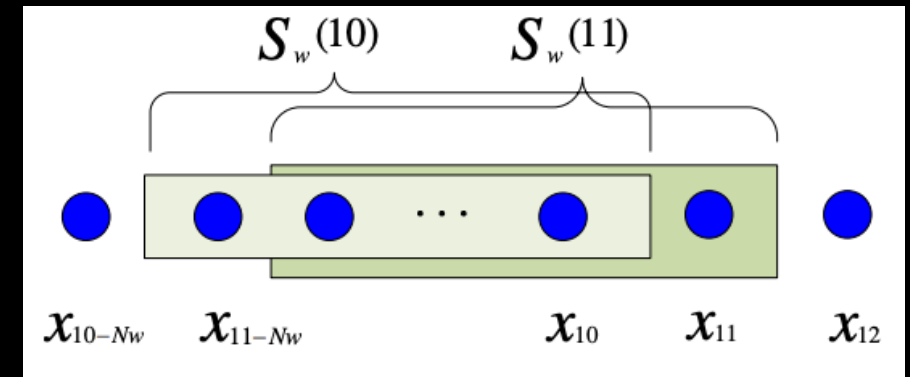
n_1 Number of observations for days 1, 2, & 3

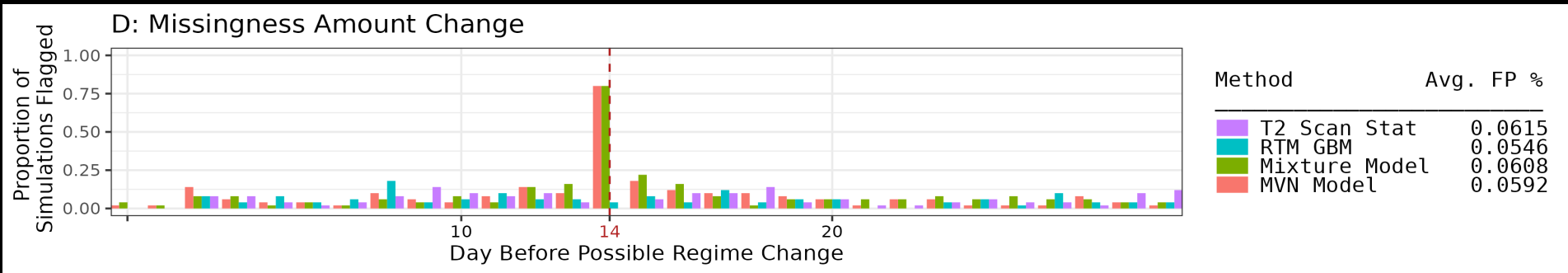
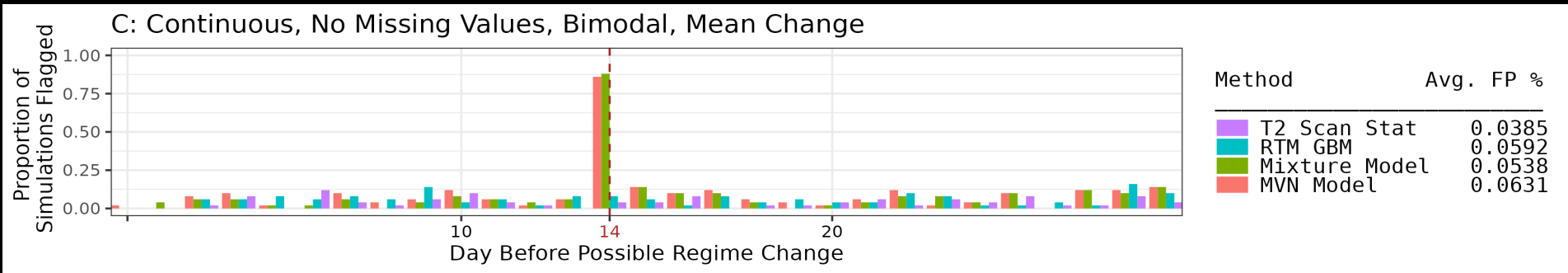
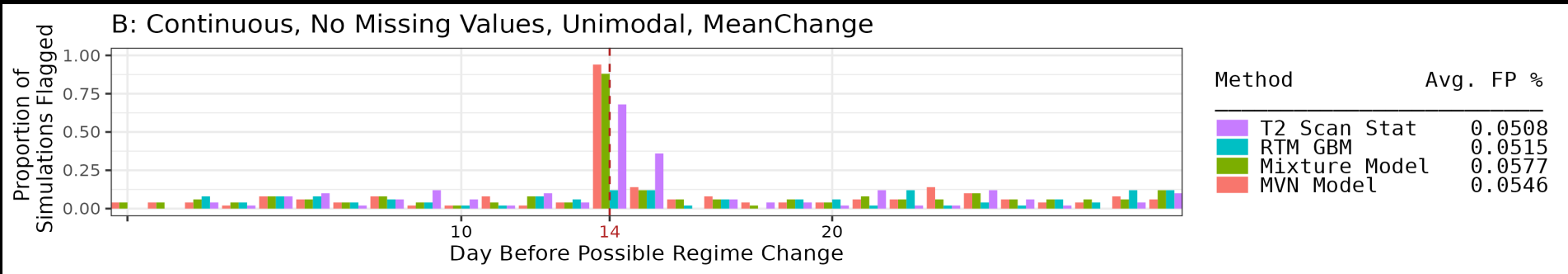
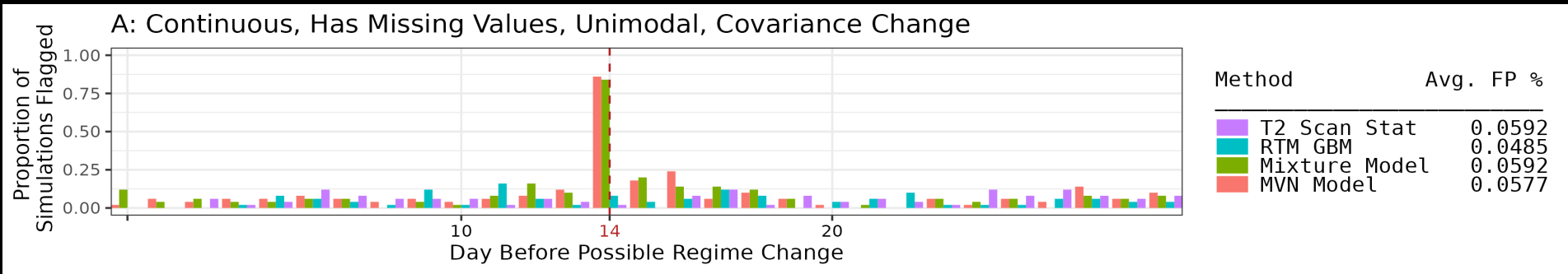
n_2 Number of observations for day 4

$T_{p,v}^2$ Hotelling T² statistic, with pooled degrees of freedom v and number of dimensions p

Real Time Contrasts (RTC) with a Gradient Boosting Machine (GBM) Classifier

- Given a classifier $c(x)$ learned to a hold-out sample, does this classifier have significant predictive power?
- We simulate a distribution of AUC values on $c(x)$'s ability to distinguish day 4 from days 1, 2, & 3.
- If 0.5 is NOT in the center 95% confidence region, a changepoint is flagged.





FAULT DETECTION ANALYSIS

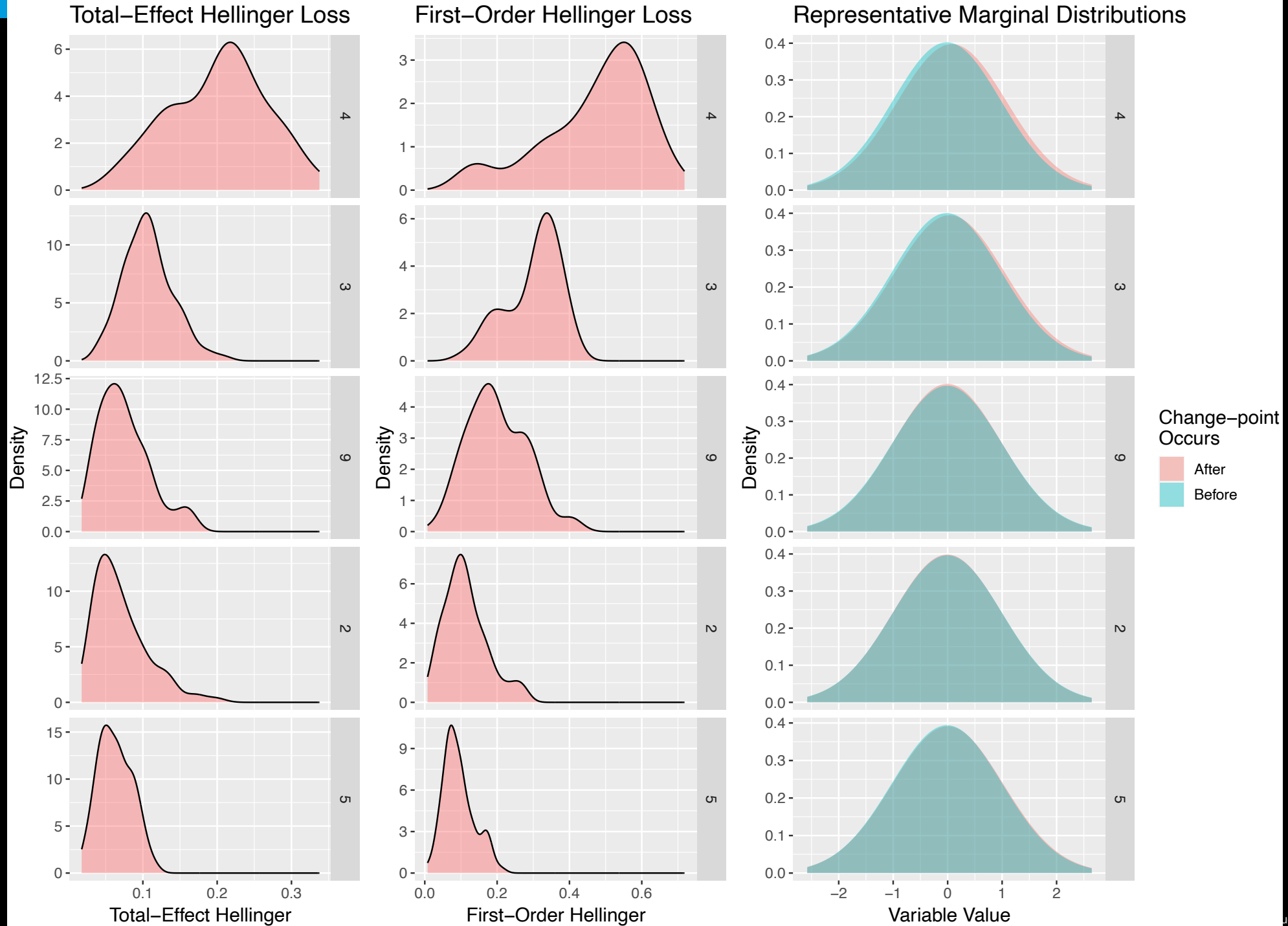
Once a changepoint is found, how do we determine what *caused* it?

Let	<u>Dist'n Before: X</u>	and	<u>Marginal Dist'n w/o</u>	then we	<u>Total-Effect Loss of i:</u>
	<u>Dist'n After: Y</u>		<u>Variable i: $X \setminus i$</u>		$1 - H(X \setminus i, Y \setminus i)/H(X, Y)$
	<u>Hellinger</u>		<u>Marginal Dist'n of</u>		<u>First-Order Loss of i:</u>
	<u>Distance: $H(X, Y)$</u>		<u>Variable i: $X : i$</u>	Define:	$H(X : i, Y : i)/H(X, Y)$

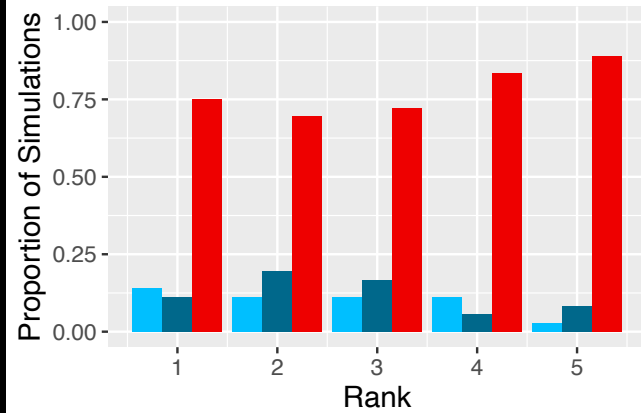
These metrics *decompose* the effects of each individual variable on the changepoint



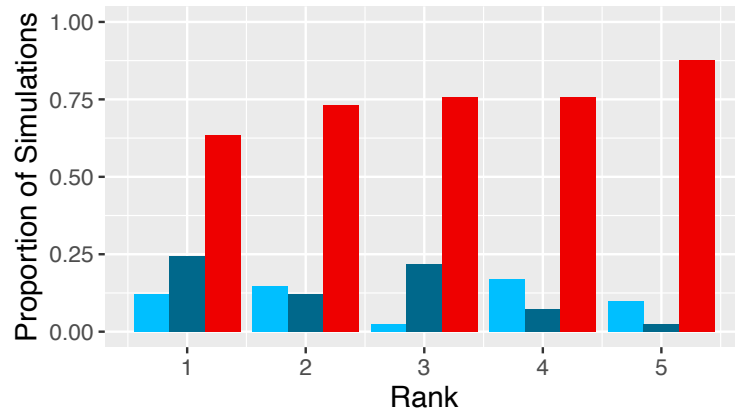
Simulation



GBM RTC, Simulation A



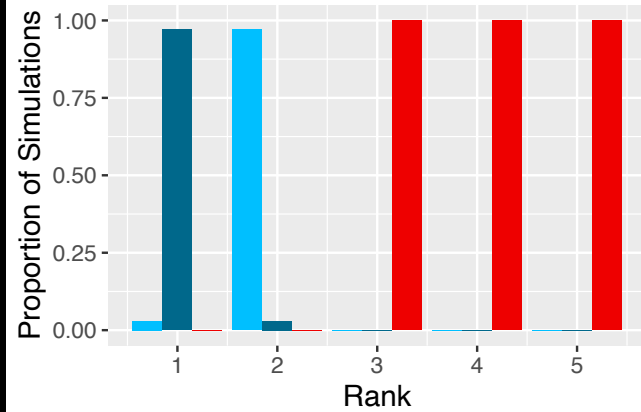
GBM RTC, Simulation B



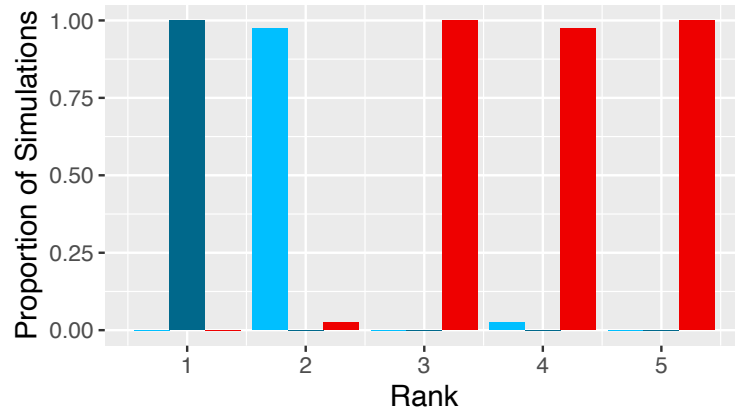
Variable

3
4
Other Vars

MVN Model, Simulation A



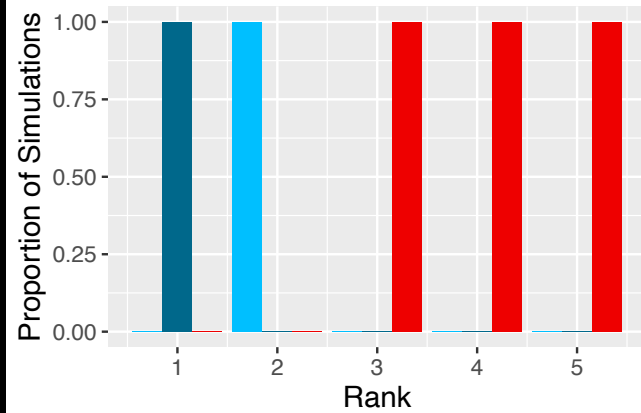
MVN Model, Simulation B



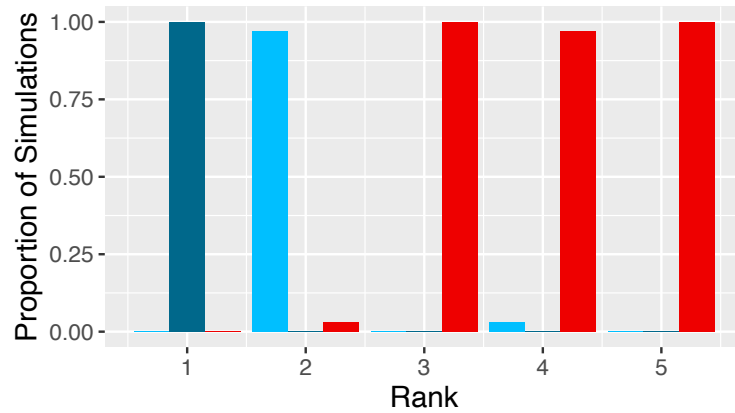
Variable

3
4
Other Vars

Mixture Model, Simulation A



Mixture Model, Simulation B



Variable

3
4
Other Vars

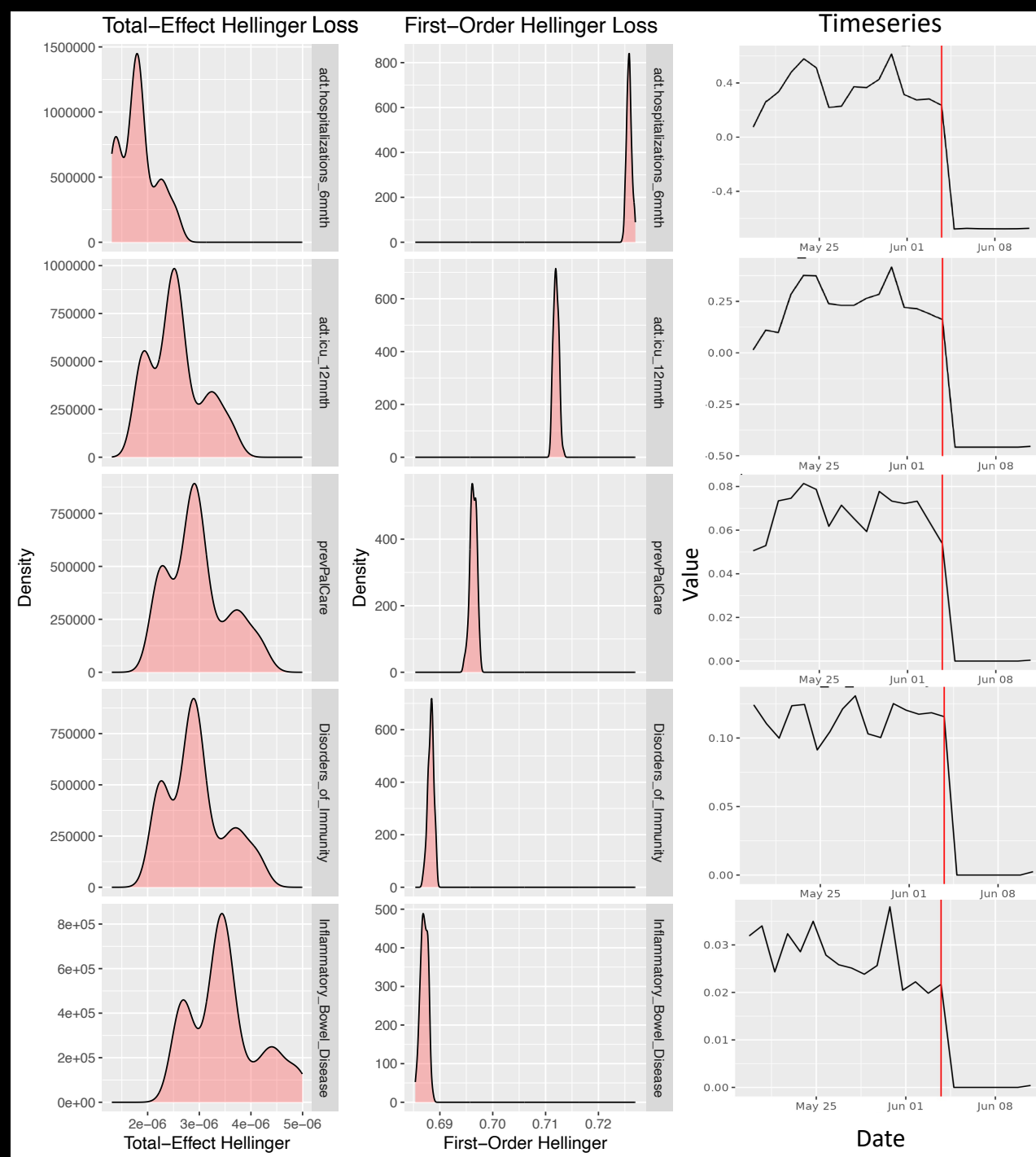
DATA APPLICATION

Let's investigate a month (mid-May to mid-June 2020) of the data from the Palliative Care Model

- Each observation in the data is a palliative care prediction that occurs for a patient at the Mayo Clinic in Rochester, MN.
- Variables include patient labs, the presence/absence of specific chronic diseases, basic demographics, and information about a patient's movement within the hospital.
- There are **123 variables**, 35 of which are continuous, with 27 variables having some amount missing values.
- During the month investigated, there were **298039 data observations**.

This is a *big* dataset with *many* data challenges; there are missing values, and various data types (censored & discrete variables).

Application



QUESTIONS?