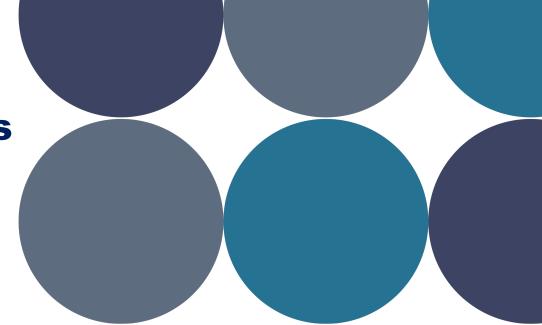




Adaptive Disease Modelling for COVID-19 decision-making in South Africa



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Modelling and Simulation Hub, Africa (MASHA)

**University of Cape Town** 



## South African COVID-19 Modelling Consortium (SACMC)

- Group of researchers from academic, nonprofit, and government institutions across South Africa
- Epi and Econ models developed by MASHA, SACEMA and HE2RO in conjunction with the NICD
- Extensive and ongoing input from clinicians, virologists, intensivists and epidemiologists to refine key model assumptions and parameters
- Purpose: support govt planning

#### **COVID-19 STATISTICS IN SA**



Learn more to Be READY for #COVID19 www.sacoronavirus.co.za

Covid-19 public hotline: 0800 029 999 WhatsApp 'Hi' to 0600 123 456

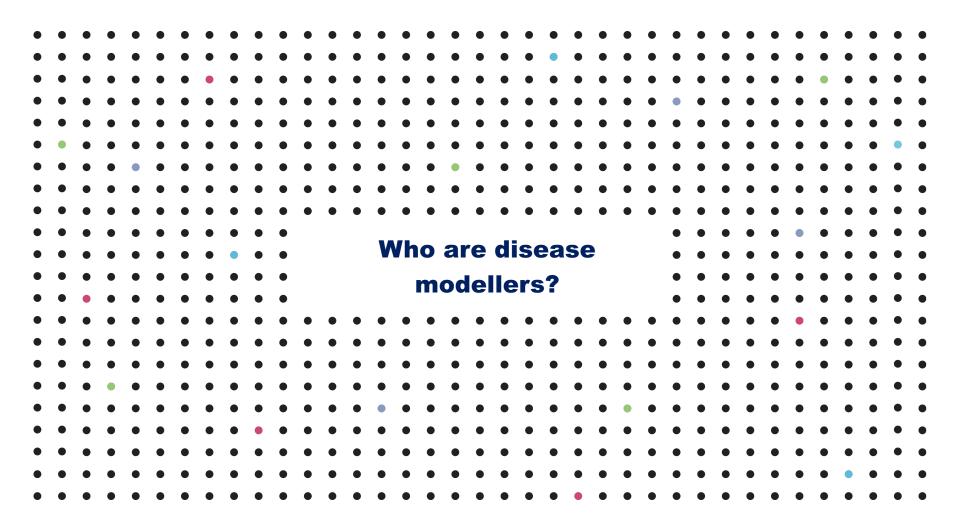


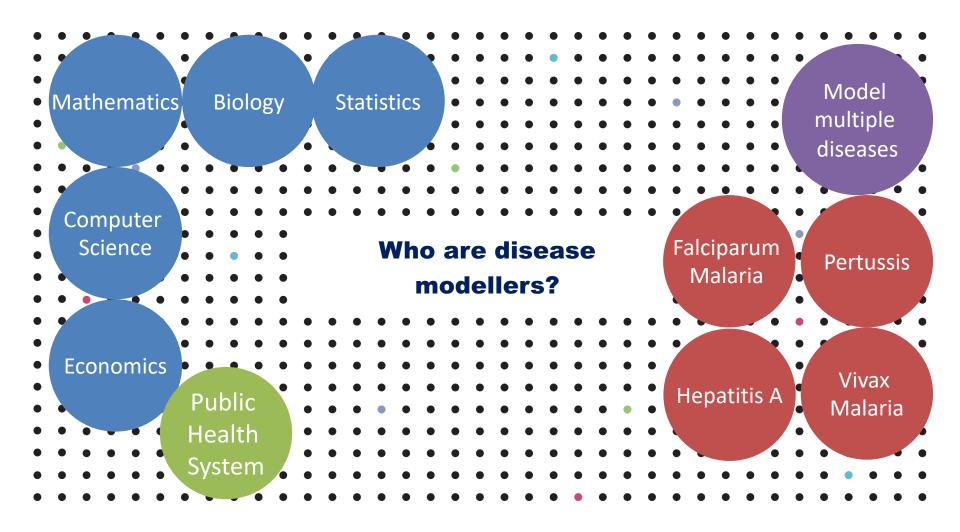
#### DOWNLOAD the COVID Alert SA app

The COVID Alert SA app can notify you if you have been exposed to another app user with coronavirus. Download it now to protect yourself and others









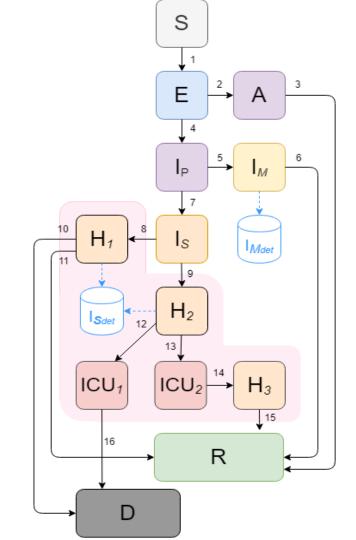
# Adaptive modelling for COVID-19 in South Africa



#### NCEM model structure (v1)

#### Modelling Method?

- Statistical prediction
- Compartmental models
- Metapopulation models
- Agent-based models



## **Modelling in Context**

- Absence of local level data
- Not yet possible to estimate impact of lockdown
- Models project estimated need for services
  - Utilisation of services is likely to be lower
  - Criteria for entry into hospital (general and ICU) likely to be stricter
- Behavioural changes in response to the epidemic unaccounted for
- Long run expectation of epidemic assumed to be the same across country (after accounting for population/age distribution)
- Budgets calculated before anything known on cost/resources needed
- UNCERTAINTY



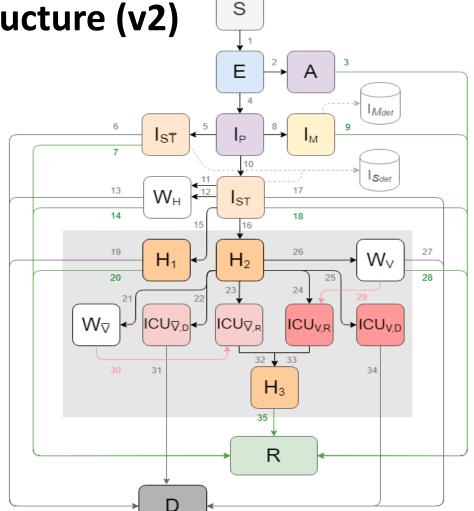
### NCEM model structure (v2)

#### **Model States**

S	Susceptible
Е	Exposed (not infectious)
А	Infected, asymptomatic (A)
Ι <sub>P</sub>	Infected, pre-symptomatic (Ip)
I <sub>M</sub>	Infected, mild
ls∓	Infected, severe, untreated
I <sub>ST</sub>	Infected, severe, seeking treatment
W <sub>H</sub>	Infected, severe, waiting for hospital bed
Hospitalised	Infected, severe, general ward (H <sub>1</sub> )
	Infected, severe, general ward pre-ICU (H <sub>2</sub> )
	Infected, critical, waiting for ICU, no ventilation (W $\bar{\sl}$ )
	Infected, critical, in ICU, not ventilated, non-survivor $(ICU_{\overline{V},D})$
	Infected, critical, in ICU, not ventilated, survivor (ICU <sub>V,R</sub> )
	Infected, critical, waiting for ICU & ventilation ( $W_V$ )
	Infected, critical, in ICU, ventilated, survivor (ICU $_{\rm V,R}$ )
	Infected, critical, in ICU, ventilated, non-survivor (ICU <sub>V,D</sub> )
	Infected, severe, general ward post-ICU (H <sub>3</sub> )
R	Removed (recovered)
D	Died
I <sub>Mdet</sub>	Detection of mild cases (laboratory confirmed)
I <sub>Sdet</sub>	Detection of severe cases (laboratory confirmed)

#### **Model Flows**

- Force of infection
- Latent period (until asymptomatic infectiousness)
- 3. Recovery: duration of asymptomatic infectiousness)
- 4. Latent period (until symptomatic infectiousness)
- 5. Development of severe symptoms, does not seek treatment
- 6. Death of severe, untreated case
- 7. Recovery: duration of severe case's infectiousness
- 8. Development of mild symptoms
- 9. Recovery: duration of mild case's infectiousness
- 10. Development of severe symptoms, seeks treatment
- 11. Severe case waiting for a hospital bed (if bed capacity reached)
- 40. O TO TO THE TOTAL TO
- 12. Critical case waiting for a hospital bed (if bed capacity reached)
- 13. Death while waiting for hospital bed (excess mortality)
- 14. Recovery while waiting for hospital bed
- 15. Severe case admitted to hospital
- 16. Critical case admitted to hospital (pre-ICU progression)
- 17. Death of severe case while seeking treatment
- 18. Recovery of severe case while seeking treatment
- 19. Death of severe case in general hospital bed
- 20. Recovery of severe case in general hospital bed
- 21. Critical case in hospital, waiting for ICU admission (no ventilation)
- 22. Progression to ICU admission (no ventilation), non-survivor
- 23. Progression to ICU admission (no ventilation), survivor
- 24. Progression to ICU admission (with ventilation), survivor
- 25. Progression to ICU admission (with ventilation), non-survivor
- 23. I regression to 100 damicolon (with ventilation), non sarvivor
- 26. Critical case in hospital, waiting for ICU admission (ventilation)
- 27. Death of critical case while awaiting ICU & ventilation
- 28. Recovery of critical case while awaiting ICU & ventilation
- 29. Waiting critical case needing ventilation admitted to ICU
- 30. Waiting critical case not needing ventilation admitted to ICU
- 31. Death of critical case from ICU (not ventilated)
- 32. Critical case discharged from ICU (non-ventilated) to general ward
- 33. Critical case discharged from ICU (ventilated) to general ward
- 34. Death of critical case from ICU (ventilated)
- 35. Recovery of critical case (discharged from hospital)



Source: NICD COVID-19 Surveillance Reports

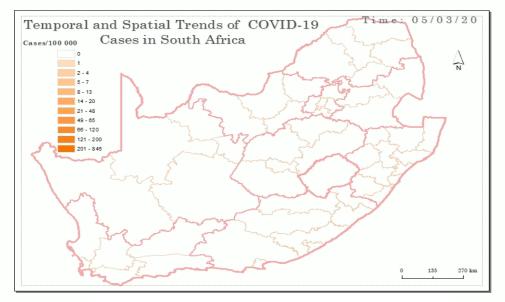
https://www.nicd.ac.za/diseases-a-z-index/covid-19/surveillance-reports/

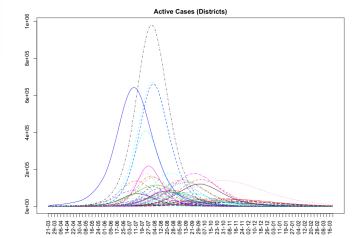
#### **NCEM Model Structure (v3)**

- Preparation of DATCOV and NICD surveillance data
- Analysis of mobility data from different sources
- Revision of Bed Availability



Source: SA COVID-19 Modelling Consortium (Preliminary output)

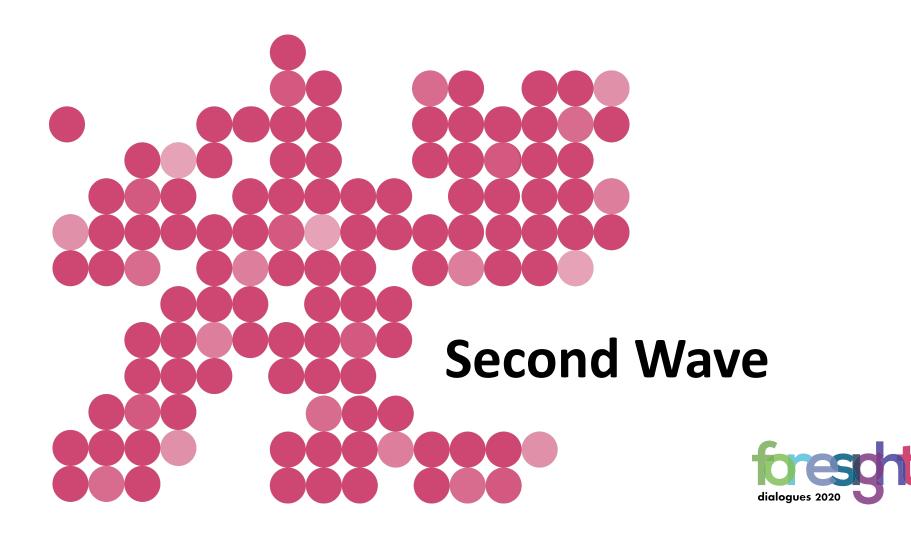




### **Stakeholders**

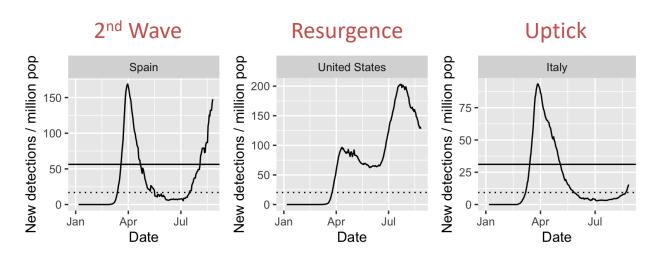
- NDOH Ministerial Advisory Committee
- Facility preparedness and expansion
- Staff required at facilities
- Ventilators
- Oxygen planning team
- Drug quantities
- Number of diagnostic test kits and testing algorithms
- Isolation facilities
- Number and placement of mortuary containers
- National and provincial treasuries: Covid-19 health budget





## • Will it really be a second wave?

- Second Wave
- Resurgence
- Uptick
- Upswing



https://covid.ourworldindata.org/data/owid-covid-data.csv [2020-08-30]



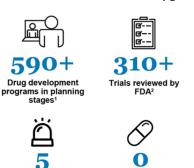
## Vaccine & Therapeutics

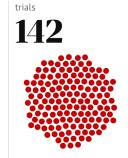
#### **Treatment & Cure**

- No cure yet.
- No fully licensed treatments. FDA granted emergency authorization to Remdesivir & Cytosorb
- Promising evidence:
  - Remdesivir (antiviral) for reducing hospital stay duration (possibly mortality)
  - Dexamethasone (steroid) for reducing mortality in ventilated & oxygenated patients

#### **Vaccination**

- R&D: over 140 teams with 231 candidates in development
- Timeline: aiming for a licensed vaccine by end-2020 to mid-2021
- Distribution: likely to be challenges with production constraints, hoarding and roll-out





Source: WHO. Last updated 30 Sep

vaccines not yet in human

**Pre-clinical** 

Phase 1
vaccines in small-scale
safety trials

Phase 2 Phase 3 vaccines in expanded vaccines in

safety trials

vaccines in large-scale efficacy trials

1

Approved

vaccines approved for general use



https://www.theguardian.com/world/ng-interactive/2020/oct/02/covid-vaccine-tracker-when-will-a-coronavirus-vaccine-be-ready

Treatments currently

approved by FDA for

use in COVID-19

COVID 19 treatments

currently authorized

for Emergency Use<sup>3</sup>

## **Imm**unity Dynamics

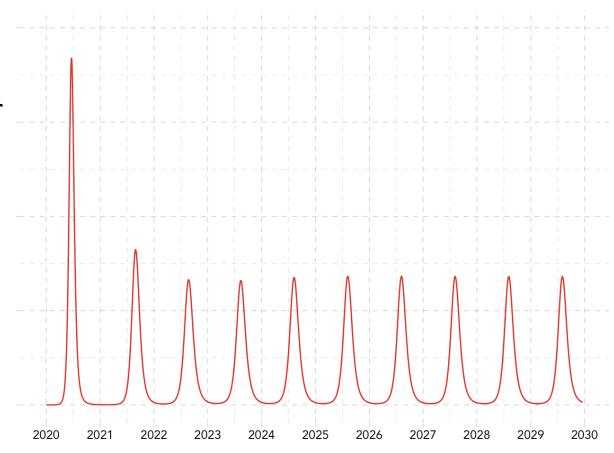
- A lot of uncertainty (and a lot of research underway):
- Antibodies, T-cells and memory cells detected during and post-infection
- Major question: What is the duration and strength of protective immunity?
- **Implications for vaccine development:** Uncertain what level of immune response is needed to protect against the virus, which is crucial for evaluating vaccine efficacy.
- **Life-long immunity:** Possible that neither a vaccine nor infection will produce life-long immunity (similar to influenza).



#### Scenario A: Short duration of immunity (40 weeks)

**Expect**: Annual epidemics with smaller peaks than the initial epidemic

COVID-19 becomes a regular occurence



#### Scenario B: Longer duration of immunity (2 years)

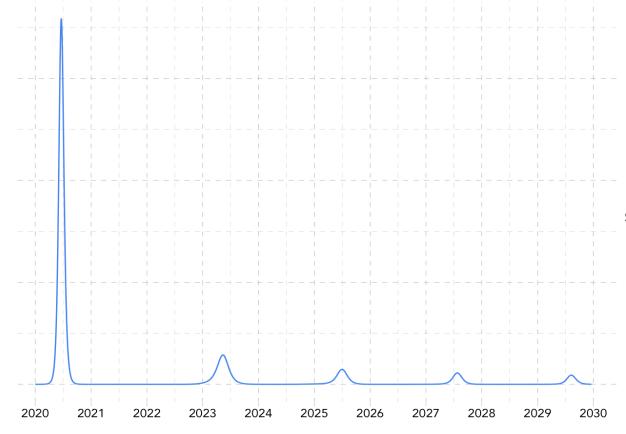
**Expect:** Two-year gap before the next epidemic (2023) followed by epidemics every 2 years with substantially smaller peaks than the initial epidemic.

Adapted from Kissler et al. 2020 Science

2030

# Scenario C:Longer duration immunity (2 years) with reduced disease upon reinfection

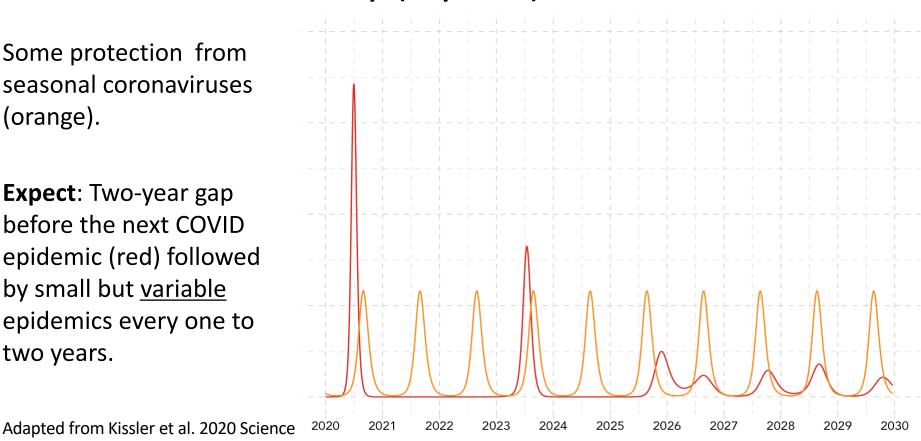
Expect: COVID-19
becomes a regular
occurrence BUT
severity of disease
(burden on
hospitals) decreases



## Scenario D: Weak cross immunity with long duration of immunity (2 years)

Some protection from seasonal coronaviruses (orange).

**Expect**: Two-year gap before the next COVID epidemic (red) followed by small but variable epidemics every one to two years.







### **Future Outlook**

- Invest in data for decision support
  - Cluster surveillance, Pandemic preparedness, Linked data systems
- Hospital Preparedness
  - Fungibility of wards, Healthcare Workers, Fully Equipped beds, Emergency services
- Procurement flexibility
- Improved National and Provincial coordination
- Vaccine Allocation and Implementation
- Planning for the 2<sup>nd</sup> Wave

