

COVID-19

Investor briefing

22 March 2022

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Agenda





ResApp's COVID-19 scientific advisory board



Prof. Elizabeth Talbot Geisel School of Medicine Dartmouth



Prof. Antonio Anzueto

University of Texas Health San Antonio



Prof. Catherine Bennett

Chair in Epidemiology Deakin University



Assoc. Prof. Joan Soriano

Hospital Universitario de la Princesa Madrid



Assoc. Prof. Mark Howard

Director of the Victorian Respiratory Support Service



Assoc. Prof. Paul Porter

Curtin University & Joondalup Health Campus

COVID-19

- Coronavirus disease (COVID-19) is an infectious disease caused by the SARS-CoV-2 virus
- Globally there has been >6m deaths, at a mortality rate estimated to be x10 higher than flu²
- At the peak of Omicron, Australians were waiting for up to 7 days for PCR test results (after waiting in queue for up to 6 hours) and rapid antigen tests were difficult to source³
- PCR and rapid antigen tests generally require a nasal swab as shown (right)
- Rapid antigen tests are widely used to screen and detect COVID-19



Obtaining the Nasopharyngeal Swab Specimen.



1. https://www.nejm.org/doi/full/10.1056/nejmvcm2010260

https://www.hopkinsmedicine.org/health/conditions-and-diseases/coronavirus/coronavirus-disease-2019-vs-the-flu

https://www.theage.com.au/politics/victoria/test-samples-no-longer-suitable-after-seven-day-wait-20220108-p59ms1.html
https://covid19.who.int (data to 18/3/2022)

COVID-19 latest trends

Market CNBC

Global conditions perfect for more Covid variants to emerge, WHO's Tedros says

Chloe Taylor Monday, 24 Jan 2022 12:54 PM EST



- Conditions are ripe for Covid-19 to mutate into more new variants, and it is dangerous to assume the pandemic is approaching its endgame, the WHO's top official warned Monday.
- Last week, an average 100 cases were reported every three seconds, Tedros Adhanom Ghebreyesus added, and someone lost their life to the virus every 12 seconds.
- However, Tedros was optimistic that with the right course of action, the pandemic could reach a turning point in 2022.





Do not assume COVID pandemic reaching 'end game', warns WHO

By Emma Farge and Mrinalika Roy

GENEVA, Jan 24 (Reuters) - The head of the World Health Organization (WHO) warned on Monday that it was dangerous to assume the Omicron variant would herald the end of COVID-19's acutest phase, exhorting nations to stay focused to beat the pandemic.

South Korea reports record Covid deaths as daily cases surge past 600,000

Despite record infections and fatalities, public opinion appears to support plans to ease Covid curbs in the coming days





How concerning is it that Covid infections are rising in the UK?

Nicola Davis Science Correspondent

♥@NicolaKSDavis

▶ Wed 16 Mar 2022

04.40 AFDT

Experts say rise was expected but further case increases and new variants are still a threat

The Observer China

With 37 million in lockdown and Covid plans under fire, Chinese ask: what comes next?

Elderly residents are wary of the jab even as Omicron spreads, and critics say zero-Covid policy is not sustainable

Helen Davidson in Taipei ©@heldavidson > Sun 20 Mar 2022 01.00 AEDT f y 😰



🛱 A resident is tested for Covid in Changzhou, China. Photograph: Sheldon Cooper/SOPA Images/REX/Shutterstock

Germany hits record Covid infection Philip rate since start of pandemic Berlin

Over past 24 hours 262,593 cases have been recorded, but with testing facilities at capacity, number could be higher.

Philip Oltermann in Berlin

✓ @philipoltermann
▶ Wed 16 Mar 2022 23.24
AEDT

Thu 17 Mar 2022 17.16 AEDT

Reuters

Future public health needs to manage COVID-19





Leaders in audio based diagnosis of respiratory health Why our 10 years of experience matters now more than ever



>3,500

cough sounds matched to gold standard PCR test results for COVID-19 infection from studies run in 2021/22



Breathe Easy

the **only** substantial pre-COVID-19 cough dataset in the world.

Enables us to be confident that we are detecting COVID-19, not just general respiratory disease.



3 Approved Products

CE Marked & TGA approved products in respiratory disease diagnosis and screening using audio



COVID-19 study design

R

741 Total Participants

US Arm 1: Decentralised, at-home, partnered with Phosphorus

US Arm 2: COVID-19 testing clinics, partnered with Covid Clinic

India Arm 1: COVID-19 testing clinics, partnered with Triomics

India Arm 2: Mild-moderate hospitalised patients, partnered with Triomics



446 COVID-19 Positive Participants



Comparator

Gold standard PCR test vs coughs recorded using a smartphone



Study results



Area under the Curve (AUC) 0.93*

*>0.9 is outstanding³

>90% Specificity vs Breathe Easy

Consistent across

study arm and location, age, gender and vaccination status, Delta & Omicron variants



Dinnes J., et al. (2021). Rapid, point-of-care antigen and molecular-based tests for diagnosis of SARS-CoV-2 infection. Cochrane Database of Systematic Reviews. https://doi.org/10.1002/14651858.CD013705.pub2 Hayer J., Kasapic D. and Zemmrich, C. (2021). Real-world clinical performance of commercial SARS-CoV-2 rapid antigen tests in suspected COVID-19: A systematic meta-analysis of available data as of November 20, 2020. International Journal of Infectious Diseases, 108, 592-602. https://doi.org/10.1016/j.ijid.2021.05.029

B Hosmer Jr D.W., Lemeshow S. and Sturdivant R.X. (2013). Applied Logistic Regression, Third Edition. John Wiley & Sons

COVID-19 mass-screening: *high sensitivity* algorithm Test with 92% sensitivity, 80% specificity @ 5% incidence of COVID-19



Post-test action

236 people receive a **positive** test result (**190** false positives)

Further testing with RAT or PCR required

764 people are ruled out by a negative test result (4 false negatives)0.5% chance of having COVID-19

No further testing required

COVID-19 diagnostics, a large and growing market

Potential to expand into patient management and other infectious disease therapy areas



Benefits of a smartphone-based test



Unlimited & instant

Smartphone only No supply chain issues Entire test process: ~30 seconds

Non-invasive

Cough sounds only, no uncomfortable swabs Likely better compliance





Resource sparing

Reduction in PCR & RAT testing Savings for governments insurers and individuals

Environmentally friendly

Reduces manufacturing, transport, chemicals & biosecurity waste



Additional options for public health management Noting these would be at the discretion of public health officials



Flexibility

- **Implementation**: App, SDK and AppClips/Android instant Apps.
- Flexible patient journey instructions post-test.

 Ability to verify test taker and their results via a photograph during the test (eg high risk settings, nursing home visitors etc).

Security

- Reporting
- Automated reporting of results to assist in public health management.
- Surveillance programs de-identified, aggregate level, patient data.



COVID-19 go-to-market: 3 key opportunities

1. Mass screening

Potential reduction in requirement for RAT/PCR tests – direct to consumer, government & insurance payers



2. Surveillance

Public health monitoring of outbreaks, prevalence rates etc

3. Diagnostic

High prevalence, low resource settings (eg LIMIC/emerging markets), high volume situations where RAT/PCR testing is not practical

Strategy: Global exclusive distribution license, instant scale and reach

Value proposition: Infinitely scalable, instant, non-invasive screening test to dramatically reduce COVID-19 rapid antigen and PCR testing, providing significant savings and reduced environmental impact



COVID-19 strategy





ResApp's overall strategy





Future implications of today's announcement

COVID-19 patient management



Enrolment and preliminary data collection completed. Review of data and algorithm development to follow.



Other infectious diseases

Potential to detect other infectious respiratory diseases – flu, RSV etc.

Global market of USD18Bn by 2026¹.



Pre-pandemic preparedness

Potential to develop candidate algorithms and further POC's to support management of future pandemics.

Next steps



Prospective trial:

Prove our mass screening data in a double-blinded, prospective clinical trial



Regulatory:

Engagement with key regulatory bodies on the data we have, Breathe Easy data context and future development path

Paediatric:

Initiate research in children <18 years



Patient monitoring:

Analyse data and develop algorithms



Commercial Engagement:

Already talking to a number of parties.





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Thank you!