

Network Systems
Science & Advanced
Computing

Biocomplexity Institute
& Initiative

University of Virginia

Estimation of COVID-19 Impact in Virginia

March 24th, 2021

(data current to March 20th – 23rd)

Biocomplexity Institute Technical report: TR 2021-029



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biocomplexity.virginia.edu

About Us

- Biocomplexity Institute at the University of Virginia
 - Using big data and simulations to understand massively interactive systems and solve societal problems
- Over 20 years of crafting and analyzing infectious disease models
 - Pandemic response for Influenza, Ebola, Zika, and others



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Overview

- **Goal:** Understand impact of COVID-19 mitigations in Virginia
- **Approach:**
 - Calibrate explanatory mechanistic model to observed cases
 - Project based on scenarios for next 4 months
 - Consider a range of possible mitigation effects in "what-if" scenarios
- **Outcomes:**
 - Ill, Confirmed, Hospitalized, ICU, Ventilated, Death
 - Geographic spread over time, case counts, healthcare burdens

Key Takeaways

Projecting future cases precisely is impossible and unnecessary.

Even without perfect projections, we can confidently draw conclusions:

- **Case rates in Virginia have flattened and now have some growth**
- VA mean weekly incidence slightly up to 17/100K from 15/100K, US flat (to 16.5 from 16 per 100K)
- Significant progress made in last month, however 81% of VA counties above mean rate of Summer 2020
- Projections are flattening out across Commonwealth, with growth on the horizon boosted by B.1.1.7
- Recent updates:
 - Slowed ramp up for new variant scenario given lack of clear evidence it has reached predominance this week
 - Adjusted Seasonal Effects scenarios to account for spring and summer weather
 - Accelerated vaccine schedule with Johnson & Johnson added as base case in anticipation of boost in vaccine supplies
- The situation continues to change. Models continue to be updated regularly.



Situation Assessment

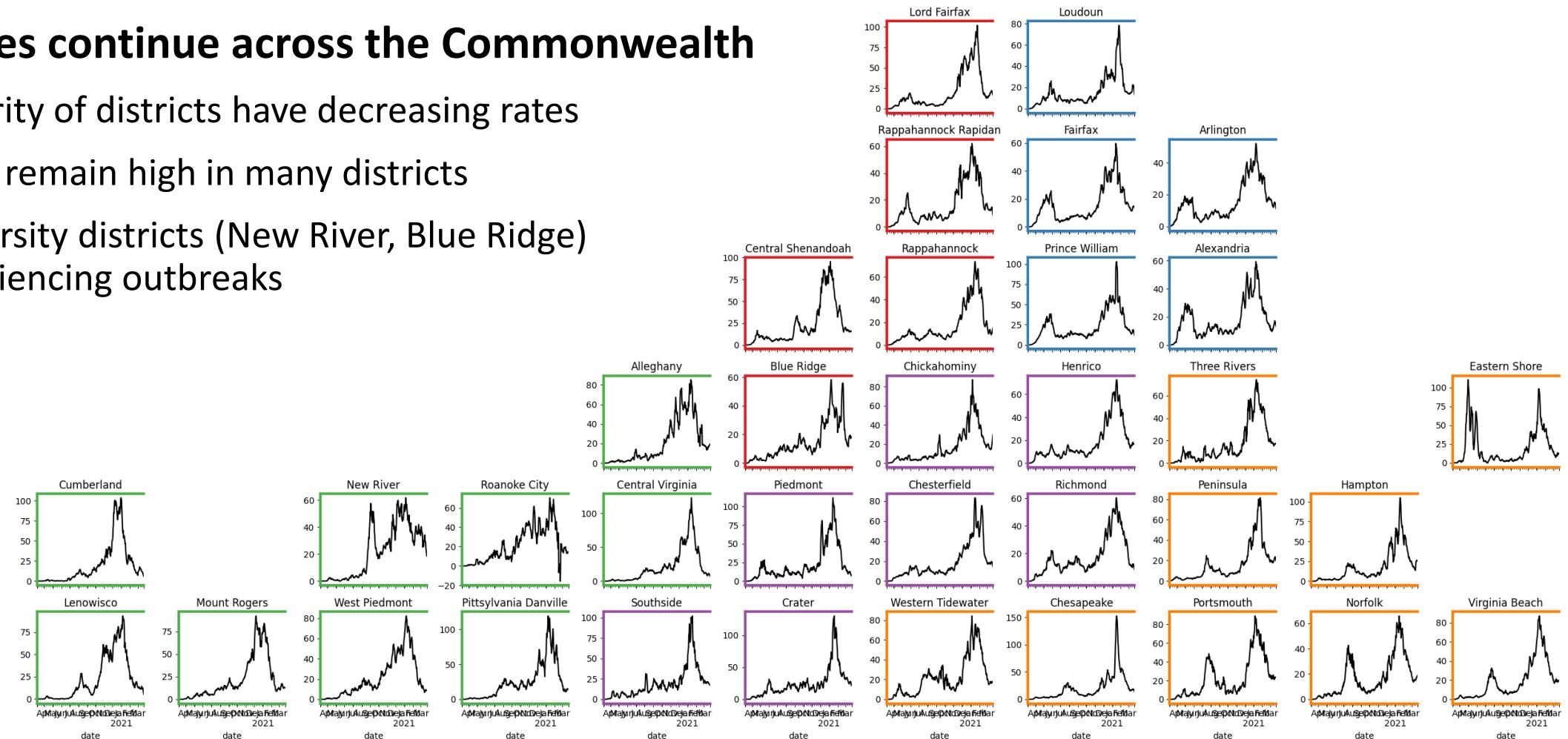


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Case Rate (per 100k) by VDH District

Declines continue across the Commonwealth

- Majority of districts have decreasing rates
- Rates remain high in many districts
- University districts (New River, Blue Ridge) experiencing outbreaks

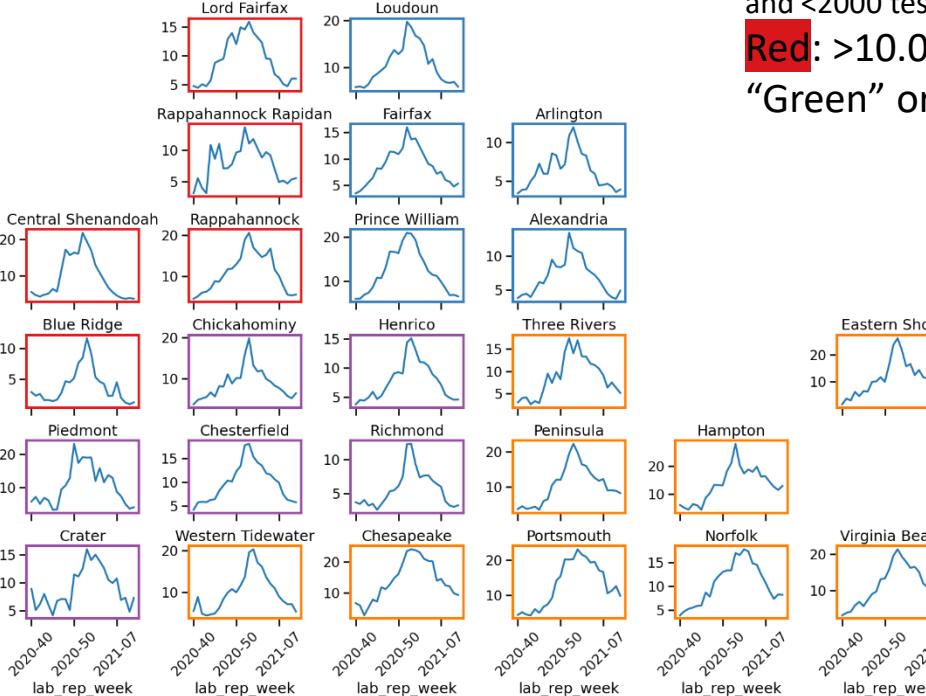
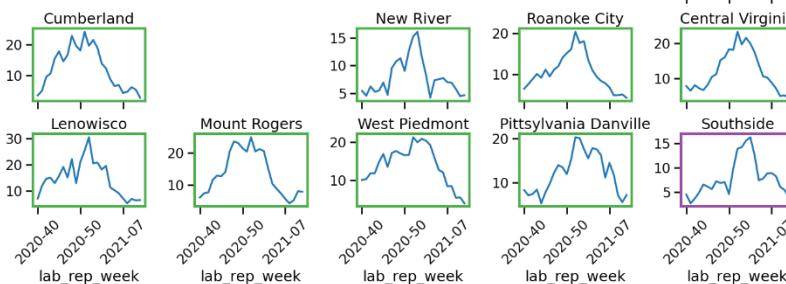


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Test Positivity by VDH District

Weekly changes in test positivity by district

- Rates continue to broadly decline, however some leveling off seen in some districts
- More counties are below 5%, 5-10% starts to decline as well



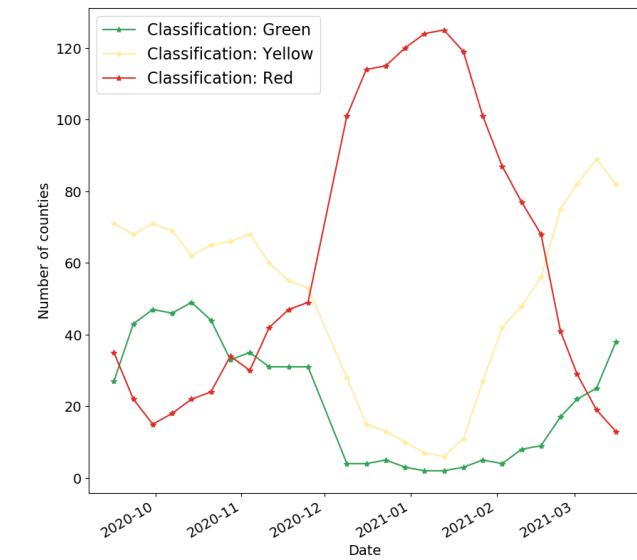
County level test positivity rates for RT-PCR tests.

Green: Test positivity <5.0%

(or with <20 tests in past 14 days)

Yellow: Test positivity 5.0%-10.0% (or with <500 tests and <2000 tests/100k and >10% positivity over 14 days)

Red: >10.0% and not meeting the criteria for “Green” or “Yellow”

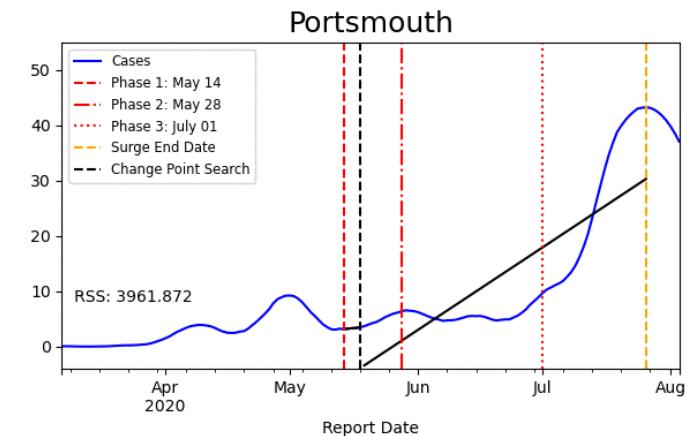


District Trajectories

Goal: Define epochs of a Health District's COVID-19 incidence to characterize the current trajectory

Method: Find recent peak and use hockey stick fit to find inflection point afterwards, then use this period's slope to define the trajectory

Hockey stick fit



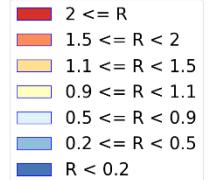
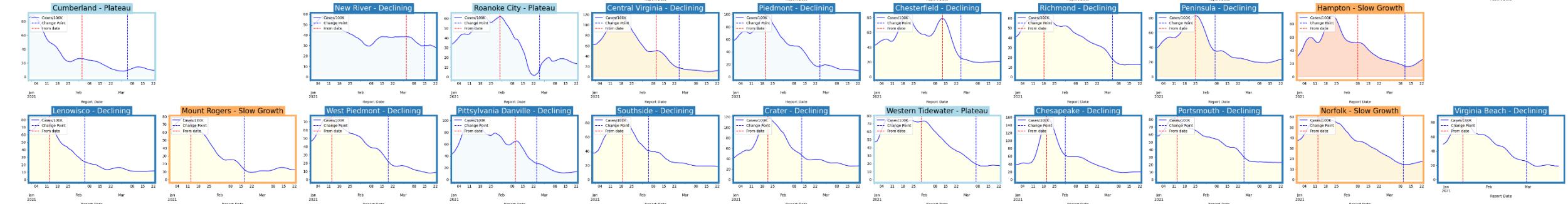
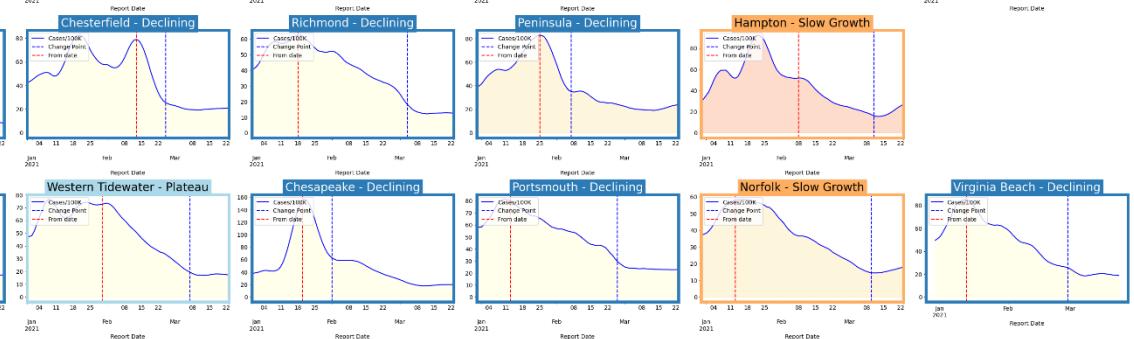
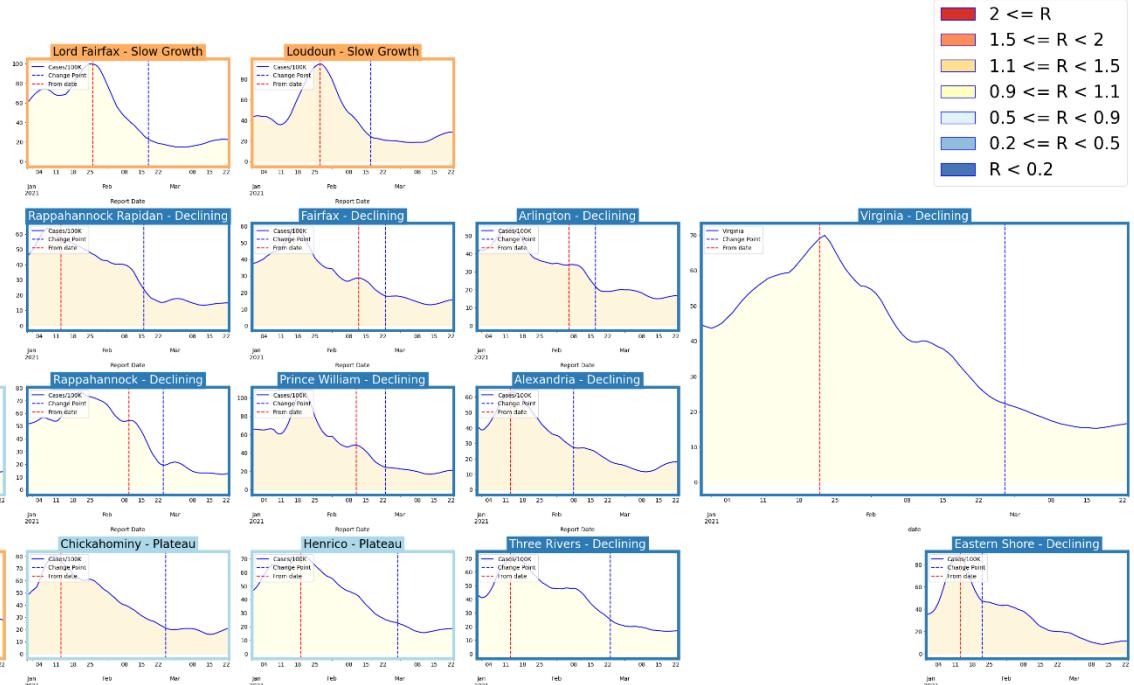
Trajectory	Description	Weekly Case Rate (per 100K) bounds	# Districts (prev week)
Declining	Sustained decreases following a recent peak	below -0.9	23 (31)
Plateau	Steady level with minimal trend up or down	above -0.9 and below 0.5	6 (2)
Slow Growth	Sustained growth not rapid enough to be considered a Surge	above 0.5 and below 2.5	6 (2)
In Surge	Currently experiencing sustained rapid and significant growth	2.5 or greater	0 (0)



District Trajectories – last 10 weeks

Status	# Districts (prev week)
Declining	23 (31)
Plateau	6 (2)
Slow Growth	6 (2)
In Surge	0 (0)

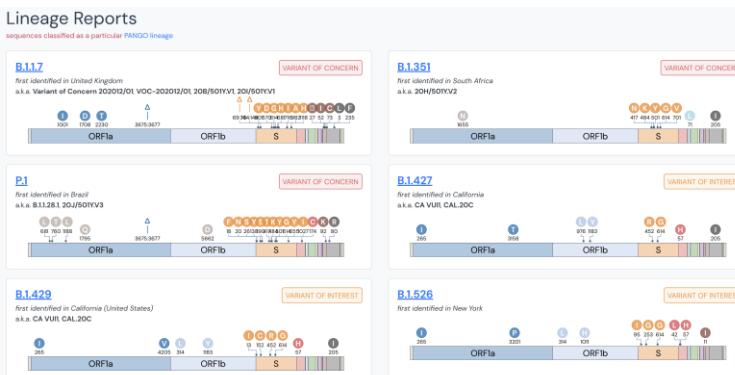
Curve shows smoothed case rate (per 100K)
 Trajectories of states in label & chart box
 Case Rate curve colored by Reproductive



SARS-CoV2 Variants of Concern

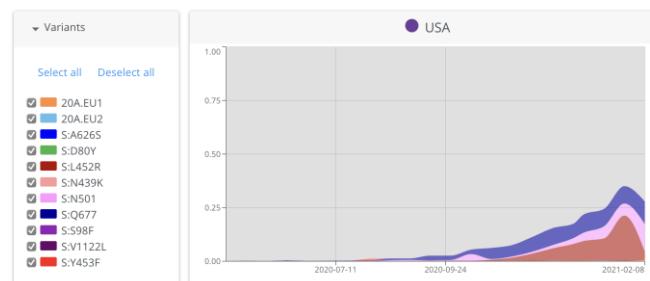
Emerging new variants will alter the future trajectories of pandemic and have implications for future control

- Current evidence supports that new variants can:
 - Increase transmissibility
 - Increase severity (more hospitalizations and/or deaths)
 - Limit immunity provided by prior infection and vaccinations
- Genomic surveillance remains very limited
 - Challenges ability to estimate impact in US to date and estimation of arrival and potential impact in future
 - B.1.1.7 is most frequent and well studied



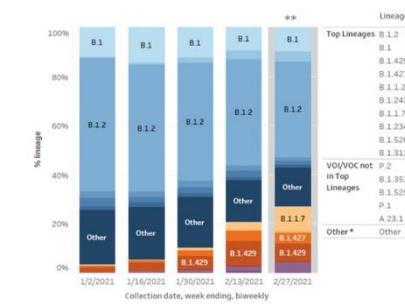
outbreak.info

[Outbreak Info](#)

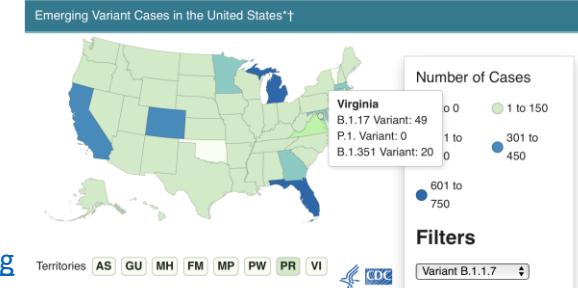


Lineages Of Concern							
LoC name	PANGO lineage	NextStrain lineage	Other synonyms	Emergence date	Emergence location	Key AA substitutions in spike protein	Impact
B.1.1.7	B.1.1.7	20I/501Y.V1	VOC 202012/01, UK variant	September 2020	Southeast England	H69-, V70-, N501Y, D614G, P681H	Increased transmissibility; S gene target failure (SGTF)
B.1.351	B.1.351	20H/501Y.V2	South African variant	October 2020	Nelson Mandela Bay, South African	L241-, L242-, A243-, K417N, E484K, N501Y, D614G	loss of serum antibody neutralization
P.1	B.1.1.28	20J/501Y.V3	Brazilian variant	July 2020	Brazil	K417T, E484K, N501Y, D614G	Increased transmissibility; loss of serum antibody neutralization
CAL.20C	B.1.429			July 2020	Southern California, USA	W152C, L452R, D614G	loss of monoclonal antibody binding
B.1.375	B.1.375			September 2020	Massachusetts, USA	H69-, V70-, D614G	S gene target failure (SGTF)

[NIH-NIAID Bacterial-Viral Bioinformatics Resource Center](#)



Variant	Reported Cases in US	Number of Jurisdictions Reporting
B.1.1.7	7,501	51
B.1.351	219	27
P.1	61	18



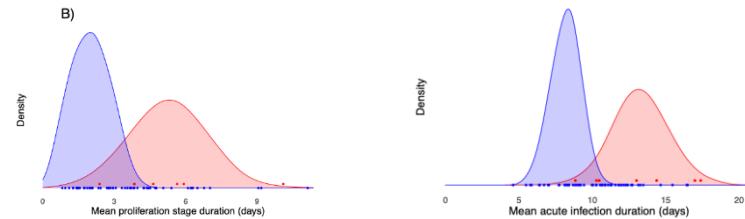
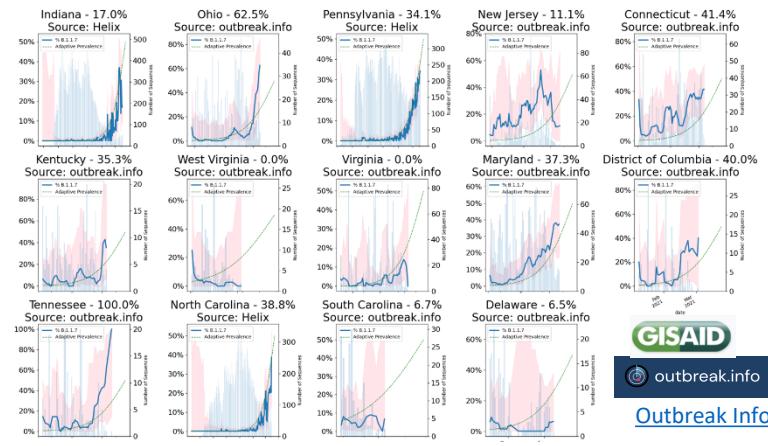
[CDC Variant Tracking](#)

Territories AS GU MH FM MP PW PR VI CDC

SARS-CoV2 Variants of Concern

Lineage B.1.1.7

- B.1.1.7 has been detected in Virginia and has continued to rapidly grow. Reporting delays don't permit a current estimate this week, however, neighboring states suggest VA may be around 35% (national frequency at ~45%)
- Virginia remains behind but still within bounds of estimates based on growth rates indicating it will predominate (eg reach 50% frequency) by late March and is more transmissible
- A cluster with the E484K mutation has been described indicating multiple independent acquisitions in UK, potential for aiding immune escape
- A recent study finds B.1.1.7 to have longer duration which may be the source of increased transmissibility and has implications for isolation durations
- Evidence continues to mount supporting increased risks of hospitalization and mortality for B.1.1.7 infected individuals
- Update to Rasmussen et al. study estimates B.1.1.7 to have the highest "fitness" advantage of all observed variants and mutations of note

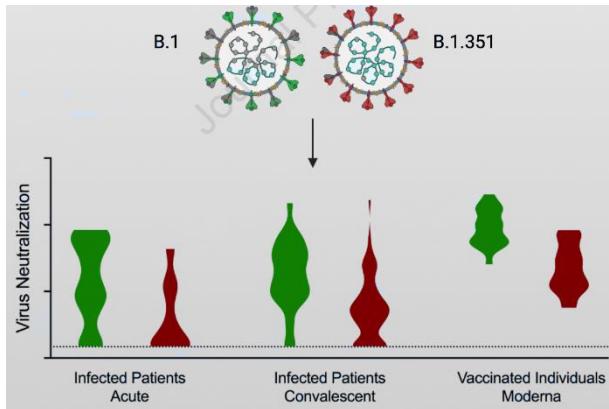


Variant B.1.1.7 may cause longer infections with similar peak viral concentration compared to non-B.1.1.7. May contribute to B.1.1.7's increased transmissibility.
<https://dash.harvard.edu/handle/1/37366884>

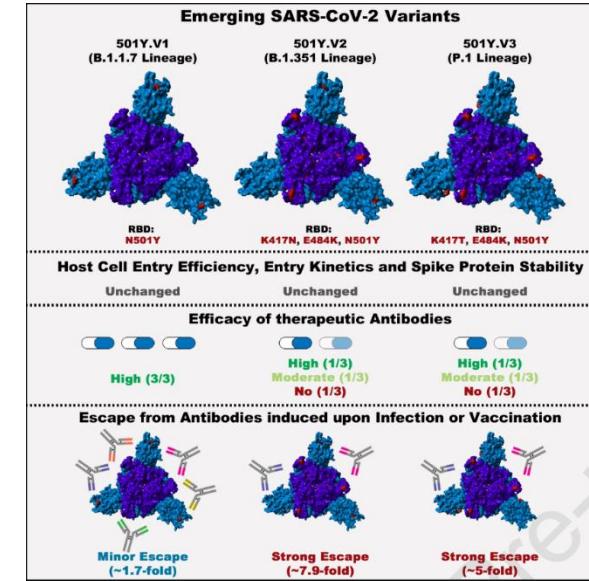
SARS-CoV2 Variants of Concern

Lineage B.1.351

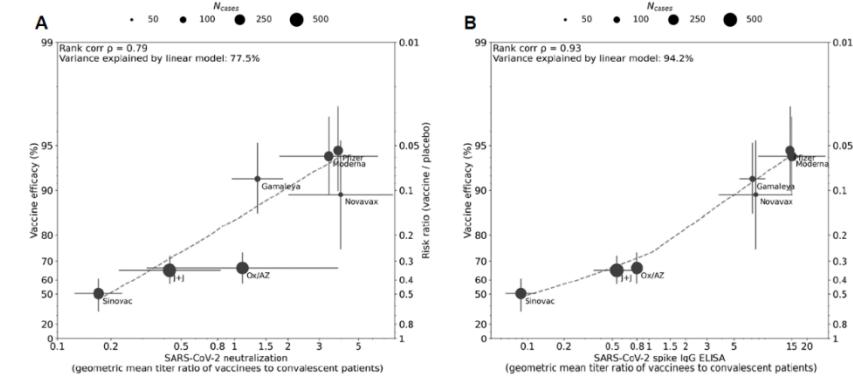
- Emerging strain initially identified in South Africa shows signs of vaccine escape, currently 219 reported cases in 27 states (including 26 now in Virginia) as of Mar 23rd
- An additional study corroborates recent study based on clinical trial data shows that convalescent serum neutralization is highly predictive of actual immune protection for infection, thus B.1.351 may require booster vaccinations, and provides estimates for timing
- Study in Cell shows B.1.351 resists neutralization from anti-body treatments and another further shows resistance in pseudo-virus neutralization
- Another study in Cell supports previous report that demonstrated that despite reduced antibody binding the Moderna vaccinated individuals able to neutralize the B.1.351 variant



Despite reduced antibody binding to the B.1.351 RBD, sera from infected (acute and convalescent) and Moderna (mRNA-1273) vaccinated individuals were still able to neutralize the SARS-CoV-2 B.1.351 variant. [Cell](#)



Entry driven by S-proteins of all variants were inhibited by Imdevimab. B.1.351 and P.1 showed resistance to neutralization by Casirivimab. A cocktail of both REGN-COV2 efficiently inhibited cell entry. P.1 and B.1.351 S-protein mutations demonstrated markedly reduced inhibition on sera from Pfizer vaccinated patients. [Cell](#)

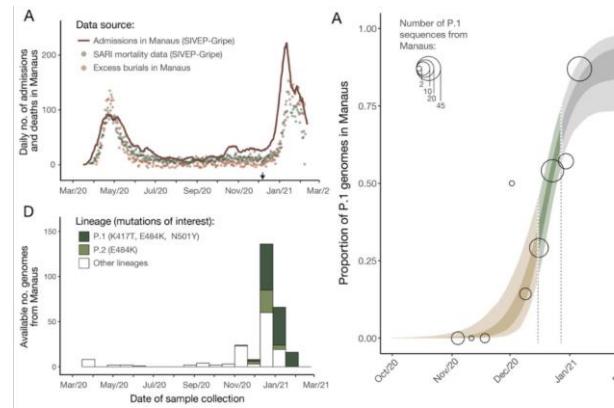


Corroborating: calibrated to titers of human convalescent sera reported in each study, a robust correlation was seen between neutralizing titer and efficacy ($p=0.79$) and binding antibody titer and efficacy ($p=0.93$) [MedArxiv](#)

SARS-CoV2 Variants of Concern

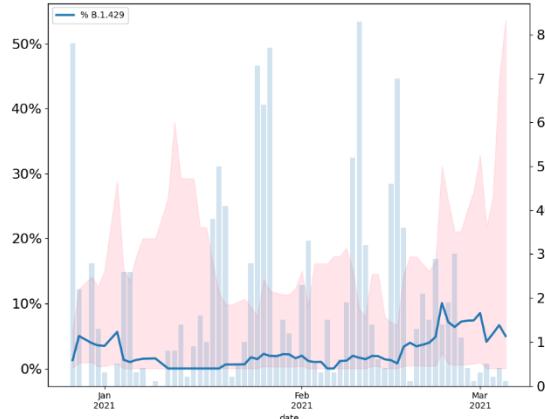
Lineage P.1

- Present in at least 61 cases in 18 states, shows signs of increased transmissibility and ability to evade immunity
- Caused a [resurgence of hospitalizations in Manaus, Brazil](#) which has now caused more deaths in last 3 months than all of 2020
- [Recent study](#) estimates it to be 1.4-2.2 times more transmissible and able to partially evade protective immunity

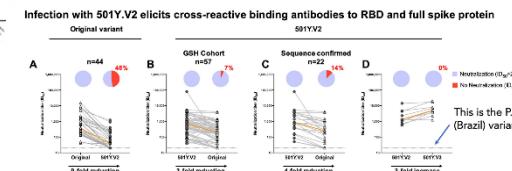
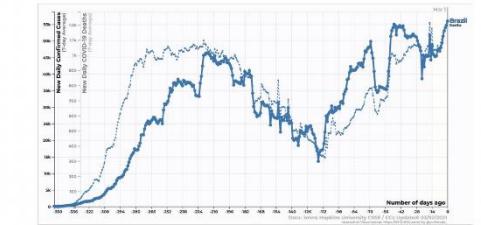


[Estimate](#) that P.1 may be 1.4–2.2 times more transmissible and able to evade 25–61% of protective immunity elicited by previous infection with non-P.1 lineages.

Virginia - 5.0% (B.1.429)
Source: outbreak.info



The tragedy in Brazil <https://t.co/GCCxhrlf04> by @terrence_mccoy "The variant known as P.1, which was discovered earlier this year, has stamped the Amazonian city of Manaus, leading to more deaths in January and February than in all of 2020."



Limited neutralization from COVID-19 patient sera. [BioRxiv](#)



Estimating Daily Reproductive Number

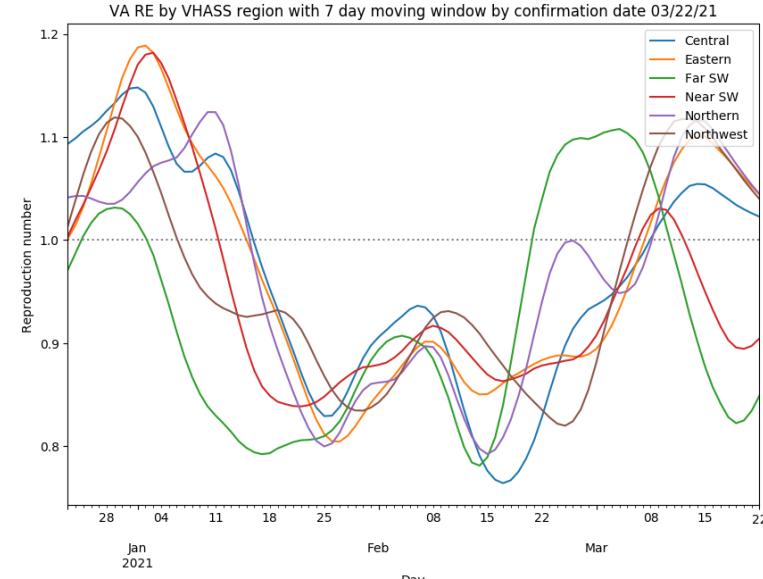
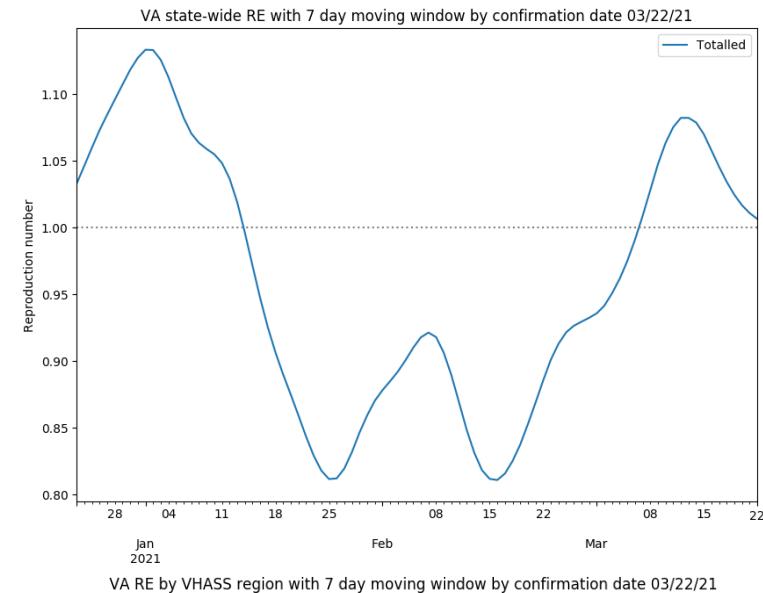
March 22nd Estimates

Region	Date Confirmed R _e	Date Confirmed Diff Last Week
State-wide	1.006	0.052
Central	1.023	0.099
Eastern	1.044	0.056
Far SW	0.849	-0.335
Near SW	0.905	-0.071
Northern	1.045	0.142
Northwest	1.040	0.057

Methodology

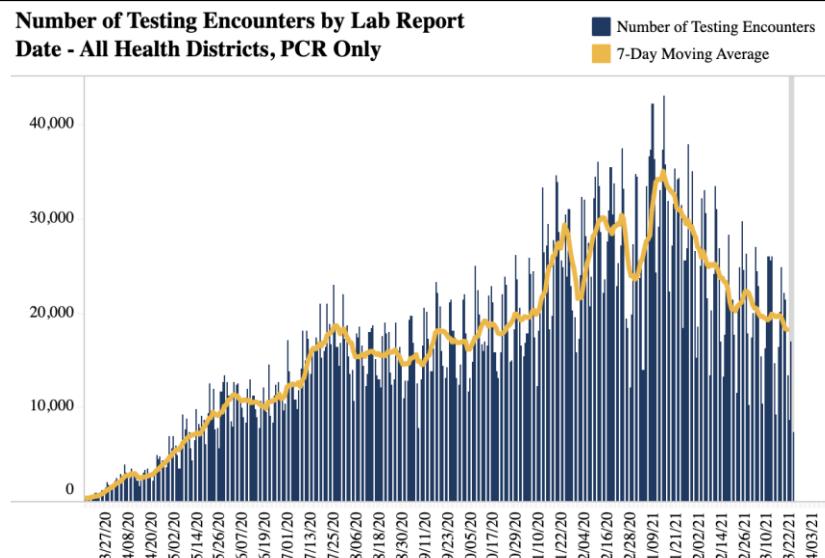
- Wallinga-Teunis method (EpiEstim¹) for cases by confirmation date
- Serial interval: updated to discrete distribution from observations (mean=4.3, Flaxman et al, Nature 2020)
- Using Confirmation date since due to increasingly unstable estimates from onset date due to backfill

1. Anne Cori, Neil M. Ferguson, Christophe Fraser, Simon Cauchemez. A New Framework and Software to Estimate Time-Varying Reproduction Numbers During Epidemics. American Journal of Epidemiology, Volume 178, Issue 9, 1 November 2013, Pages 1505–1512, <https://doi.org/10.1093/aje/kwt133>



Changes in Case Detection

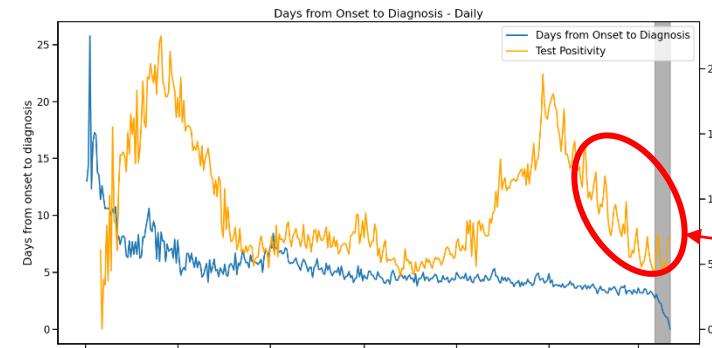
Timeframe (weeks)	Mean days	% difference from overall mean
July (26-30)	6.2	-7%
Aug (31-34)	4.9	-26%
Sept (35-38)	4.5	-32%
Oct (39-43)	4.5	-33%
Nov (44-47)	4.5	-32%
Dec (48-49)	4.2	-37%
Jan (00-04)	3.9	-41%
Feb (05-08)	3.4	-49%
Mar (09)	3.3	-50%
Overall (13-09)	6.7	--



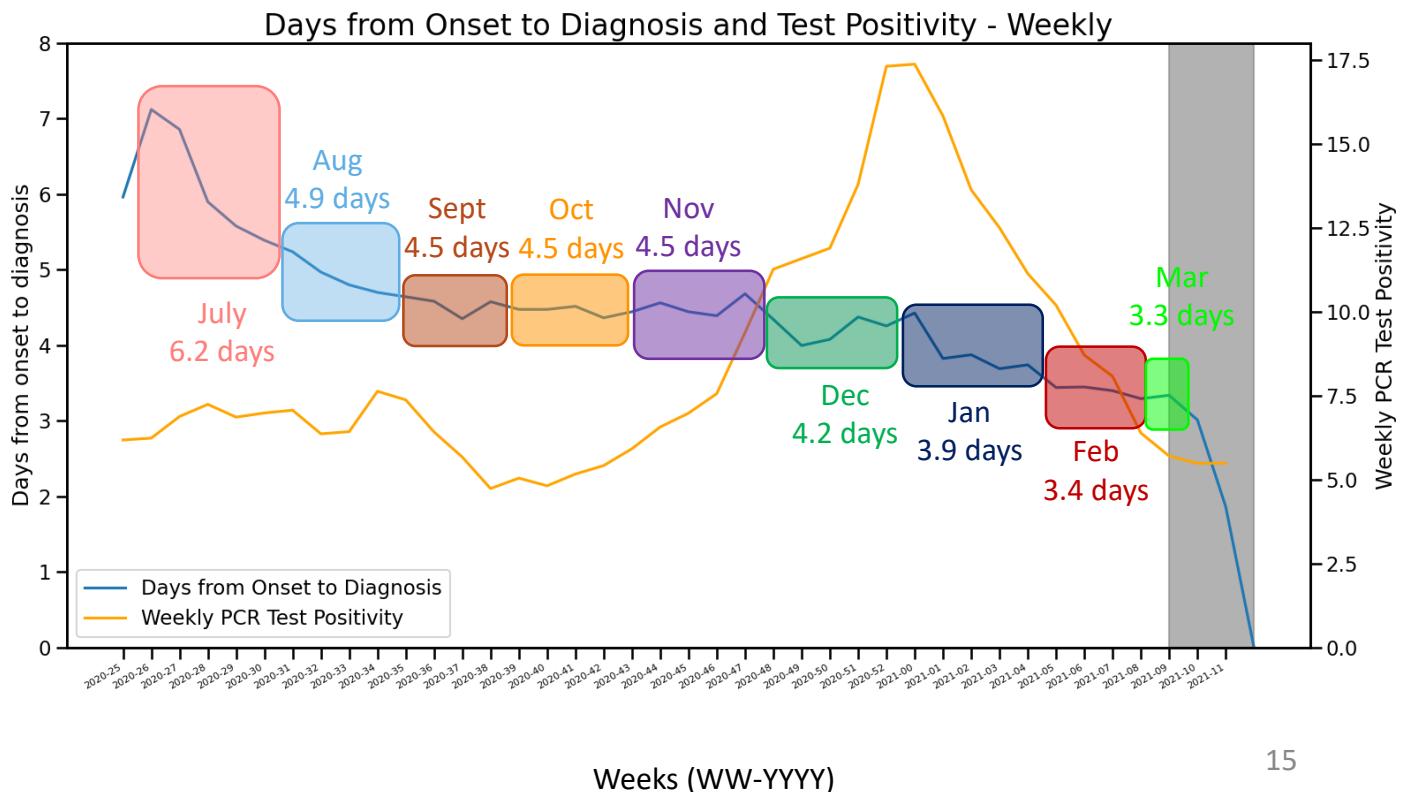
25-Mar-21

Accessed 9:00am March 24, 2021
<https://www.vdh.virginia.gov/coronavirus/>

Test positivity vs. Onset to Diagnosis

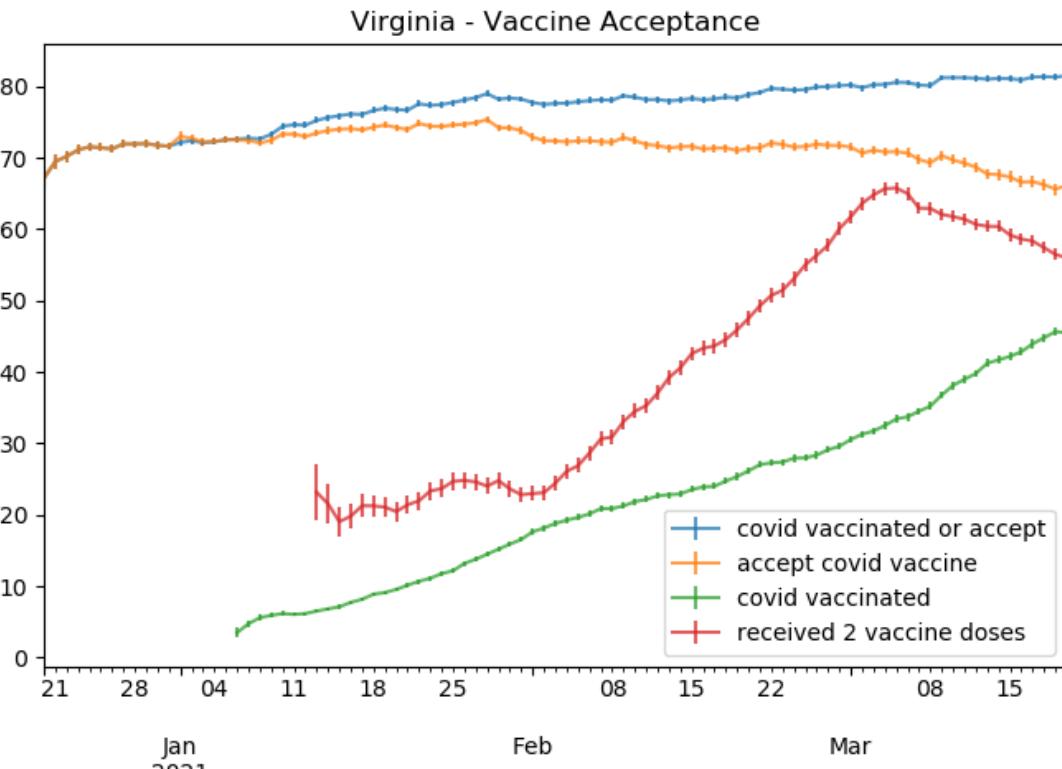


Positivity continues its rapid decline



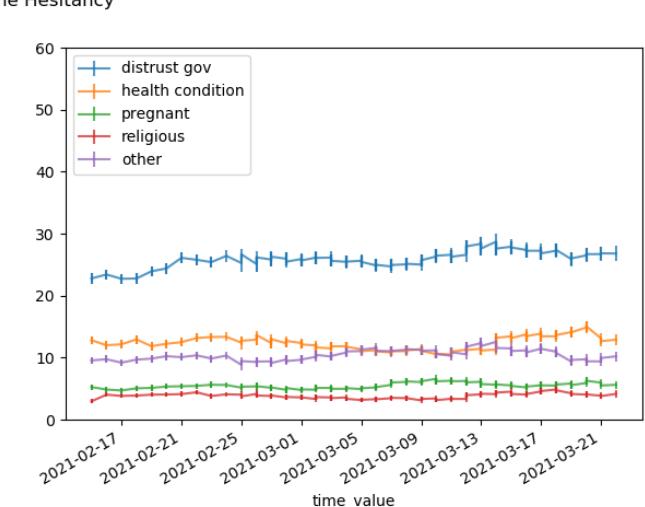
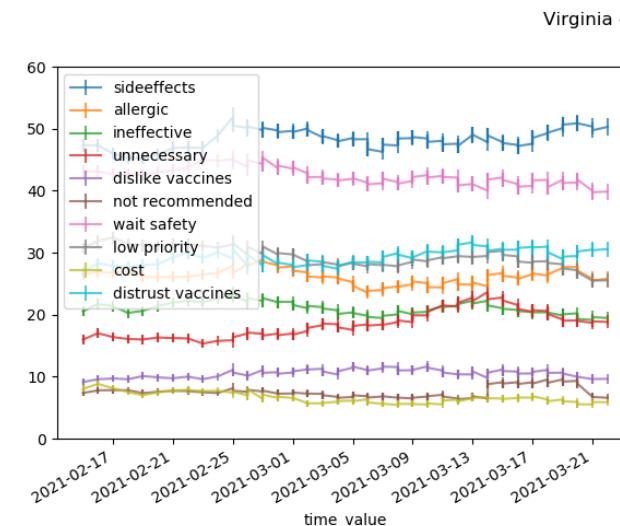
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Vaccine Acceptance in Virginia



Acceptance remains high:

- Proportion of Virginians that would definitely or probably accept vaccination if offered today
- Nearly 80% Virginians have already or will choose to be vaccinated
- Top reasons for hesitancy: side effects, safety, distrust
- Some slight movement up in side effects, and decreases in unnecessary and ineffective



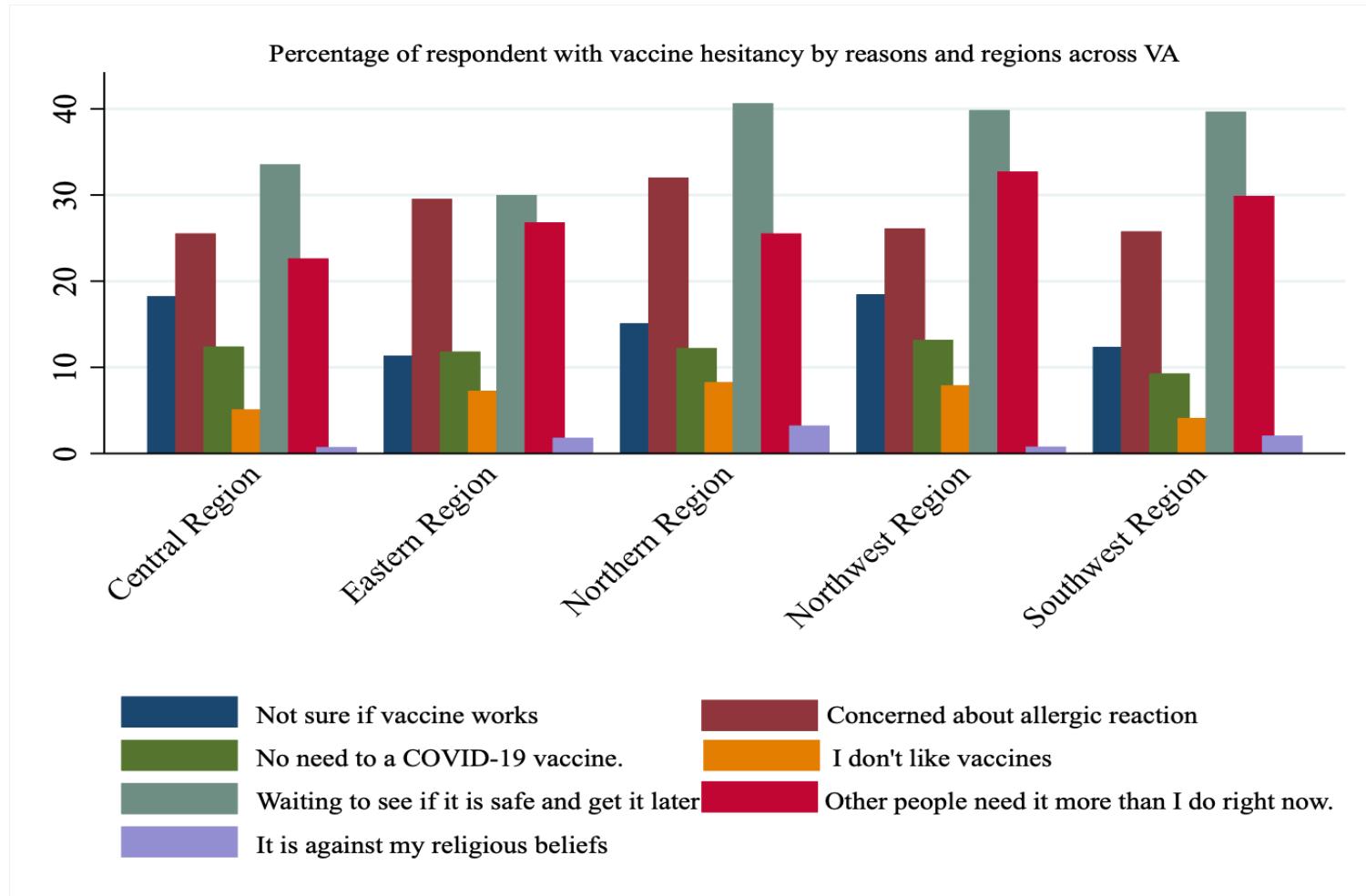
Data Source: <https://covidcast.cmu.edu>



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Vaccine Hesitancy in Virginia



Reasons over the past month by region:

- Preliminary analysis using unweighted responses
- Waiting to assess safety and allergic reactions cited as health-related reasons to hold off.
- Concerns that others should be vaccinated before them, additionally cited as a reason for hesitancy

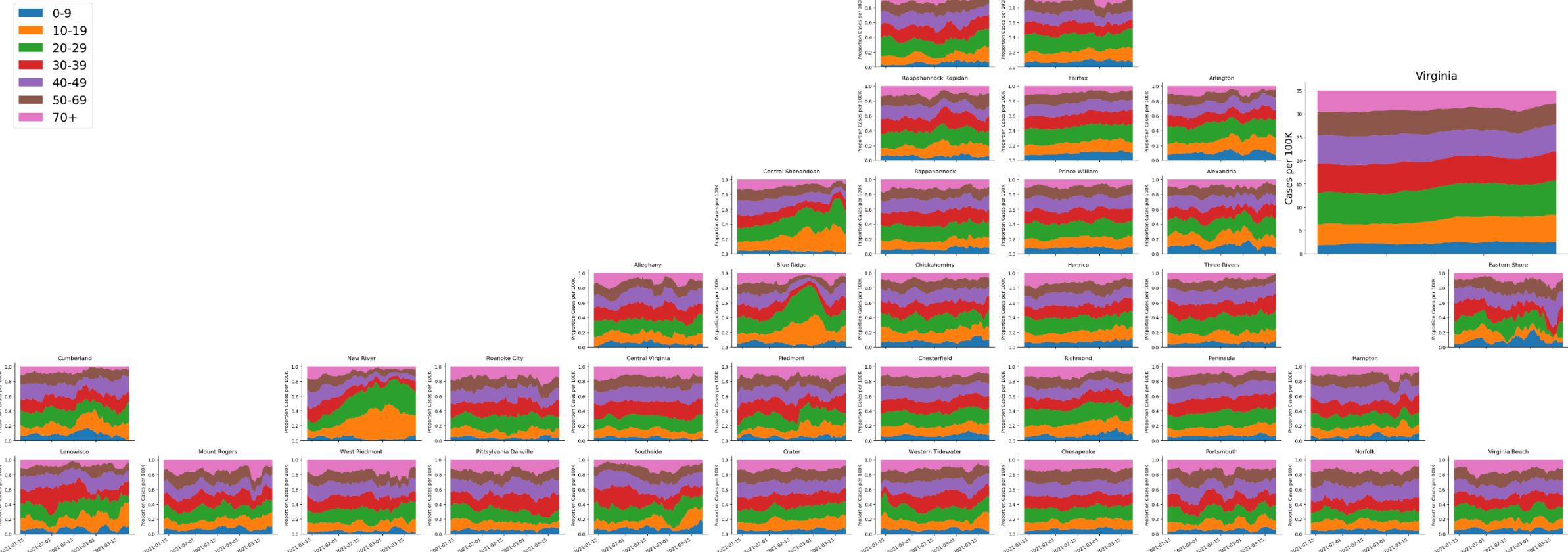
Data Source: <https://covidcast.cmu.edu>



Age-Specific Case Rates

Rates per 100K of each Age-Group, compared proportionally over time by district (last 10 weeks)

- Highlights changes in epidemiology and the overall shift in burden
- In many districts younger age-groups now have proportionally higher case rates

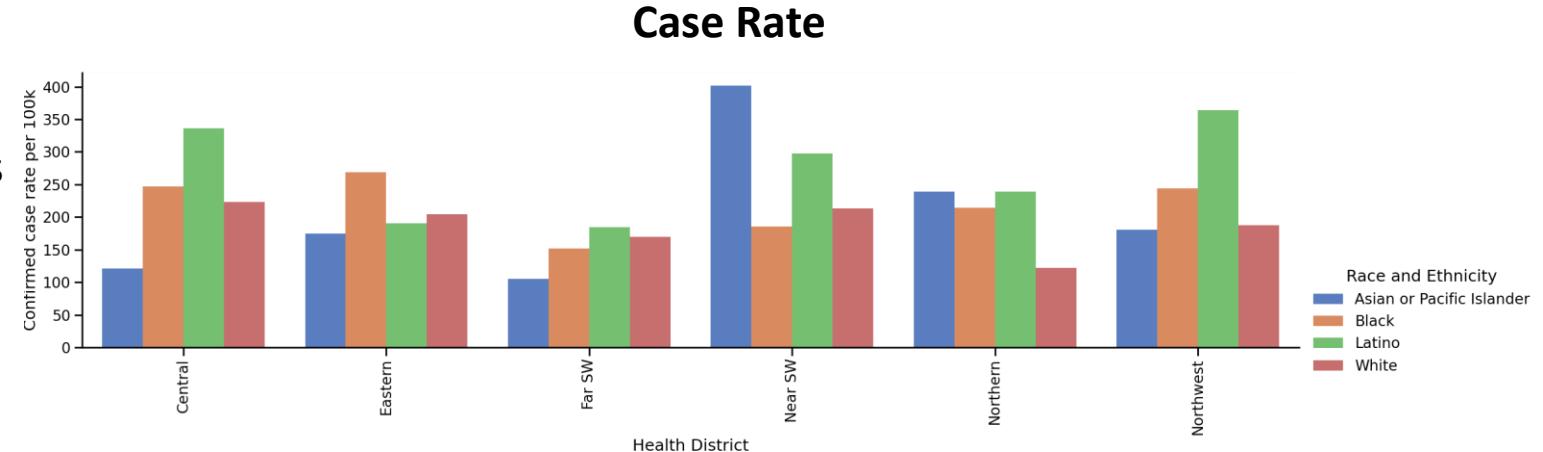


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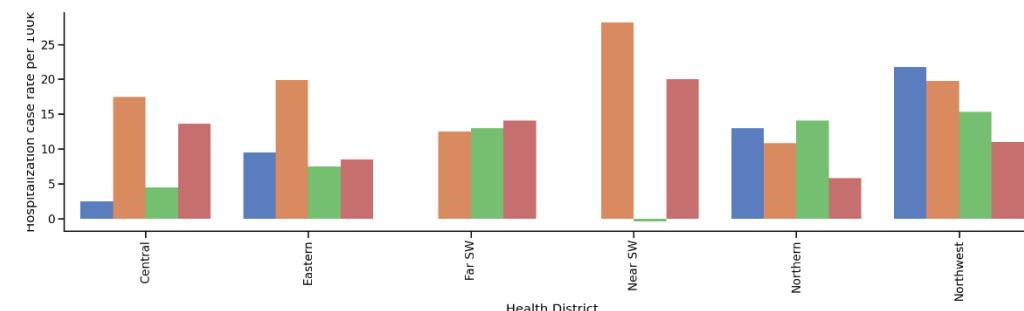
Race and Ethnicity – Recent Rate Changes (per 100K)

Changes in Race and Ethnicity Rates (per 100k) in past two weeks

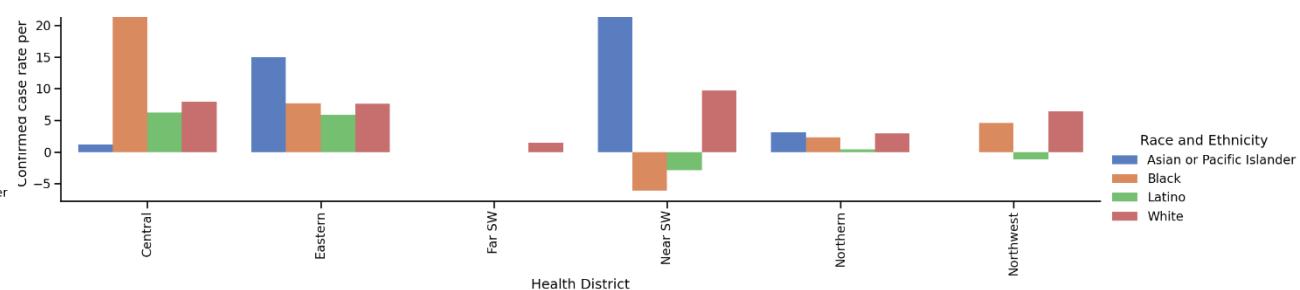
- Two week change in population level rates
- Black, Latinx and 2 or more races populations have much higher changes in rates; disparity is more pronounced in some regions than others
- Based on 2019 census race-ethnicity data by county



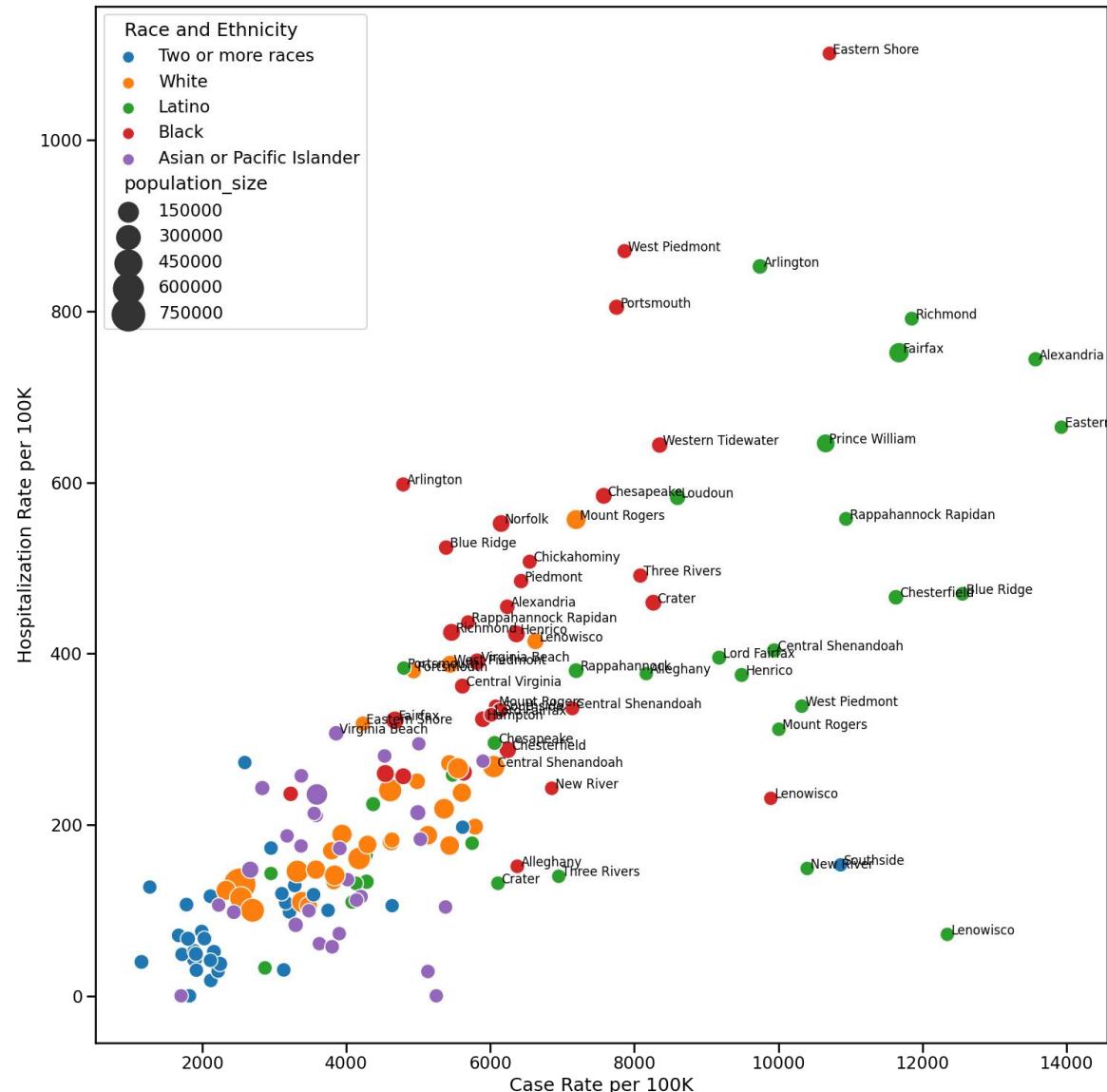
Hospitalization Rate



Death Rate



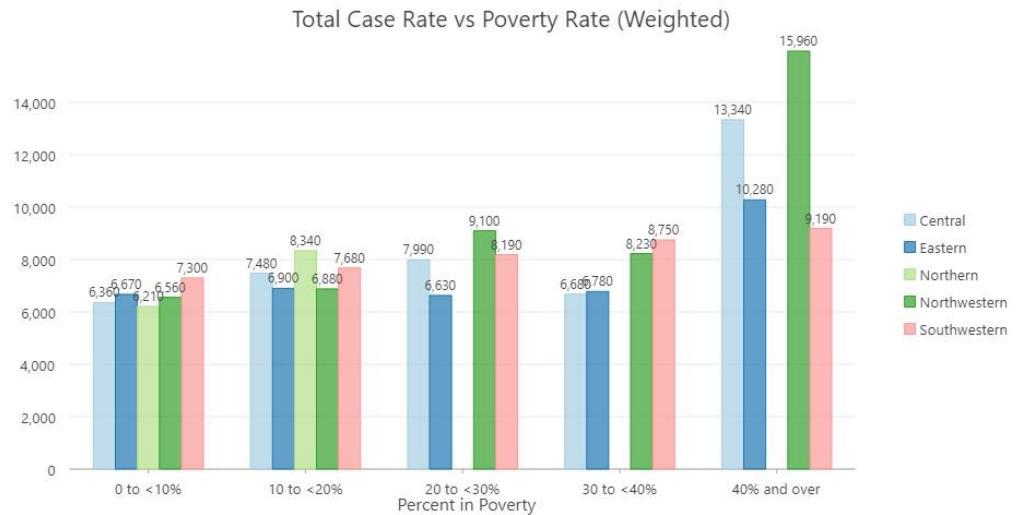
Race and Ethnicity cases per 100K



Rates per 100K of each Racial-Ethnic population by Health District

- Each Health District's Racial-Ethnic population is plotted by their Hospitalization and Case Rate
- Points are sized based on their overall population size (overlapping labels removed)

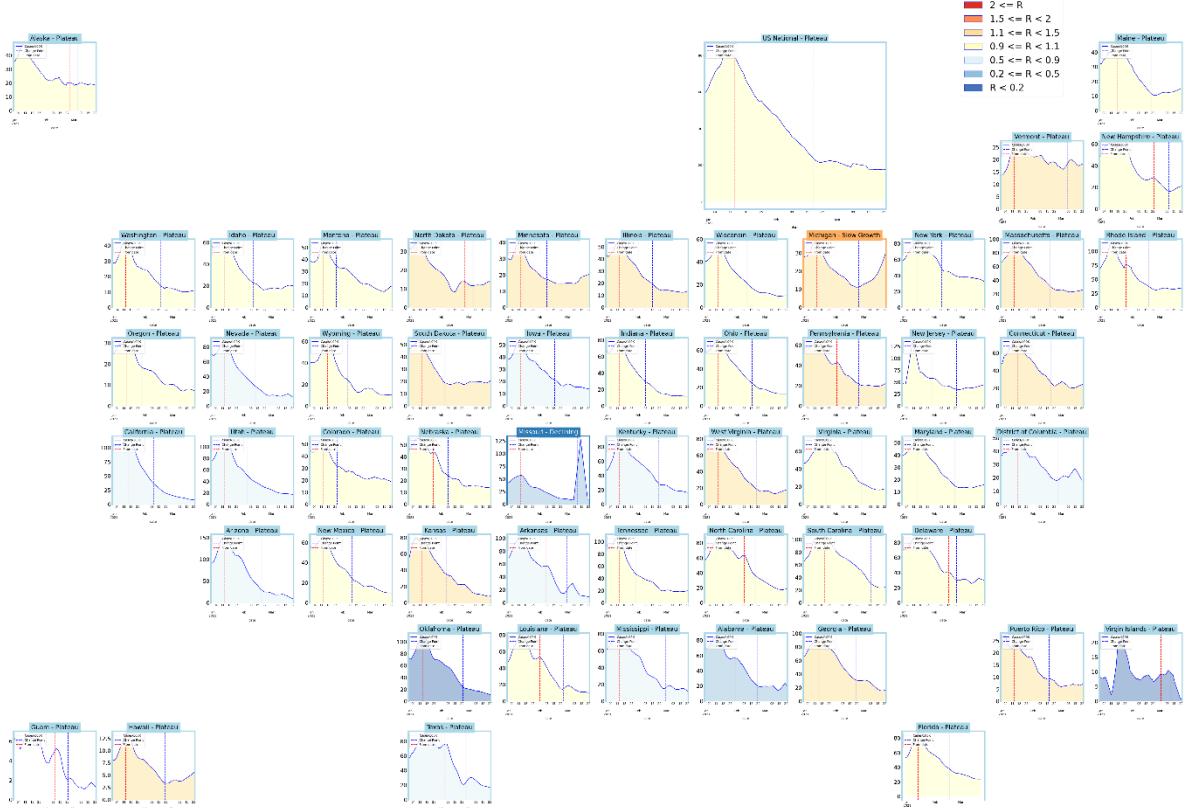
Case Rates are associate with Poverty levels



Total cumulative cases from zip codes with different levels of poverty by population. This association may be stronger at the census tract level.

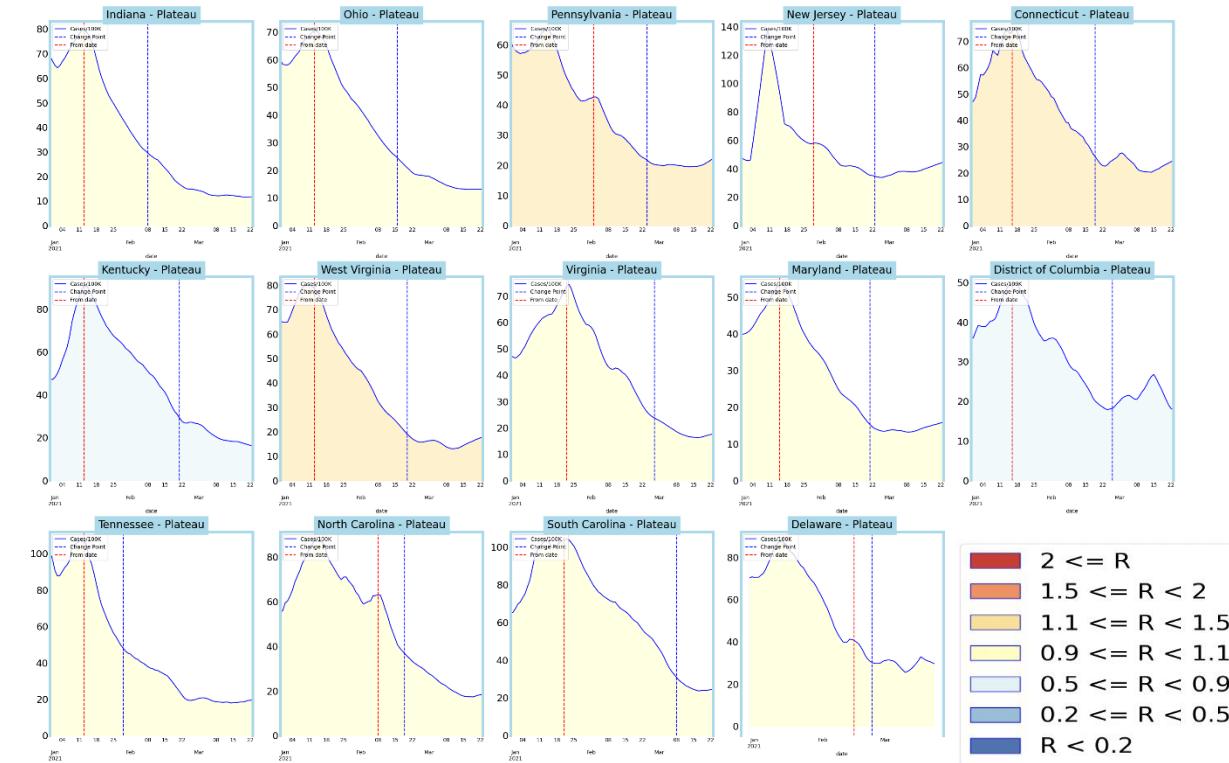
Other State Comparisons

Trajectories of States



- Nearly all states are plateaued, with slow growth in Michigan
- Missouri in decline only because of data artifact in reporting

Virginia and her neighbors

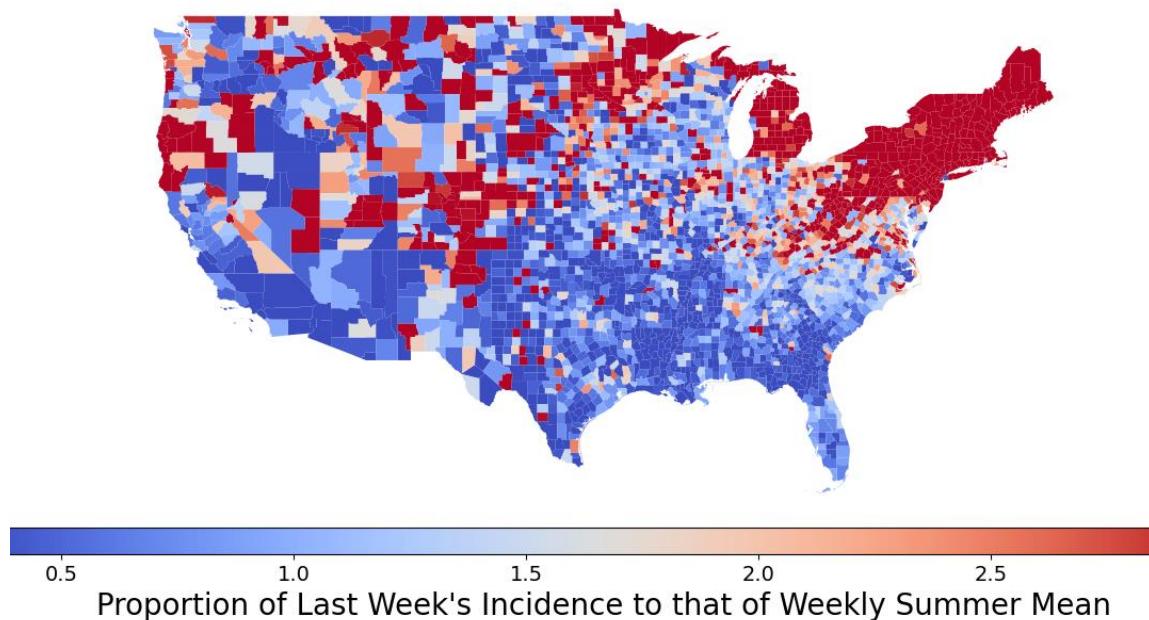


- VA and nearly all in plateau with slight upward trend
- Rates remain elevated, but significantly down from peaks in Jan

Current Week vs. Summer Mean (June-Aug 2020)

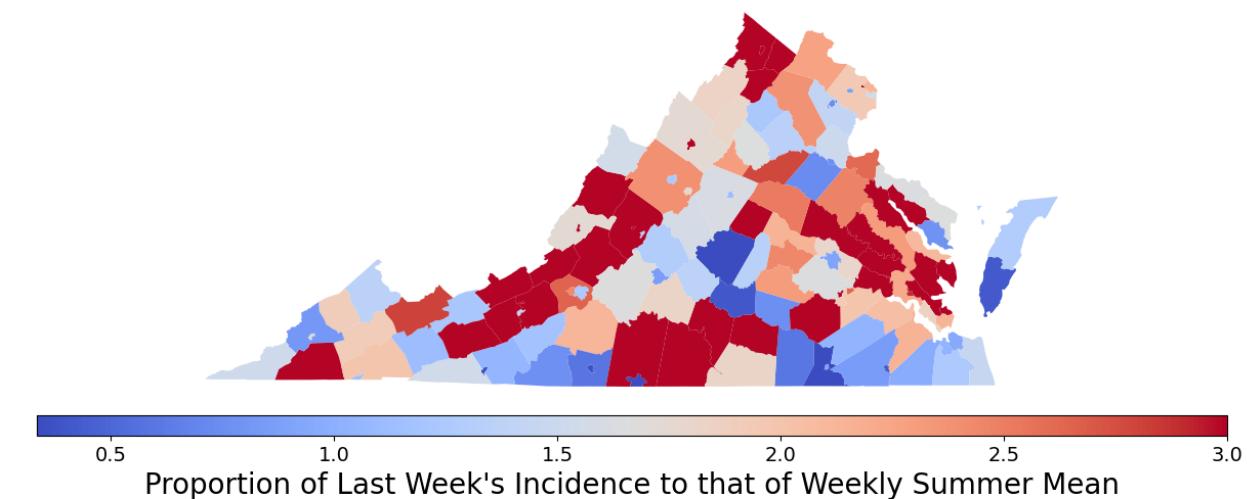
Still some way to go to return to rates experienced during the summer of 2020 (June through August)

Recent Incidence Compared to Weekly Summer Mean by County
Mean: 15.35; Median: 1.12; IQR: 0.49-2.49



- 53% of US counties are above the summer mean case rate compared to 59% last week

Recent Incidence Compared to Weekly Summer Mean by County
Mean: 2.35; Median: 1.82; IQR: 1.23-3.01



- 81% of VA counties are above the average rate for the summer compared to 82% last week

Zip code level weekly Case Rate (per 100K)

Case Rates in the last week by zip code

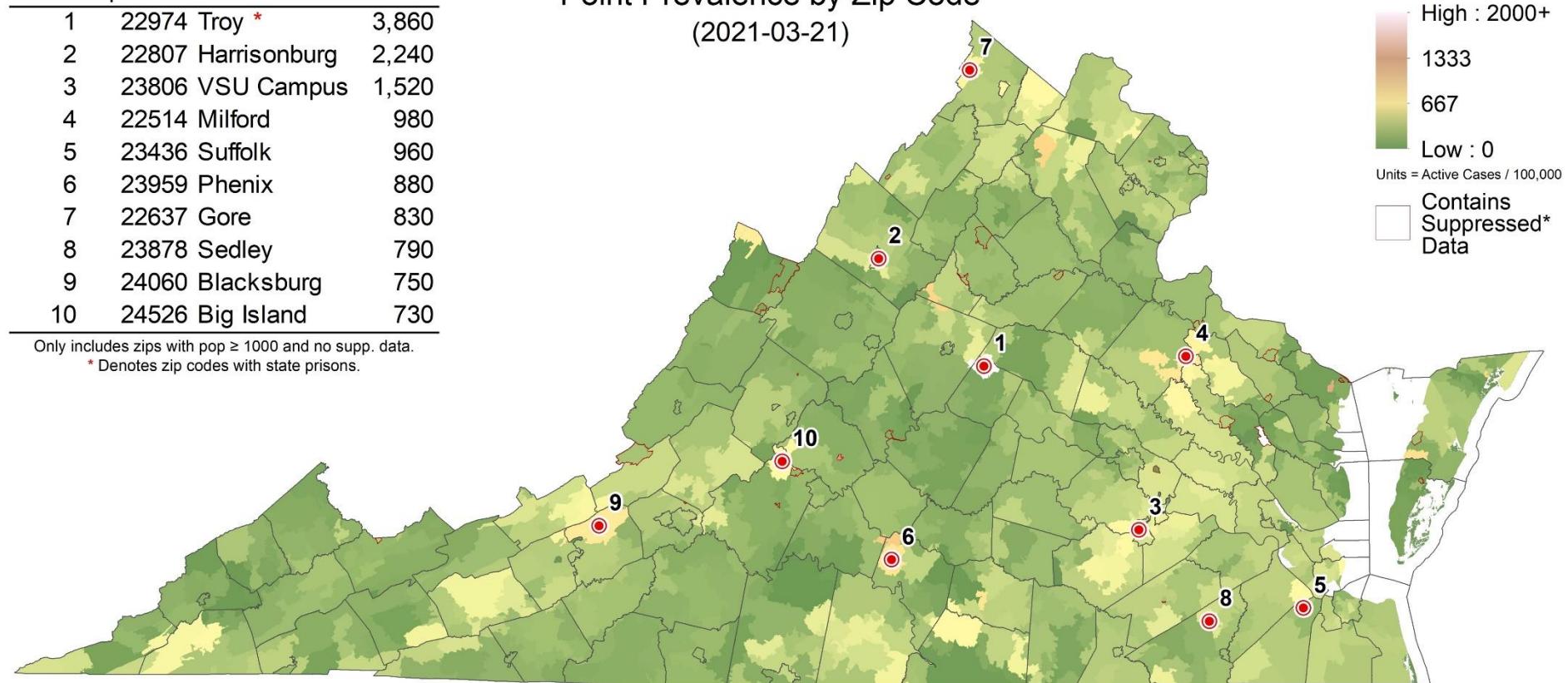
- Universities still dominate the top 10 list
- Concentrations of high rates scattered across the Commonwealth
- Some counts are low and suppressed to protect anonymity, those are shown in white

Rank	Zip Code Name	Prev
1	22974 Troy *	3,860
2	22807 Harrisonburg	2,240
3	23806 VSU Campus	1,520
4	22514 Milford	980
5	23436 Suffolk	960
6	23959 Phenix	880
7	22637 Gore	830
8	23878 Sedley	790
9	24060 Blacksburg	750
10	24526 Big Island	730

Only includes zips with pop ≥ 1000 and no supp. data.

* Denotes zip codes with state prisons.

Point Prevalence by Zip Code
(2021-03-21)



Based on Spatial Empirical Bayes smoothed point prevalence for week ending 2021-03-21.

Note: New color ramp scale and new ascertainment ratio of 2:1.



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Risk of Exposure by Group Size and HCW prevalence

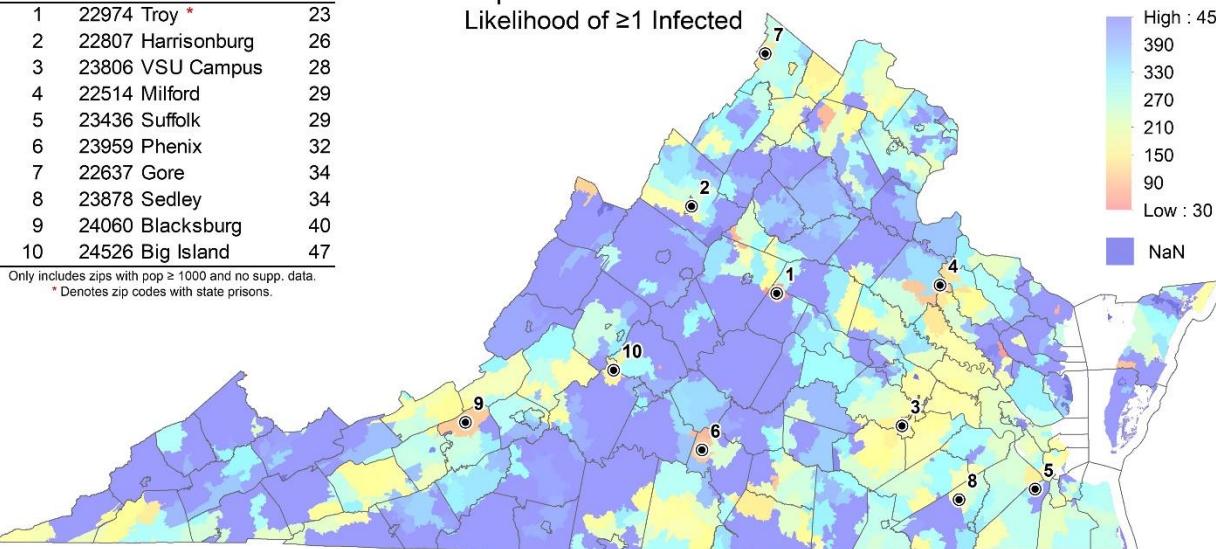
Case Prevalence in the last week by zip code used to calculate risk of encountering someone infected in a gathering of randomly selected people (group size 25)

- **Group Size:** Assumes 2 undetected infections per confirmed case (ascertainment rate from recent seroprevalence survey), and shows minimum size of a group with a 50% chance an individual is infected by zip code (eg in a group of 26 in Harrisonburg, there is a 50% chance someone will be infected)
- **HCW prevalence:** Case rate among health care workers (HCW) in the last week using patient facing health care workers as the denominator

Rank	Zip Code Name	Size
1	22974 Troy *	23
2	22807 Harrisonburg	26
3	23806 VSU Campus	28
4	22514 Milford	29
5	23436 Suffolk	29
6	23959 Phenix	32
7	22637 Gore	34
8	23878 Sedley	34
9	24060 Blacksburg	40
10	24526 Big Island	47

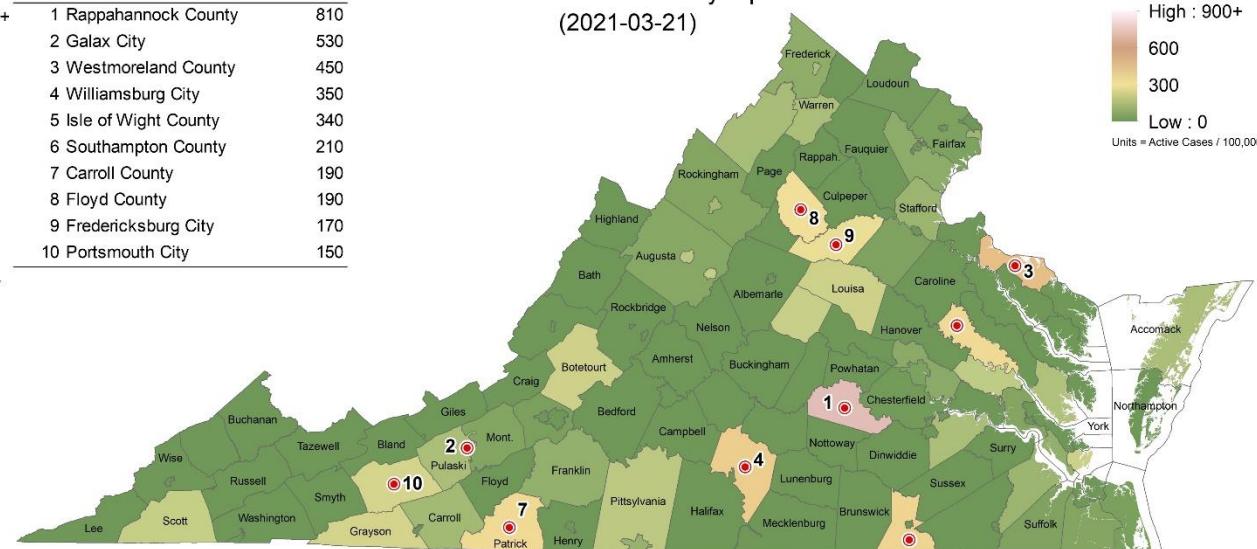
Only includes zips with pop ≥ 1000 and no supp. data.
* Denotes zip codes with state prisons.

Group Size Needed for 50% Likelihood of ≥ 1 Infected



Group Size	Rank	Name	Prevalence
High : 450+	1	Rappahannock County	810
390	2	Galax City	530
330	3	Westmoreland County	450
270	4	Williamsburg City	350
210	5	Isle of Wight County	340
150	6	Southampton County	210
90	7	Carroll County	190
Low : 30	8	Floyd County	190
	9	Fredericksburg City	170
	10	Portsmouth City	150
NaN			

HCW Point Prevalence by Zip Code
(2021-03-21)

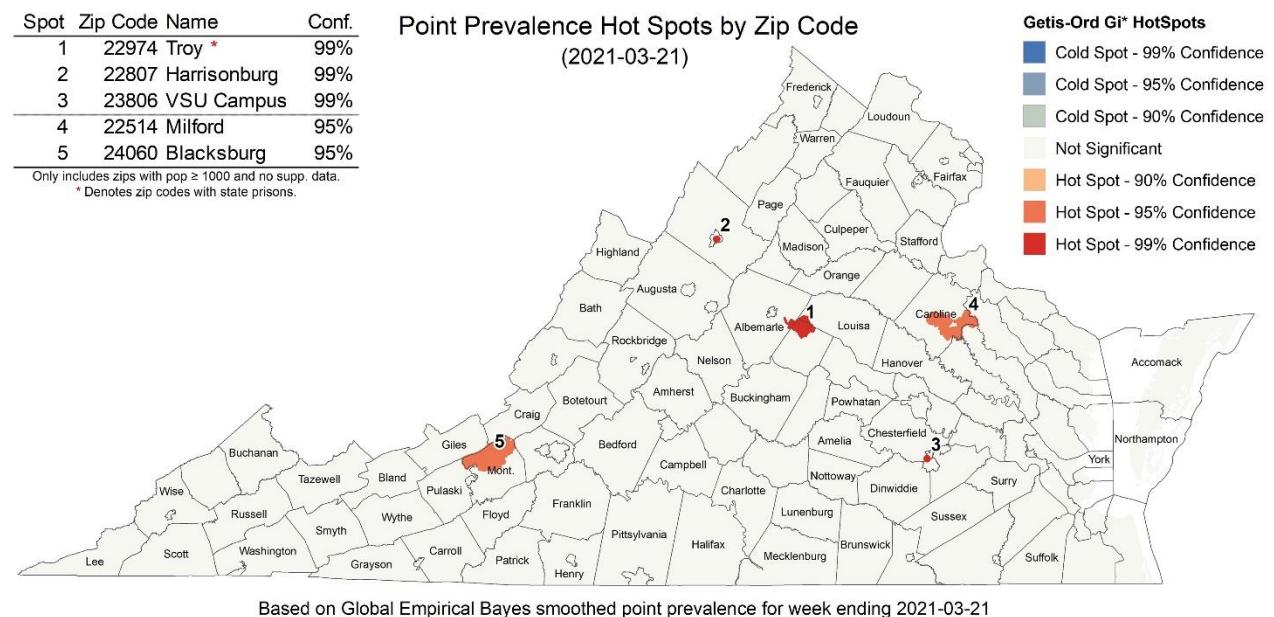


Current Hot-Spots

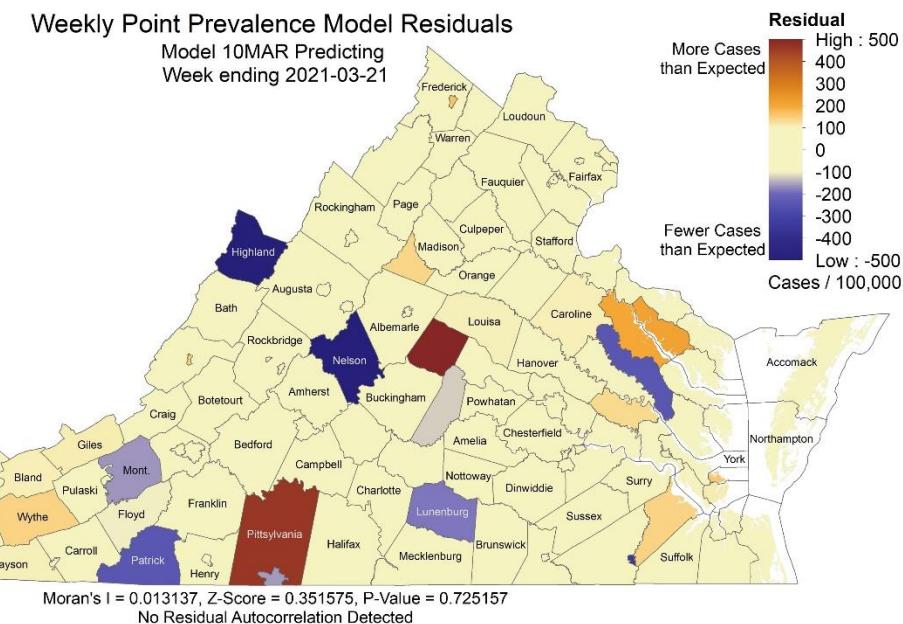
Case rates that are significantly different from neighboring areas or model projections

- **Spatial:** Hot Spots compare the weekly case prevalence to nearby zip codes to identify areas with statistically significant deviations
- **Temporal:** The weekly case rate (per 100K) projected last week compared to observed by county, which highlights temporal fluctuations that differ from the model's projections

Spatial Hotspots



Temporal Hotspots



Model Update – Adaptive Fitting



Adaptive Fitting Approach

Each county fit precisely, with recent trends used for future projection

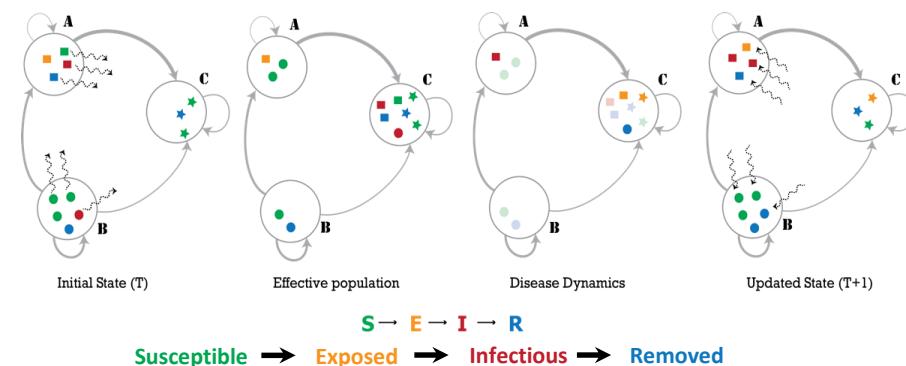
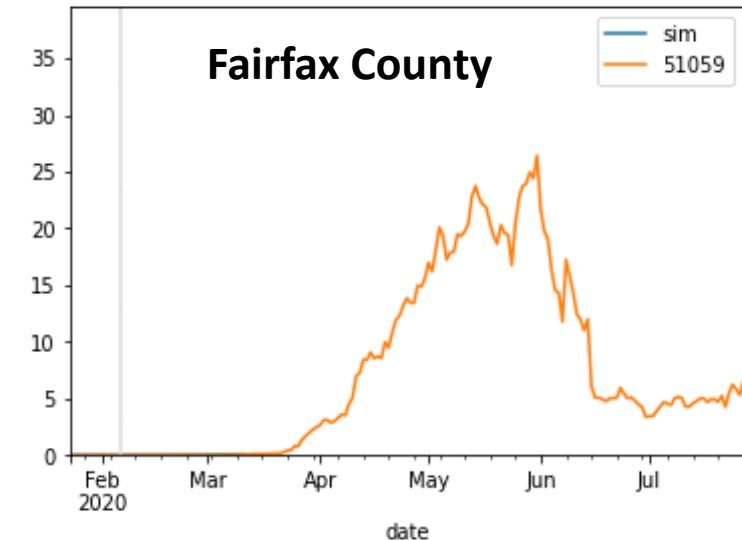
- Allows history to be precisely captured, and used to guide bounds on projections

Model: An alternative use of the same meta-population model, PatchSim

- Allows for future “what-if” Scenarios to be layered on top of calibrated model
- Eliminates connectivity between patches, to allow calibration to capture the increasingly unsynchronized epidemic

External Seeding: Steady low-level importation

- Widespread pandemic eliminates sensitivity to initial conditions
- Uses steady 1 case per 10M population per day external seeding



Using Ensemble Model to Guide Projections

Ensemble methodology that combines the Adaptive with machine learning and statistical models such as:

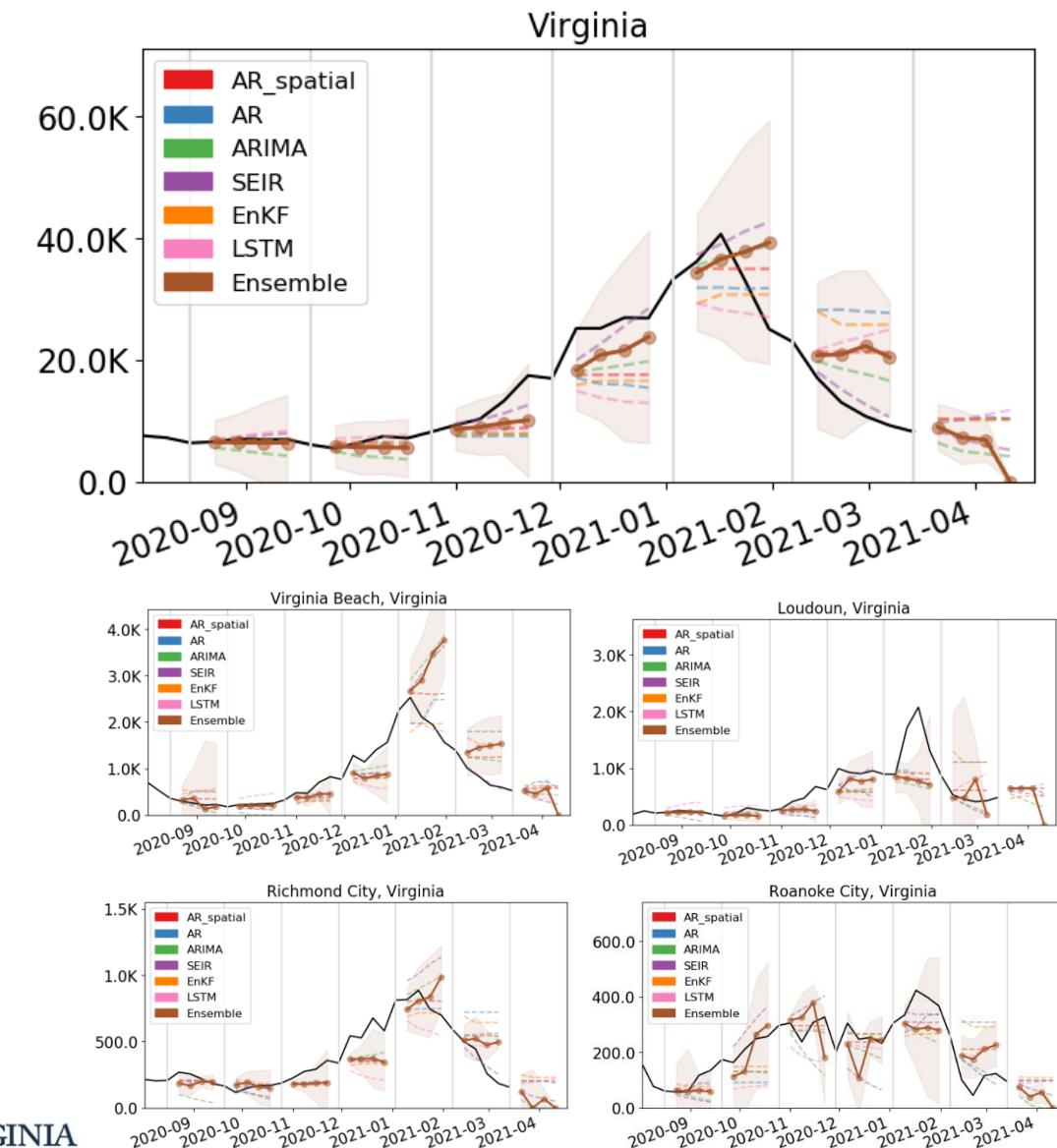
- Autoregressive (AR, ARIMA)
- Neural networks (LSTM)
- Kalman filtering (EnKF)

Weekly forecasts done at county level.

Models chosen because of their track record in disease forecasting and to increase diversity and robustness.

Ensemble forecast provides additional ‘surveillance’ for making scenario-based projections.

Also submitted to CDC Forecast Hub.



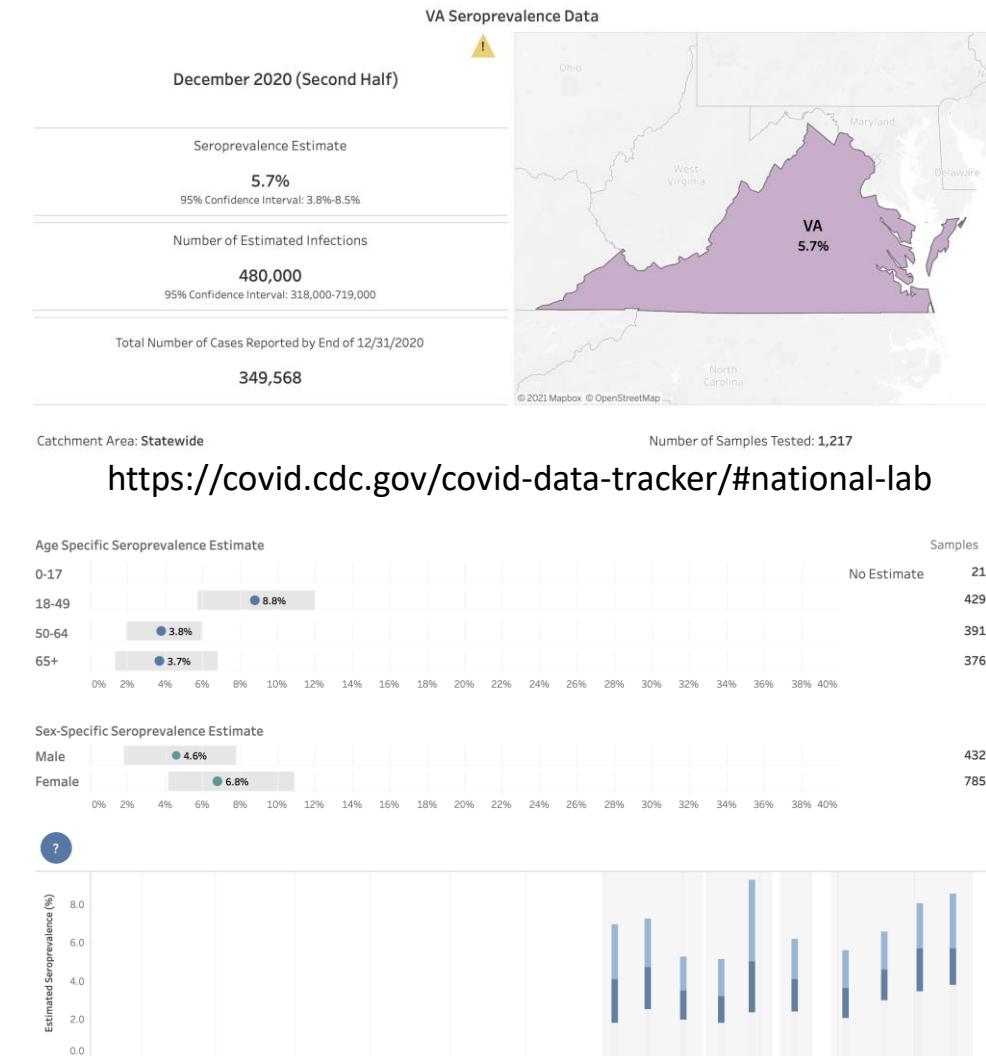
Seroprevalence updates to model design

Several seroprevalence studies provide better picture of how many actual infections have occurred

- CDC Nationwide Commercial Laboratory Seroprevalence Survey estimated 7.6% [5.6% – 9.8%] seroprevalence as of Jan 7th – 21st up from 5.7% a month earlier

These findings are equivalent to an ascertainment ratio of ~2x in the future, with bounds of (1.3x to 3x)

- Thus for 2x there are 2 total infections in the population for every confirmed case recently
- This measure now fully tracks the estimated ascertainment over time
- Uncertainty design has been shifted to these bounds (previously higher ascensions as was consistent earlier in the pandemic were being used)



Calibration Approach

- **Data:**
 - County level case counts by date of onset (from VDH)
 - Confirmed cases for model fitting
- **Calibration:** fit model to observed data and ensemble's forecast
 - Tune transmissibility across ranges of:
 - Duration of incubation (5-9 days), infectiousness (3-7 days)
 - Undocumented case rate (1x to 7x) guided by seroprevalence studies
 - Detection delay: exposure to confirmation (4-12 days)
 - Approach captures uncertainty, but allows model to precisely track the full trajectory of the outbreak
- **Project:** future cases and outcomes generated using the collection of fit models run into the future
 - **Mean trend from last 7 days of observed cases and first week of ensemble's forecast used**
 - Outliers removed based on variances in the previous 3 weeks
 - 2 week interpolation to smooth transitions in rapidly changing trajectories

COVID-19 in Virginia:

Cases, Hospitalizations and Deaths									
Total Cases* 608,704		Total Hospitalizations** 26,037		Total Deaths 10,143					
(New Cases: 1,470) [▲]		Confirmed† 476,936	Probable† 131,768	Confirmed† 24,712	Probable† 1,325				
Outbreaks									
Total Outbreaks*		Outbreak Associated Cases							
2,808		68,260							
* At least two (2) lab confirmed cases are required to classify an outbreak.									
Testing (PCR Only)									
Testing Encounters PCR Only*			Current 7-Day Positivity Rate PCR Only**						
6,356,719			5.6%						
Multisystem Inflammatory Syndrome in Children									
Total Cases*		Total Deaths							
48		0							

* Cases defined by CDC HAN case definition: <https://emergency.cdc.gov/han/2020/han00432.asp>

** Lab reports may not have been received yet. Percent positivity is not calculated for days with incomplete data.

Accessed 9:00am March 24, 2021

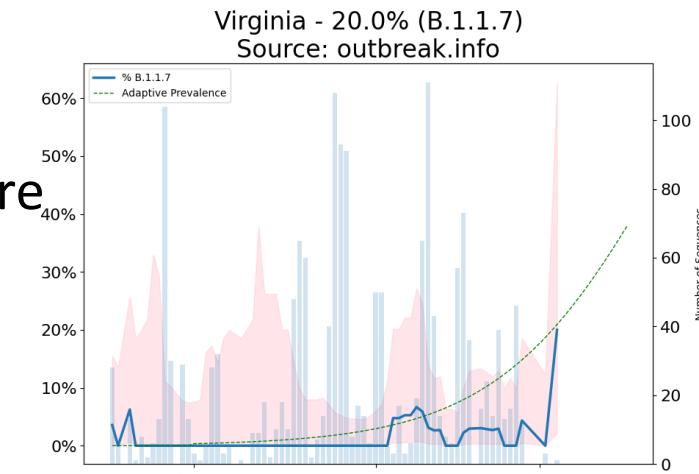
<https://www.vdh.virginia.gov/coronavirus/>

Scenarios – Seasonal Effects

- Variety of factors continue to drive transmission rates
 - Seasonal impact of weather patterns, travel and gatherings, fatigue and premature relaxation of infection control practices
- Plausible levels of transmission can be bounded by past experience
 - Assess transmission levels at the county level since May 1, 2020 through September 30, 2020
 - Use the highest and lowest levels experienced (excluding outliers) as plausible bounds for levels of control achievable
 - Transition from current levels of projection to the new levels over 2 months
- Projection Scenario:
 - **Fatigued Control:** Highest level of transmission (95th percentile) increased by additional 5%

Scenarios – Novel Variants

- Several novel variants of SARS-CoV2 are being tracked
 - Some are more transmissible, some may escape immunity from previous natural infection and/or vaccination, others may be more severe
- New Variant B.1.1.7 is best understood and is in Virginia
 - **Transmission increase:** [Several different studies](#) have estimated the increase in transmission to be 30-55%, we use 50% increase from the current baseline projection
 - **Emergence timing:** Gradual frequency increase reaching 50% frequency March 30th, one week after the national estimate in [MMWR report from CDC](#) and refined by [Andersen et al.](#)
- Variant planning Scenario:
 - **VariantB117:** Current projected transmissibility continues to increase through June to a level 50% more transmissible



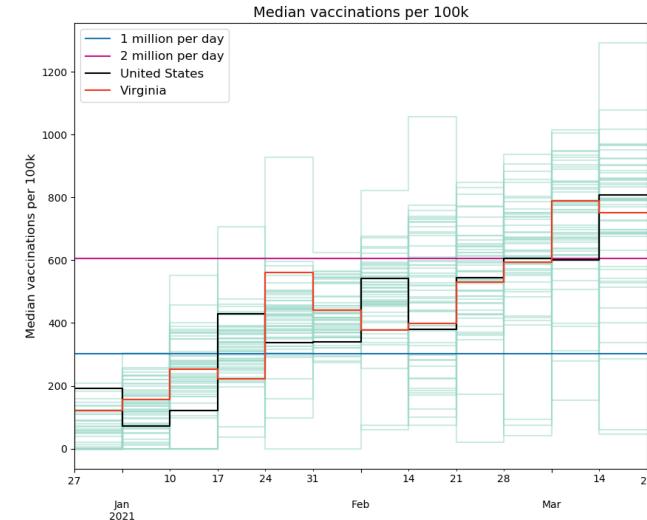
Estimated frequency from public genome repository with added analysis: 35%
Current frequency used in model: 35%



Scenarios – Vaccines

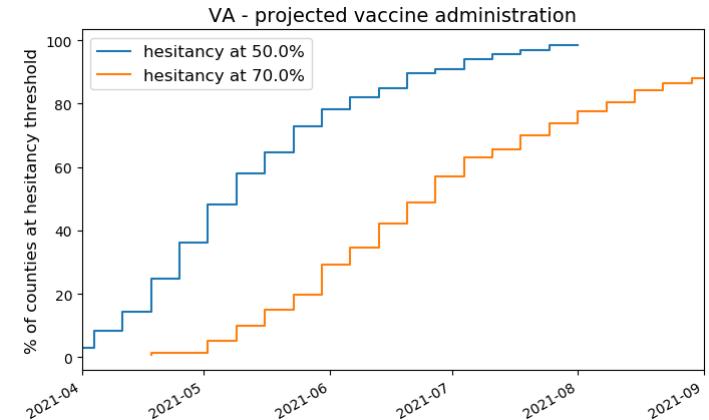
- Vaccination is well underway and accelerating its pace
- Vaccine efficacy varies over course of vaccine
 - FDA EUAs show 50% efficacy achieved 2 weeks after 1st dose, and 95% 2 weeks after 2nd dose
 - Assuming 3.5 week (average of Pfizer and Moderna) gap between doses
 - Johnson & Johnson included with 67% efficacy 2 weeks after 1st (and only) dose
- Accelerated administration pace will reach vaccine hesitancy thresholds more quickly
 - Demand still outpaces supply in many regions
 - Estimate based on current rates that some counties may reach thresholds as soon as early April, with potentially half by late June (increased rates have moved these thresholds sooner)

VA Vaccination Rates



Lines represent 1M & 2M total doses
administered a day (rate of 303/100K & 606/100K)

Anticipated Vax Hesitancy Impact

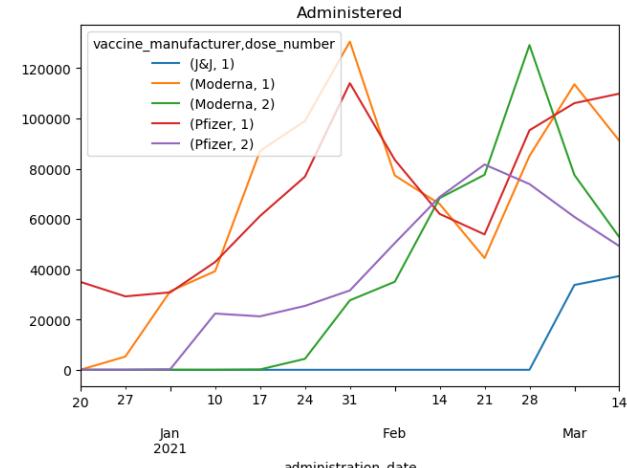


Scenarios – Vaccines

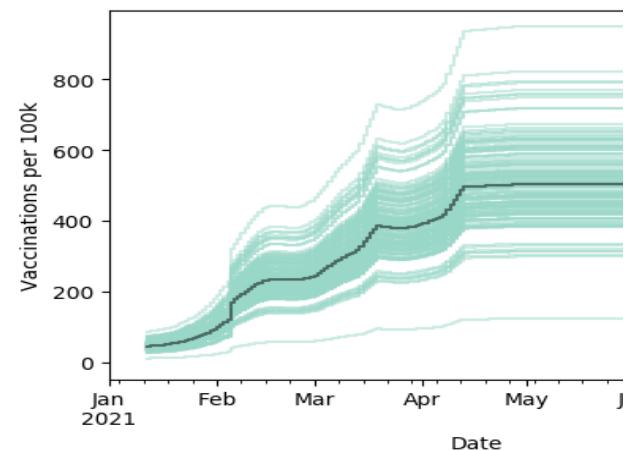
- Administration schedule uses reported administrations and anticipated supplies to generate vaccine schedule (past and future)
 - Data from [VDH](#) used to assess county level variations and dosage (these data are in data package)
- Current administration rate used as baseline
Rate: 573 FIRST DOSES per 100K per day
 - Total of ~50K 1st doses / day (some are J&J thus only dose)
 - **Total Administrations:** This pace leads to eventually reaching ~87K administered a day, implying 50K become fully vaccinated a day
 - **Location:** Per capita distribution across all counties

Current rollouts and scenarios inspired by MIDAS Network COVID-19 Scenario Hub: <https://github.com/midas-network/covid19-scenario-modeling-hub>

Weekly dose administrations



Modeled Vaccine Induced Immunity



All VA counties, state in black

Scenarios – Seasonal Effects and Vaccines

Three scenarios combine these seasonal effects and use the current vaccine schedule

- **Adaptive:** No seasonal effects from base projection
 - If things continue as they are
- **Adaptive-FatigueControl:** Fatigued control seasonal effects
 - If we revert to slightly worst transmission experienced in last 6 months
- **Adaptive-VariantB117:** Boosting of transmissibility from the emergence of B.1.1.7
 - If new variants begin to predominate and boost transmission, this assumes current seasonal affects remain the same (eg like Adaptive)
- **Adaptive-FatigueControl-VariantB117:** Fatigued control and txm boost from B.1.1.7

Counterfactuals with no vaccine (“NoVax”) are provided for comparison purposes

Model Results



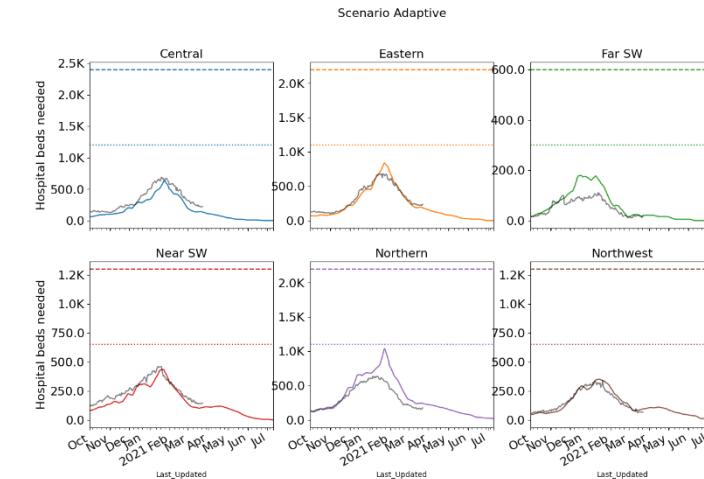
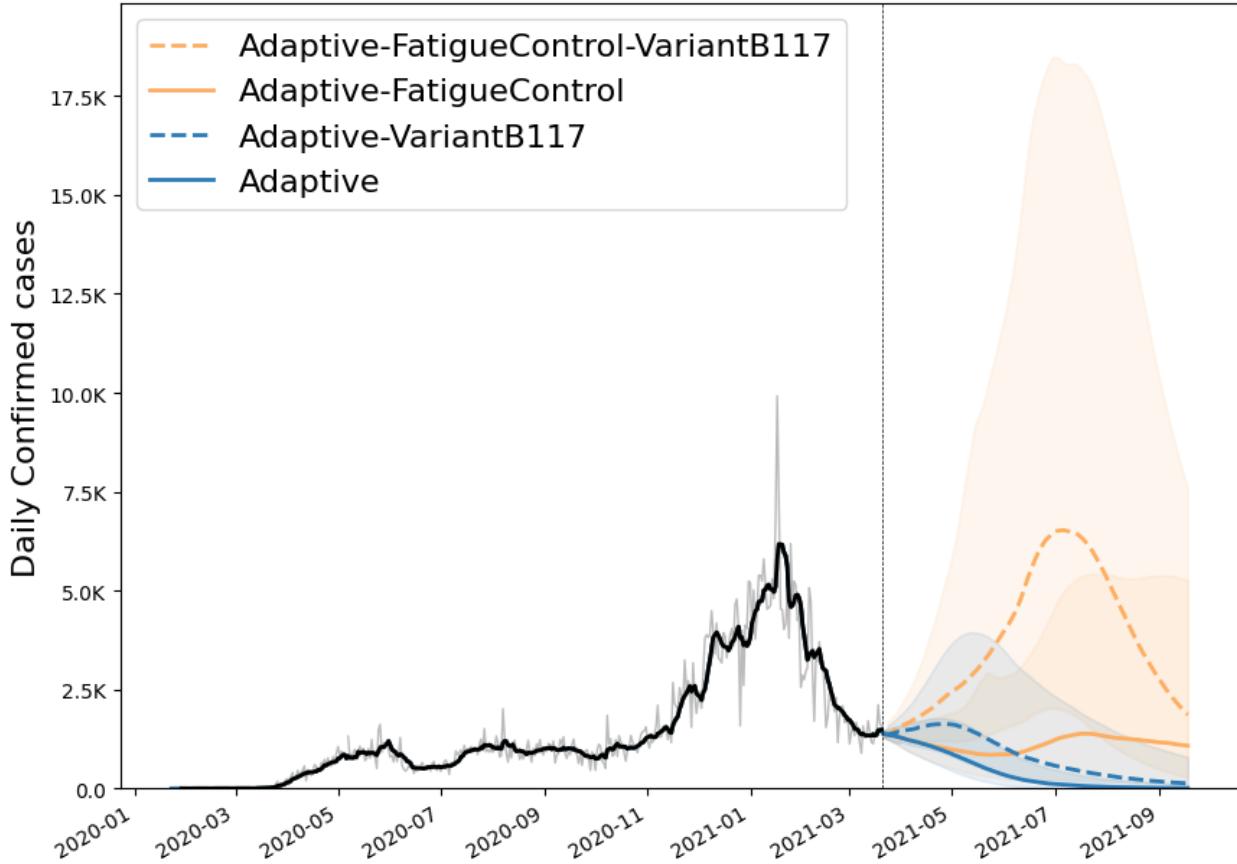
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Outcome Projections

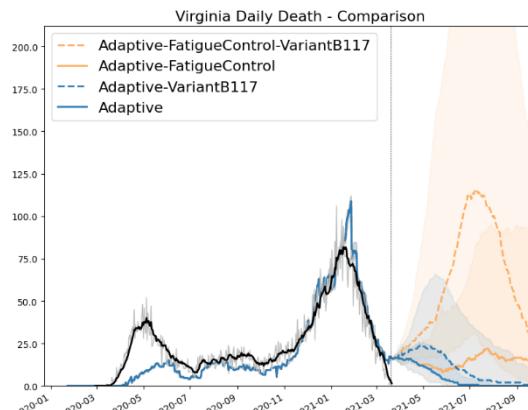
Estimated Hospital Occupancy

Confirmed cases

Virginia Daily Confirmed - Comparison

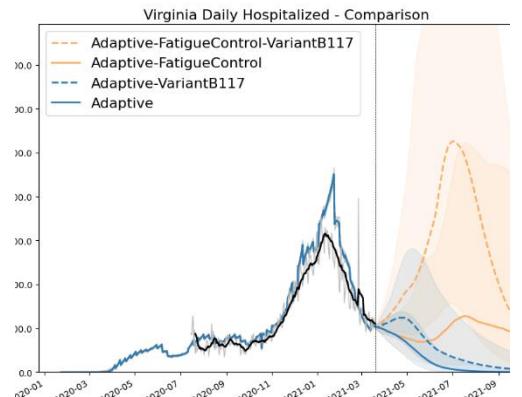


Daily Deaths



Death ground truth from VDH "Event Date" data, most recent dates are not complete

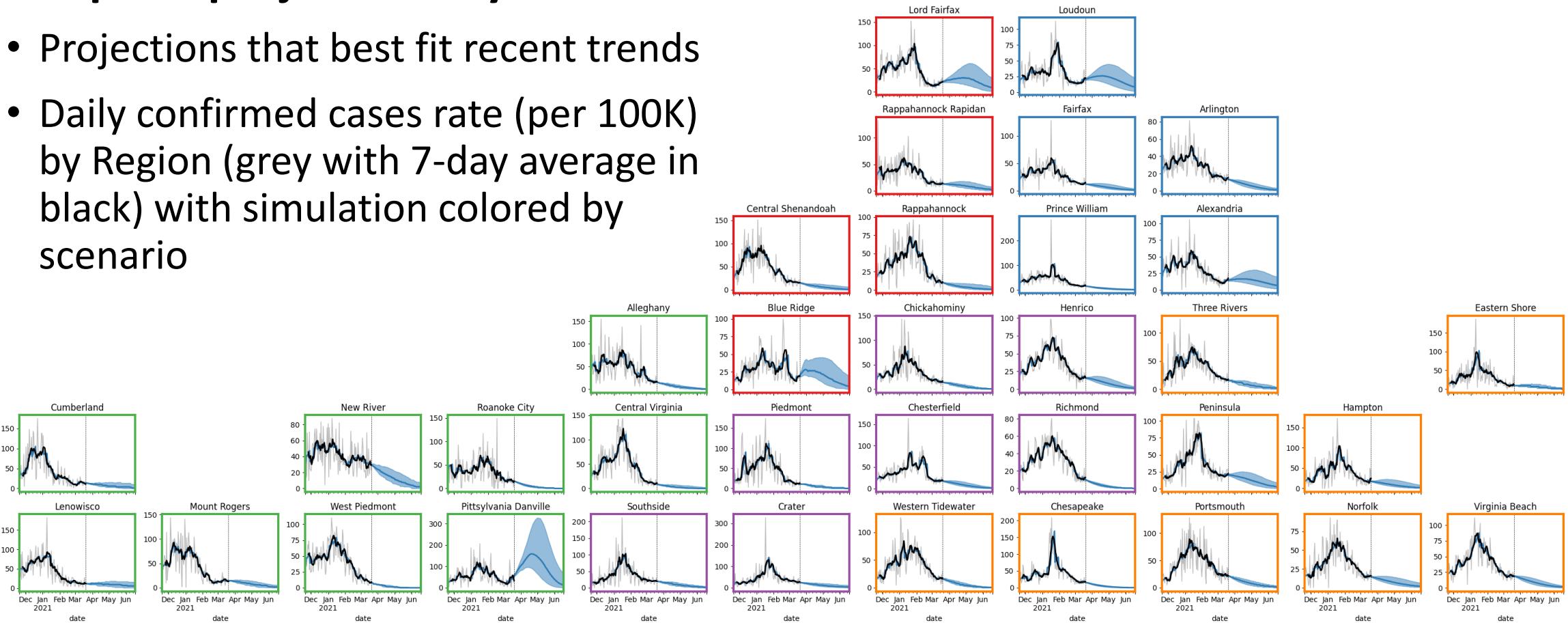
Daily Hospitalized



District Level Projections: Adaptive

Adaptive projections by District

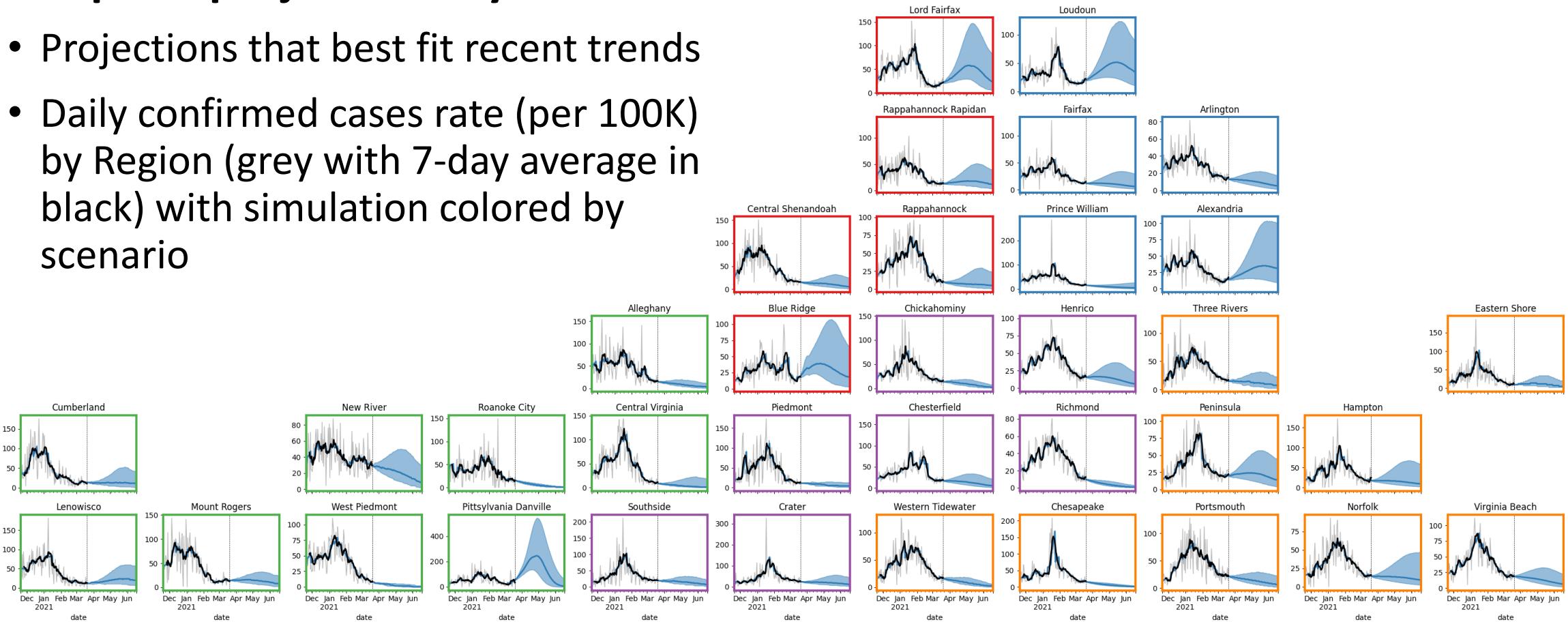
- Projections that best fit recent trends
- Daily confirmed cases rate (per 100K) by Region (grey with 7-day average in black) with simulation colored by scenario



District Level Projections: Adaptive-VariantB117

Adaptive projections by District

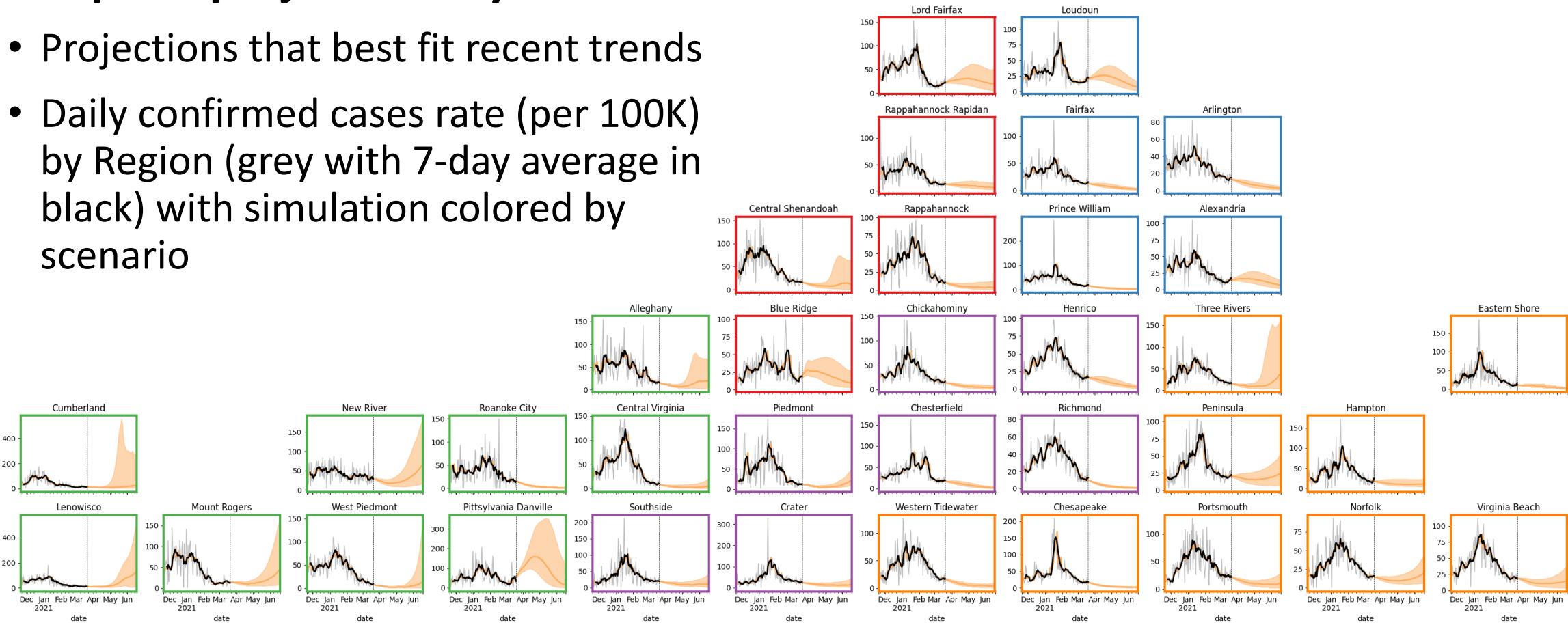
- Projections that best fit recent trends
- Daily confirmed cases rate (per 100K) by Region (grey with 7-day average in black) with simulation colored by scenario



District Level Projections: Adaptive-FatigueControl

Adaptive projections by District

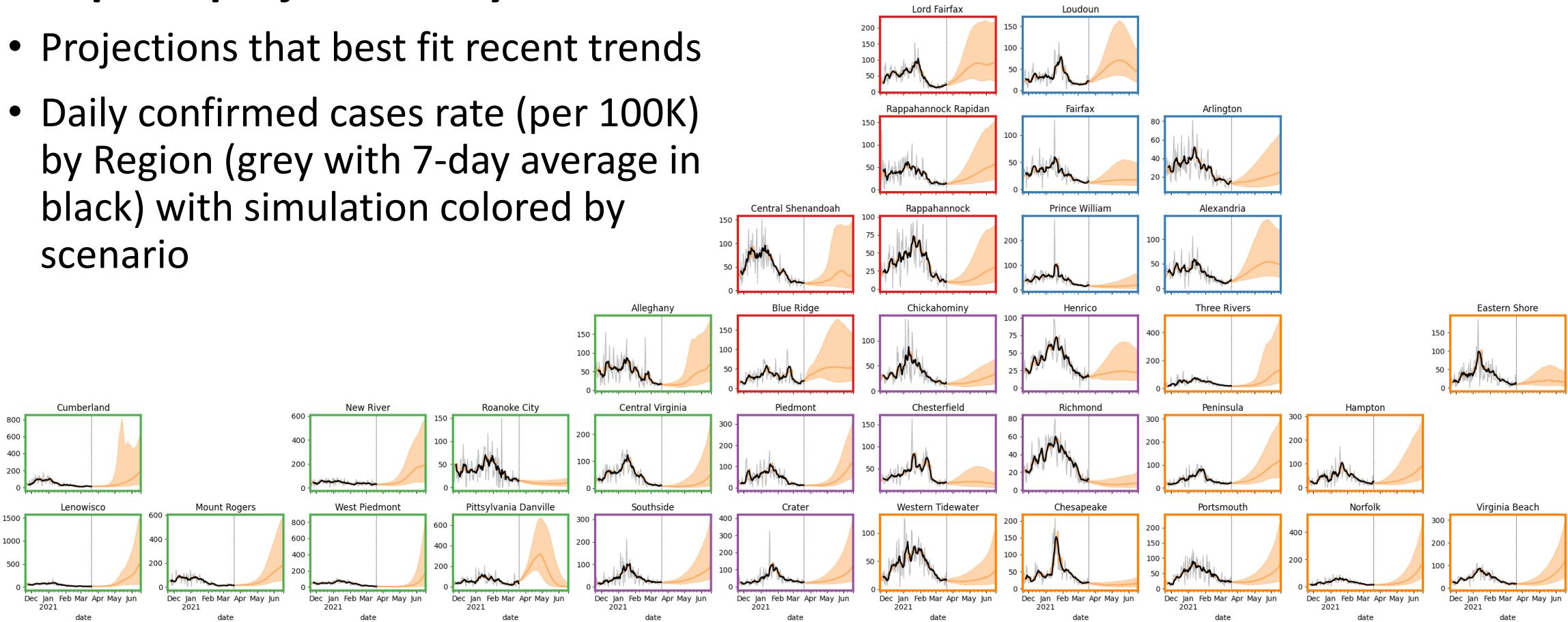
- Projections that best fit recent trends
- Daily confirmed cases rate (per 100K) by Region (grey with 7-day average in black) with simulation colored by scenario



District Level Projections: Adaptive-FatigueControl-VariantB117

Adaptive projections by District

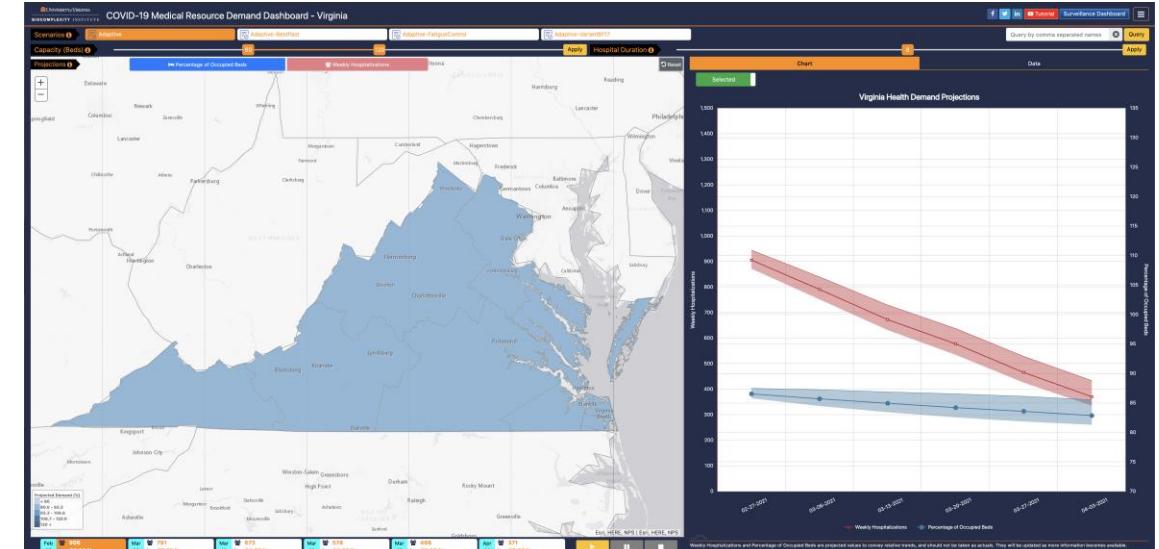
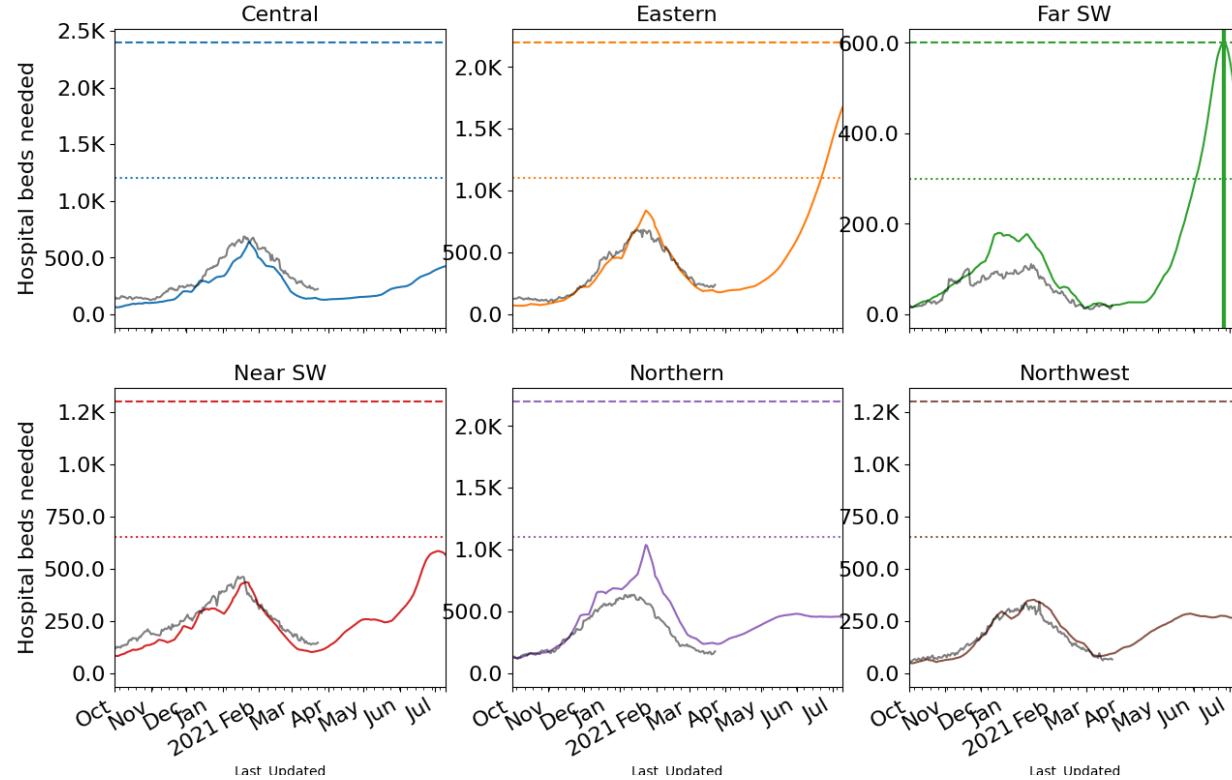
- Projections that best fit recent trends
- Daily confirmed cases rate (per 100K) by Region (grey with 7-day average in black) with simulation colored by scenario



Hospital Demand and Bed Capacity by Region

Capacities* by Region – Adaptive-FatigueControl-VariantB117

COVID-19 capacity ranges from 80% (dots) to 120% (dash) of total beds



<https://nssac.bii.virginia.edu/covid-19/vmrddash/>

If Adaptive-FatigueControl-VariantB117 scenario:

- Far SW may reach surge bed capacity in late June
- Eastern, Near SW approach initial bed capacity in June as well

* Assumes average length of stay of 8 days

Weekly Cases and Hospitalizations

Weekly confirmed cases

Week Ending	Adaptive	Adaptive-Fatigued Control	Adaptive-VariantB117	Adaptive-Fatigued Control -VariantB117
3/21/21	9,925	9,925	9,925	9,925
3/28/21	9,420	9,428	9,667	9,819
4/4/21	8,931	8,950	10,159	11,075
4/11/21	8,361	8,384	10,549	12,127
4/18/21	7,783	7,844	11,041	13,416
4/25/21	7,090	7,397	11,362	14,976
5/2/21	6,314	7,008	11,375	16,711
5/9/21	5,453	6,627	10,911	18,229
5/16/21	4,577	6,273	10,106	20,162
5/23/21	3,694	6,032	9,084	22,626
5/30/21	2,907	6,002	7,969	25,655
6/6/21	2,221	6,074	6,887	28,891

Weekly Hospitalizations

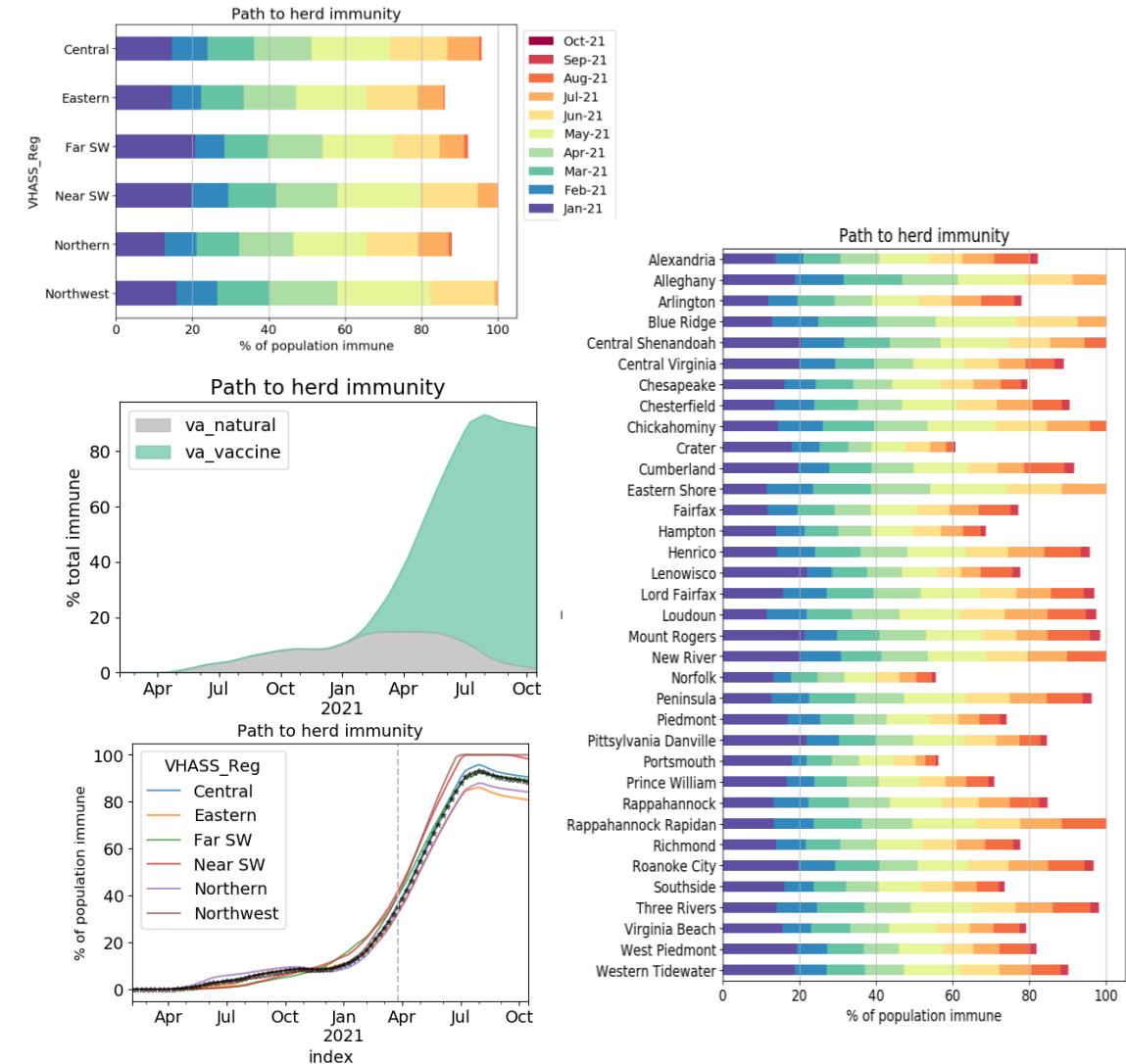
Week Ending	Adaptive	Adaptive-Fatigued Control	Adaptive-VariantB117	Adaptive-Fatigued Control -VariantB117
3/21/21	693	693	693	693
3/28/21	659	659	680	695
4/4/21	615	616	725	790
4/11/21	580	584	749	857
4/18/21	540	546	780	955
4/25/21	481	503	813	1,067
5/2/21	409	463	813	1,194
5/9/21	342	427	780	1,308
5/16/21	280	395	705	1,454
5/23/21	218	375	622	1,658
5/30/21	179	381	550	1,869
6/6/21	140	403	470	2,082



Virginia's Progress on Population Immunity

Natural Immunity and Vaccines combine to produce a population level of immunity

- Population level immunities above 75% (assuming even distribution in the population) will be effective for preventing significant outbreaks of COVID-19
- How long immunity from infection with SARS-CoV2 lasts is not well understood but may vary based on severity of symptoms
 - We assume a conservative 6 month period of protection for these calculations
- Vaccine induced immunity is likely to last longer, we assume indefinite protection
 - This also assumes that all administered vaccines remain protective against current and future novel variants
- Population immunity depends on a very high proportion of the population getting vaccinated
 - We assume 90% of adults will ultimately get vaccinated in these calculations but slow rates may prevent this from happening before October 2021



Key Takeaways

Projecting future cases precisely is impossible and unnecessary.

Even without perfect projections, we can confidently draw conclusions:

- **Case rates in Virginia have flattened and now have some growth**
- VA mean weekly incidence slightly up to 17/100K from 15/100K, US flat (to 16.5 from 16 per 100K)
- Significant progress made in last month, however 81% of VA counties above mean rate of Summer 2020
- Projections are flattening out across Commonwealth, with growth on the horizon boosted by B.1.1.7
- Recent updates:
 - Slowed ramp up for new variant scenario given lack of clear evidence it has reached predominance this week
 - Adjusted Seasonal Effects scenarios to account for spring and summer weather
 - Accelerated vaccine schedule with Johnson & Johnson added as base case in anticipation of boost in vaccine supplies
- The situation continues to change. Models continue to be updated regularly.



References

Venkatramanan, S., et al. "Optimizing spatial allocation of seasonal influenza vaccine under temporal constraints." *PLoS computational biology* 15.9 (2019): e1007111.

Arindam Fadikar, Dave Higdon, Jiangzhuo Chen, Bryan Lewis, Srinivasan Venkatramanan, and Madhav Marathe. Calibrating a stochastic, agent-based model using quantile-based emulation. *SIAM/ASA Journal on Uncertainty Quantification*, 6(4):1685–1706, 2018.

Adiga, Aniruddha, Srinivasan Venkatramanan, Akhil Peddireddy, et al. "Evaluating the impact of international airline suspensions on COVID-19 direct importation risk." *medRxiv* (2020)

NSSAC. PatchSim: Code for simulating the metapopulation SEIR model. <https://github.com/NSSAC/PatchSim> (Accessed on 04/10/2020).

Virginia Department of Health. COVID-19 in Virginia. <http://www.vdh.virginia.gov/coronavirus/> (Accessed on 04/10/2020)

Biocomplexity Institute. COVID-19 Surveillance Dashboard. <https://nssac.bii.virginia.edu/covid-19/dashboard/>

Google. COVID-19 community mobility reports. <https://www.google.com/covid19/mobility/>

Biocomplexity page for data and other resources related to COVID-19: <https://covid19.biocomplexity.virginia.edu/>



Questions?

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Supplemental Slides



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Date of Onset Reproductive Number

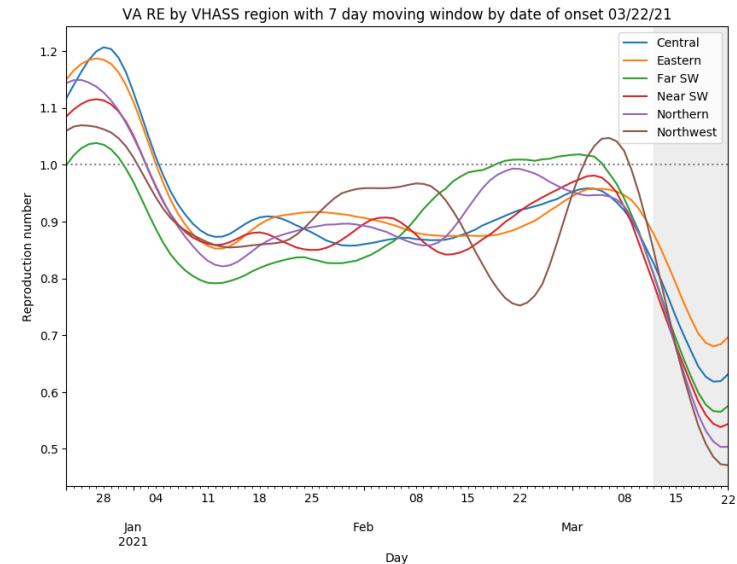
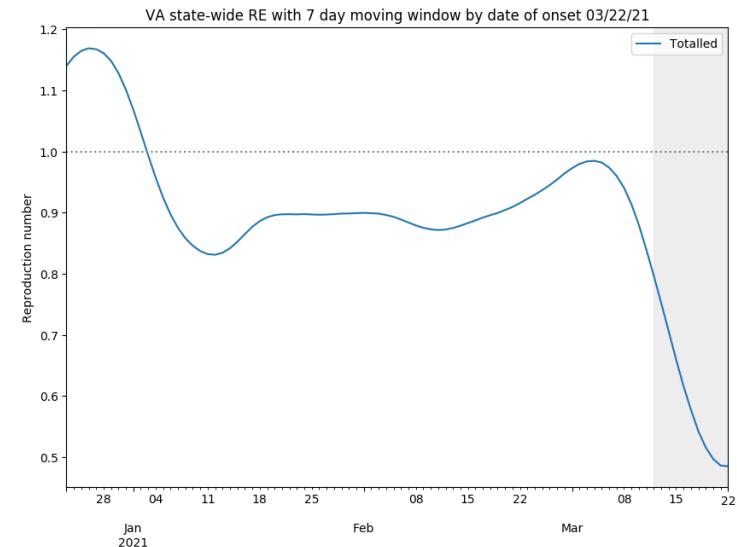
Mar 13th Estimates

Region	Date of Onset	Date Onset Diff
	R _e	Last Week
State-wide	0.752	-0.086
Central	0.797	-0.058
Eastern	0.851	0.040
Far SW	0.768	-0.151
Near SW	0.752	-0.174
Northern	0.765	-0.079
Northwest	0.791	0.049

Methodology

- Wallinga-Teunis method (EpiEstim¹) for cases by confirmation date
- Serial interval: updated to discrete distribution from observations (mean=4.3, Flaxman et al, Nature 2020)
- Using Confirmation date since due to increasingly unstable estimates from onset date due to backfill

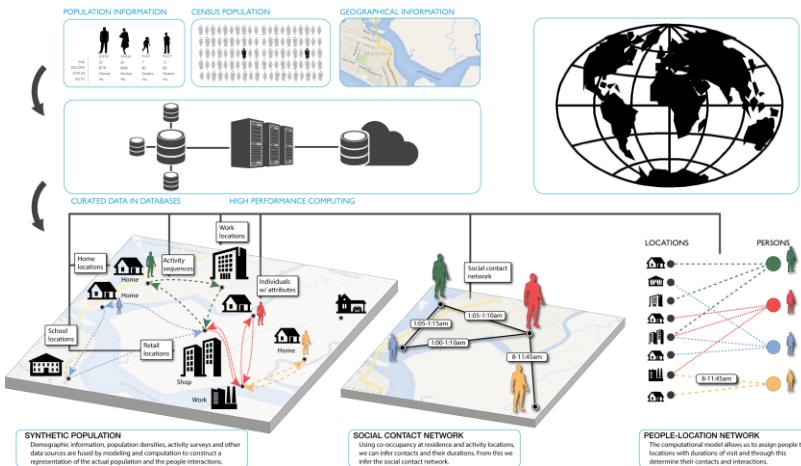
1. Anne Cori, Neil M. Ferguson, Christophe Fraser, Simon Cauchemez. A New Framework and Software to Estimate Time-Varying Reproduction Numbers During Epidemics. American Journal of Epidemiology, Volume 178, Issue 9, 1 November 2013, Pages 1505–1512, <https://doi.org/10.1093/aje/kwt133>



Agent-based Model (ABM)

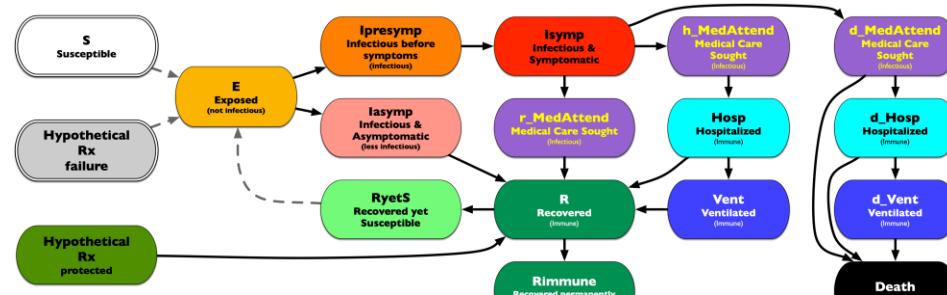
EpiHiper: Distributed network-based stochastic disease transmission simulations

- Assess the impact on transmission under different conditions
- Assess the impacts of contact tracing



Synthetic Population

- Census derived age and household structure
- Time-Use survey driven activities at appropriate locations



Detailed Disease Course of COVID-19

- Literature based probabilities of outcomes with appropriate delays
- Varying levels of infectiousness
- Hypothetical treatments for future developments

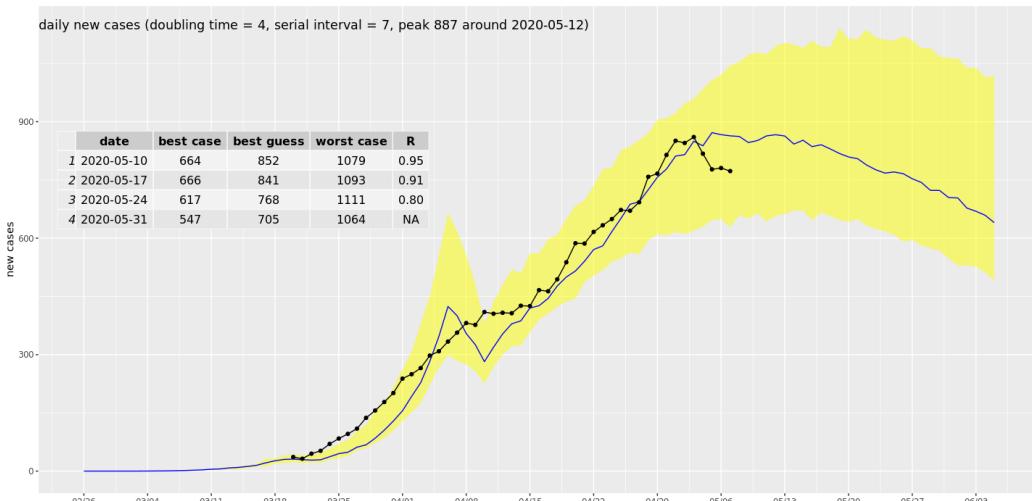


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ABM Social Distancing Rebound Study Design

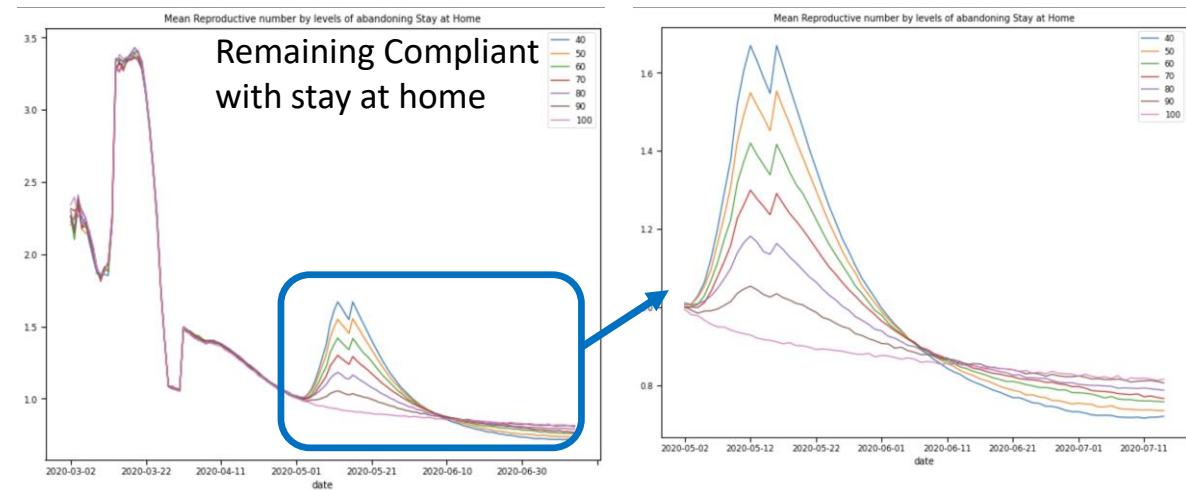
Study of "Stay Home" policy adherence

- Calibration to current state in epidemic
- Implement “release” of different proportions of people from “staying at home”



Calibration to Current State

- Adjust transmission and adherence to current policies to current observations
- For Virginia, with same seeding approach as PatchSim



Impacts on Reproductive number with release

- After release, spike in transmission driven by additional interactions at work, retail, and other
- At 25% release (70-80% remain compliant)
- Translates to 15% increase in transmission, which represents a 1/6th return to pre-pandemic levels