The emergence of the SARS-CoV-2 variant of concern Omicron (Pango lineage B.1.1.529), first identified in Botswana and South Africa, may compromise vaccine effectiveness and lead to re-infections1. Here we investigated Omicron escape from neutralization by antibodies from South African individuals vaccinated with Pfizer BNT162b2. We used blood samples taken soon after vaccination from individuals who were vaccinated and previously infected with SARS-CoV-2 or vaccinated with no evidence of previous infection. We isolated and sequence-confirmed live Omicron virus from an infected person and observed that Omicron requires the angiotensin-converting enzyme 2 (ACE2) receptor to infect cells. We compared plasma neutralization of Omicron relative to an ancestral SARS-CoV-2 strain and found that neutralization of ancestral virus was much higher in infected and vaccinated individuals compared with the vaccinated-only participants. However, both groups showed a 22-fold reduction in vaccine-elicited neutralization by the Omicron variant. Participants who were vaccinated and had previously been infected exhibited residual neutralization of Omicron similar to the level of neutralization of the ancestral virus observed in the vaccination-only group. These data support the notion that reasonable protection against Omicron may be maintained using vaccination approaches.

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The emergence of the Omicron variant of SARS-CoV-2 in November 2021, first identified in South Africa and Botswana, was first described in South Africa1, followed shortly afterwards by confirmed transmission in Hong Kong2. Owing to the large number of mutations in the spike protein and elsewhere on the virus (https://covdb.stanford.edu/page/mutation-viewer/#omicron), there is concern that this variant will exhibit substantial escape from vaccine-elicited immunity3-5. Furthermore, several mutations in the spike receptor-binding domain and S2 fusion domain are predicted to increase transmission5.

Here we have used the human lung cell line H1299-ACE2 (Extended Data Fig. 1), which overexpresses the human ACE2 receptor6, to both isolate Omicron and test its neutralization by human plasma. We isolated Omicron virus using one passage on H1299-ACE2 cells and a second passage on H1299-ACE2 cells in co-culture with the Vero E6 African green monkey kidney cell line. Sequencing of the isolated virus confirmed it was the Omicron variant bearing the R346K mutation. We observed no mutations introduced in vitro as majority or minority variants (Extended Data Table 1). H1299-ACE2 cells were similar to Vero E6 cells in that they formed infection foci during infection with ancestral D614G and Beta variant viruses; however, the H1299-ACE2 cells formed more foci than unmodified Vero E6 cells (Extended Data Fig. 2a, b). Infection by cell-free Omicron of unmodified Vero E6 cells was inefficient (Extended Data Fig. 2c) and we could not use cell-free Omicron infection in Vero E6 cells to generate a useable virus stock of this isolate (Extended Data Fig. 2d).

We observed that Omicron infected the H1299-ACE2 cells in a concentration-dependent manner but did not infect the parental H1299
cells, indicating that human ACE2 is required for Omicron entry (Fig. 1a, b). We then tested the ability of plasma from individuals vaccinated with BNT162b2 to neutralize Omicron versus ancestral D614G virus in a live virus neutralization assay. We tested plasma samples taken from 19 individuals after they had received 2 doses of BNT162b2 (Extended Data Tables 2, 3), 6 of whom had no previous record of SARS-CoV-2 infection or detectable SARS-CoV-2 nucleocapsid antibodies indicative of previous infection (Methods). We also tested samples from a later time point for two of the vaccinated-only participants (Extended Data Table 3). The previously infected and vaccinated participants were infected with either ancestral SARS-CoV-2 strains or the Delta variant (Extended Data Table 3). To quantify neutralization in the live virus neutralization assay, we calculated the focus reduction neutralization test value (FRNT$_{50}$, the inverse of the plasma dilution required for a 50% reduction in infection focus number).

Fig. 1 | ACE2 dependence and neutralization of the Omicron variant by Pfizer BNT162b2-elicited immunity. a, Representative images showing infection foci in wells of a multi-well plate with titration of live SARS-CoV-2 Omicron virus on H1299-ACE2 and H1299 parental cells. Numbers above well images denote viral stock dilution. Scale bars, 2 mm. b, Number of foci as a function of Omicron virus stock dilution. Data are mean ± s.d. of six replicates from two independent experiments. c, Neutralization of Omicron virus compared with D614G ancestral virus by plasma from participants vaccinated with two doses of BNT162b2 and previously SARS-CoV-2 infected (blue) or uninfected (orange). Numbers in black above each virus strain are geometric mean titres (GMT) of the reciprocal plasma dilution (FRNT$_{50}$) resulting in 50% reduction in infection foci. The red horizontal line denotes the most concentrated plasma used. Twenty-one samples were tested from $n=19$ participants in 2 independent experiments ($n=13$ vaccinated and previously infected; $n=6$ vaccinated only). Grey points denote measurements where 50% neutralization was not achieved with the most concentrated plasma used. $P=4.8 \times 10^{-3}$, Wilcoxon rank-sum test. d, Geometric mean and 95% confidence interval of the fold change in neutralization between ancestral D614G and Omicron neutralization in plasma. Purple denotes all participants, blue denotes vaccinated individuals with previous SARS-CoV-2 infection, orange denotes vaccinated-only individuals, and yellow denotes all participants excluding those in whom 50% neutralization was not achieved. e, Mean predicted vaccine efficacy and 95% confidence intervals against symptomatic infection with Omicron using data from previous randomized controlled trials and the 22-fold difference between D614G and Omicron observed in this study$^{22,23}$. Predictions are for vaccinated and boosted (B, red) or vaccinated-only (V, blue) individuals.
of Omicron genomes. In addition, the timing of sample collection soon after vaccination (Supplementary Tables 2, 3) does not account for the waning of neutralization capacity\textsuperscript{21,26}.

So far, a milder course of Omicron infection has been observed in South Africa relative to previous infection waves in terms of reported numbers of patients in intensive care units and needing ventilation\textsuperscript{27}. Although there may be other unidentified contributing factors that lower pathogenicity\textsuperscript{28}, pre-existing immunity would be expected to be higher in the Omicron wave because of vaccination as well as immunity elicited by previous infection during one of three preceding infection waves in South Africa\textsuperscript{28}. Therefore, the incomplete Omicron escape from previous immunity described here may be an important factor accounting for the milder course of infection. Despite the extensive neutralization escape of Omicron, residual neutralization levels may still be sufficient to protect from severe disease\textsuperscript{28,29}. Other facets of the adaptive immune response elicited by vaccination and previous infection may increase protection. Furthermore, we observed that vaccination combined with previous infection elicits similar neutralization capacity against Omicron as vaccination without previous infection elicits against ancestral virus. This indicates that protection from symptomatic Omicron infection may occur when vaccination is combined with previous infection or boosting. This may explain why Pfizer BNT162b2 vaccination has been shown to substantially decrease the risk of hospital admission caused by Omicron infection in South Africa\textsuperscript{30} and supports the use of further vaccination and boosting to combat Omicron.

Online content
Any methods, additional references, Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at https://doi.org/10.1038/s41586-021-04387-1.
Methods

Whole-genome sequencing, genome assembly and phylogenetic analysis
cDNA synthesis was performed on the extracted RNA using random primers followed by gene-specific multiplex PCR using the ARTIC V3 protocol (https://www.protocols.io/view/covid-19-artic-v3-illumina-library-construction-an-bibtканм). In brief, extracted RNA was converted to cDNA using the Superscript IV First Strand synthesis system (Life Technologies) and random hexamer primers. SARS-CoV-2 whole-genome amplification was performed by multiplex PCR using primers designed using Primal Scheme (http://primal.zibaproject.org/) to generate 400-bp amplicons with an overlap of 70 bp that covers the 30-kb SARS-CoV-2 genome. PCR products were cleaned up using AmpureXP purification beads (Beckman Coulter) and quantified using the Qubit dsDNA High Sensitivity assay on the Qubit 4.0 instrument (Life Technologies). We then used the Illumina Nextera Flex DNA Library Prep kit according to the manufacturer’s protocol to prepare indexed paired-end libraries of genomic DNA. Sequencing libraries were normalized to 4 nM, pooled and denatured with 0.2 N sodium acetate. Then, a 12-pM sample library was spiked with 1% PhiX (a PhiX Control v.3 adaptor-ligated library was used as a control). We sequenced libraries on a 500-cycle v.2 MiSeq Reagent Kit on the Illumina MiSeq instrument (Illumina). We assembled paired-end fastq reads using Genome Detective L126 (https://www.genomedetective.com) and the Coronavirus Typing Tool. We polished the initial assembly obtained from Genome Detective by aligning mapped reads to the reference sequences and filtering out low-quality mutations using the bcftools 1.7 2 mpileup method. Mutations were confirmed visually with BAM files using Geneious software ( Biomatters). P2 stock was sequenced and confirmed Omicron with the following substitutions: E:T91I, M:D33G, M:Q19E, M:A46T, N:P13L, N:R203K, N:G204R, ORF1a:K856R, ORF1a:L2084I, ORF1a:A2710T, ORF1a:T3255I, ORF1a:P3395H, ORF1a:I3758W, ORF1b:P3419L, ORF1b:I5666V, ORF9b:p10S, S:A67V, S:T95I, S:Y145D, S:L212I, S:G393D, S:R346K, S:S371L, S:S373P, S:S375F, S:S441K, S:G446K, S:S477N, S:T478K, S:E484A, S:Q493R, S:G496S, S:Q498R, S:N501Y, S:Y503H, S:T547K, S:D614G, S:H655Y, S:N679K, S:P681H, S:N764K, S:D796Y, S:N856K, S:Q954H, S:N969K and S:Y981F. Deletions: N:E3, N:R32, N:R53, ORF1a:S2083, ORF1a:L367F, ORF1a:S367, ORF1a:C366, ORF9b:e27, ORF9b:n28, ORF9b:a29, S:H69, S:V70, S:G142, S:V143, S:Y144 and S:N21L. The sequence was deposited at GISAID under accession EPI_ISL_7358094.

SARS-CoV-2 nucleocapsid enzyme-linked immunosorbent assay (ELISA)
Nucleocapsid protein (2 μg ml⁻¹) (Biotech Africa; catalogue (cat.) no. BA25-P) was used to coat 96-well, high-binding plates and incubated overnight at 4 °C. The plates were incubated in a blocking buffer consisting of 5% skimmed milk powder, 0.05% Tween 20, 1 × PBS. Plasma samples were diluted to 1:100 dilution in a blocking buffer added to the plates. Horseradish peroxidase (HRP)-conjugated IgG secondary antibody was diluted to 1:3,000 in blocking buffer and added to the plates followed by tetramethylbenzidine (TMB) peroxidase substrate (Thermo Fisher Scientific). Upon stopping the reaction with 1 M H₂SO₄, absorbance was measured at a 450-nm wavelength.

Cells
Vero E6 cells (ATCC CRL-1586, obtained from Cellonex) were propagated in complete growth medium consisting of Dulbecco’s modified Eagle medium (DMEM) with 10% fetal bovine serum (Hyclone) containing 10 mM of HEPES, 1 mM sodium pyruvate, 2 mM L-glutamine and 0.1 mM nonessential amino acids (Sigma-Aldrich). Vero E6 cells were passaged every 3–4 days. H1299 cell lines were propagated in growth medium consisting of complete Roswell Park Memorial Institute (RPMI) 1640 medium with 10% fetal bovine serum containing 10 mM of HEPES, 1 mM sodium pyruvate, 2 mM L-glutamine and 0.1 mM non-essential amino acids. H1299 cells were passaged every second day. The H1299-E3 (H1299-ACE2, clone E3) cell line was derived from H1299 (CRL-S803) as described in previous work and Supplementary Fig. 1. In brief, vesicular stomatitis virus G glycoprotein (VSVG) pseudotyped lentivirus containing ACE2 was used to spinfect H1299 cells. ACE-2 transduced H1299 cells (containing an endogenously yellow fluorescent protein labelled histone H2AZ gene) were then incubated at single-cell density in 96-well plates (Eppendorf) in conditioned medium derived from confluent cells. After 3 weeks, wells were detached using a 0.25% trypsin-EDTA solution (Gibco) and plated in 2 replicate plates, where the first plate was used to determine infectivity and the second was stock.

The first plate was screened for the fraction of mCherry-positive cells per cell clone upon infection with a SARS-CoV-2 mCherry expressing spike pseudotyped lentiviral vector. Screening was performed using a Metamorph-controlled (Molecular Devices) Nikon TIE motorized microscope (Nikon) with a 20×, 0.75 NA phase objective, 561-nm laser, and 607-nm emission filter (Semrock). Images were captured using an 888 EMCCD camera (Andor). The clone with the highest fraction of mCherry expression was expanded from the stock plate and denoted H1299-E. Infectivity was confirmed with mCherry expressing lentivirus by flow cytometry using a BD Fortessa instrument and analysed using BD FACSDiva Software (BD Biosciences). This clone was used in the outgrowth and focus forming assay. Cell lines have not been authenticated. The cell lines have been tested for mycoplasma contamination and are mycoplasma negative.

Virus expansion
All work with live virus was performed in biosafety level 3 containment using protocols for SARS-CoV-2 approved by the Africa Health Research Institute Biosafety Committee. ACE2-expressing H1299-E3 cells were seeded at 4.5 × 10⁴ cells in a well on a 6-well plate and incubated for 18–20 h. After one DPBS wash, the sub-confluent cell monolayer was inoculated with 500 μl universal transport medium diluted 1:1 with growth medium filtered through a 0.45-μm filter. Cells were incubated for 1 h. Wells were then filled with 3 ml complete growth medium. After 4 days of infection (completion of passage 1 (P1)), cells were trypsinized, centrifuged at 300g for 3 min and resuspended in 4 ml growth medium. Then, 2 ml was added to Vero E6 cells that had been seeded at 2 × 10⁴ cells per ml, 5 ml total, 18–20 h earlier in a T25 flask (approximately 1:8 donor-to-target cell dilution ratio) for cell-to-cell infection. The co-culture of ACE2-expressing H1299-E3 and Vero E6 cells was incubated for 1 h and 7 ml of complete growth medium was added to the flask and incubated for 4 days. The viral supernatant (passage 2 (P2) stock) was used for experiments. Further optimization of the viral outgrowth protocol used for subsequent Omicron isolates showed that addition of 4 ml instead of 2 ml of infected H1299-E3 cells to Vero E6 cells that had been seeded at 2 × 10⁵ cells per ml, 5 ml total, 18–20 h earlier in a T25 flask gave P2 stocks with substantially higher titres that could detectably infect Vero E6 cells. The Omicron virus isolate is available from the authors contingent on verification that it will be received and used in a biosafety level 3 facility.

Live virus neutralization assay
H1299-E3 cells were plated in a 96-well plate (Corning) at 30,000 cells per well 1 day before infection. Plasma was separated from EDTA-anticoagulated blood by centrifugation at 300g for 10 min and stored at −80 °C. Aliquots of plasma samples were heat-inactivated at 56 °C for 30 min and clarified by centrifugation at 10,000g for 5 min. Virus stocks were used at approximately 50–100 focus-forming units per microwell and added to diluted plasma. Antibody–virus mixtures were incubated for 1 h at 37 °C, 5% CO₂. Cells were infected with 100 μl of the virus–antibody mixtures for 1 h, then 100 μl of a 1× RPMI1640 (Sigma-Aldrich, R6504), 1.5% carboxymethylcellulose (Sigma-Aldrich, C4888) overlay was added without removing the inoculum. Cells were
fixed 18 h after infection using 4% PFA (Sigma-Aldrich) for 20 min. Foci were stained with a rabbit anti-spike monoclonal antibody (BS-R2B12, GenScript A02058) at 0.5 μg ml⁻¹ in a permeabilization buffer containing 0.1% saponin (Sigma-Aldrich), 0.1% BSA (Sigma-Aldrich) and 0.05% Tween-20 (Sigma-Aldrich) in PBS. Plates were incubated with primary antibody overnight at 4 °C, then washed with wash buffer containing 0.05% Tween-20 in PBS. Secondary anti-rabbit HRP conjugated antibody (Abcam ab205718) was added at 1 μg ml⁻¹ and incubated for 2 h at room temperature with shaking. TrueBlue peroxidase substrate (SeraCare 5S10-0030) was then added at 30 μl per well and incubated for 20 min at room temperature. Plates were imaged in an ImmunoSpot Ultra V S6-02-6140 Analyzer ELISPOT instrument with BioSpot Professional built-in image analysis (C.T.L).

Statistics and fitting
All statistics and fitting were performed in MATLAB v.2019b. Neutralization data were fit to:

\[ T_x = \frac{1}{1 + (D/ID_{50})} \]

Here \( T_x \) is the number of foci normalized to the number of foci in the absence of plasma on the same plate at dilution \( D \) and \( ID_{50} \) is the plasma dilution giving 50% neutralization. FRNT_{50} = 1/ID_{50}. Values of FRNT_{50} < 1 are set to 1 (undiluted), the lowest measurable value. The most concentrated plasma dilution was 1:25 and therefore FRNT_{50} < 25 were extrapolated. We have marked these values in Fig. 1c and calculate the fold-change FRNT_{50} either for the raw values or for values where FRNT_{50} > 25 in Fig. 1d.

Estimating vaccine efficacy from neutralization titres
Previously, the fold reduction in neutralization was shown to correlate and predict vaccine efficacy against symptomatic infection with ancestral SARS-CoV-2, and more recently with variants of concern in data from randomized controlled trials. The model was used here to estimate the vaccine efficacy against Omicron based on the fold drop observed in this study applied to the randomized controlled trial data. In brief, vaccine efficacy (VE) was estimated based on the \( \log_{10} \) fold drop in neutralization titre to Omicron \( f \), and the \( \log_{10} \) mean neutralization titre as a fold of the mean convalescent titre reported for BNT162b2 in phase II trials \( \mu \) using the equation:

\[ VE(\mu, f) = \int_{-\infty}^{\infty} N(x, \mu - f, \sigma) \frac{1}{1 + e^{-50(x-x_{10})}} \, dx \]

Here, \( N \) is the probability density function of a normal distribution with mean \( \mu - f \) and standard deviation \( \sigma \), and \( k \) and \( x_{10} \) are the parameters of the logistic function relating neutralization to protection for the Pfizer BNT162b2 vaccine which were fitted from randomized controlled trial data: \( \sigma = 0.46, k = 3 \) and \( x_{10} = \log_{10}0.2 \) for symptomatic infection. Importantly, \( \mu = \log_{10}2.4 \) for trial participants vaccinated with two doses of BNT162b2, and \( \mu = \log_{10}12 \) for vaccinated and boosted trial participants.

Informed consent and ethical statement
Blood samples were obtained after written informed consent from hospitalized adults with PCR-confirmed SARS-CoV-2 infection and/or vaccinated individuals who were enrolled in a prospective cohort study approved by the Biomedical Research Ethics Committee at the University of KwaZulu–Natal (reference BREC/00001275/2020). Use of residual swab sample was approved by the University of the Witwatersrand Human Research Ethics Committee (HREC) (ref. M210752).

Reporting summary
Further information on research design is available in the Nature Research Reporting Summary linked to this paper.

Data availability
Sequence of outgrown virus has been deposited in GISAID with accession EPI_ISL_7338094. Raw images of the data are available upon reasonable request.

Code availability
The sequence analysis and visualization pipeline are available on Github (https://github.com/nexstain/ncov). Image analysis and curve fitting scripts in MATLAB v.2019b are available on Github (https://github.com/sigallab/NatureMarch2021).

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Author contributions
A.S., P.L.M. and T.d.O. and R.J.L. conceived the study. A.S., S.C., K.K., T.M.-G. and L.J. designed the study and experiments. A.v.G., P.L.M. and J.N.B. identified and provided the virus sample. S.-H.H. generated and provided plaque purified Beta variant virus. M.-Y.S.M., F.K., B.I.G., M.E., K.K. and Y.G. set up and managed the cohort and cohort data. S.C., L.J., K.K., TM.-G., H.T., J.E.S., C.S., D.G.A., G.L., D.A., M.S., Y.G., Z.J. and K.R. performed experiments and sequence analysis with input from A.S., T.d.O., R.J.L and J.M.B. D.S.K., D.C. and M.P.D. performed predictions of vaccine efficacy based on the data. A.S., S.C., P.L.M., T.d.O., L.J., K.K., W.H., S.S.A.K., D.S.K., M.P.D., J.N.B., R.J.L. and M.-Y.S.M. interpreted data. A.S., L.J., D.S.K., S.C., G.L., P.L.M. and M.P.D. performed the manuscript with input from all authors.

Competing interests
Salim S. Abdool Karim is a member of the COVID advisory panel for emerging markets at Pfizer. The authors declare no other competing interests.

Additional information
Supplementary information The online version contains supplementary material available at https://doi.org/10.1038/s41586-021-04387-1. Correspondence and requests for materials should be addressed to Alex Sigal.

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Extended Data Fig. 1 | Generation of H1299-ACE2 clonal cell line. (A) The H1299 human non-small cell lung carcinoma cell line with YFP labelled histone H2AZ was spinfected with the pHAGE2-EF1a-Int-ACE2 lentivector. Cells were single cell cloned by limiting dilution in a 384-well plate. Clones were expanded into duplicate 96-well plates, where one plate was used to select infectable clones based on mCherry signal from infection with SARS-CoV-2 mCherry expressing spike pseudotyped lentivirus. Clones were chosen based on infectability and expanded from the non-infected replicate 96-well plate. (B) Flow cytometry of SARS-CoV-2 mCherry expressing spike pseudotyped lentivirus infection in H1299-ACE2 cells versus H1299 parental cells.
Extended Data Fig. 2 | Comparison of SARS-CoV-2 infection in H1299-ACE2 and Vero E6 cells. Both H1299-ACE2 and Vero E6 cells were infected with the same viral stock in the same experiment with D614G virus (A) or Beta virus (B) and a focus forming assay was performed. (C) Focus forming assay with stock of Omicron virus isolate on H1299-ACE2 and Vero E6 cells. (D) Comparison of passage 2 (P2) and passage 3 (P3) stock, where P3 stock was generated by infection of 1 mL of cell-free P2 stock in 20 mL of Vero E6 cells seeded at 2x10^5 cells per mL and incubated over 4 days. Numbers above well images denote viral stock dilution. Scale bar is 2 mm.
Extended Data Fig. 3 | Neutralization of the Beta variant by Pfizer BNT162b2 elicited immunity. Neutralization of the Beta variant virus compared to D614G ancestral virus in H1299-ACE2 (A) or Vero E6 cells (B) in participants vaccinated with BNT162b2 and infected by SARS-CoV-2 (green) or vaccinated only (orange). Numbers in black above each virus strain are geometric mean titers (GMT) of the reciprocal plasma dilution (FRNT$_{50}$) resulting in 50% reduction in the number of infection foci. Numbers in red denote fold-change in GMT between virus strain on the left and the virus strain on the right of each panel. Red horizontal line denotes most concentrated plasma used. Samples were tested from the n = 19 participants described in Table S2 and S3, where n = 6 were vaccinated only and n = 13 were vaccinated and previously infected. p = 0.006 for both (A) and (B) as determined by the Wilcoxon rank sum test.
Extended Data Table 1 | Codon frequency table

| Amino Acid Change | Nucleotide Change | Codon(s) Change | K632E23_N67 |
|-------------------|-------------------|-----------------|-------------|
| A67V              | 21762C>T          | 21761 GCT - 0   | GCT - 0     |
|                   |                   | GCT>GT T       | GT - 133    |
| *H69_V70del       | 21768_21771delACATGT | 21769_21771ACATGT | ACATGT - 0   |
|                   |                   | del - 123      | del - 123   |
| T95I              | 21846C>T          | 21845 ACT - 0   | ACT - 0     |
|                   |                   | ACT>ATT        | ATT - 164   |
| *G142D            | 21987_21986delCTG  | 21987_21986CTG  | CTG - 0     |
|                   |                   | del - 432      | del - 432   |
| *V143_Y145del     | 21990_21996delTTTATT | 21990_21996TTTATT | TTATT - 0   |
|                   |                   | del - 432      | del - 432   |
| *L212I            | 22194_22196delATT  | 22194_22196ATT  | ATT - 0     |
|                   |                   | del - 146      | del - 146   |
| *R214_D215        | 22204_22206insGAGCCAGAA | 22204_22206GAGCCAGAA | WT - 37     |
|                   |                   | insGAGCCAGAA - 74 |            |
| G338D             | 22578G>A          | 22577 GTT - 0   | GTT - 0     |
|                   |                   | GTT>CAT        | CAT - 256   |
| R346K             | 22598G>A          | 22598 AAG - 1   | AAG - 1     |
|                   |                   | AAG>AAA        | AAA - 250   |
| S371L             | 22674C>T          | 22674 TCC - 0   | TCC - 0     |
|                   |                   | TCC>GTC        | GTC - 152   |
| S373P             | 22679C>G          | 22679 TCA - 3   | TCA - 3     |
|                   |                   | TCA>CCA        | CCA - 166   |
| S375F             | 22686C>T          | 22686 TCC - 0   | TCC - 0     |
|                   |                   | TCC>TTC        | TTC - 160   |
| K417T             | 22813G>T          | 22811 AAG - 3   | AAG - 3     |
|                   |                   | AAG>AAT        | AAT - 934   |
| N440K             | 22882T>G          | 22880 AAT - 3   | AAT - 3     |
|                   |                   | AAT>AAG        | AAG - 761   |
| G444S             | 22898G>A          | 22898 GTT - 30  | GTT - 30    |
|                   |                   | GTT>AGT        | AGT - 870   |
| T478K             | 22995C>A          | 22994 ACA - 0   | ACA - 0     |
|                   |                   | ACA>AAC        | AAC - 59    |
| E484A             | 23013A>C          | 23012 GAA - 0   | GAA - 0     |
|                   |                   | GAA>GCA        | GCA - 110   |
| Q493R             | 23040A>G          | 23039 CAA - 0   | CAA - 0     |
|                   |                   | CAA>CGA        | CGA - 128   |
| G498S             | 23048G>A          | 23048 GTT - 0   | GTT - 0     |
|                   |                   | GTT>AGT        | AGT - 150   |
| Q498R             | 23055A>G          | 23054 CAA - 1   | CAA - 1     |
|                   |                   | CAA>CGA        | CGA - 144   |
| N501Y             | 23063A>T          | 23063 AAT - 0   | AAT - 0     |
|                   |                   | AAT>TAT        | TAT - 209   |
| Y505H             | 23075G>C          | 23075 TAC - 1   | TAC - 1     |
|                   |                   | TAC>CAC        | CAC - 261   |
| T547K             | 23202C>A          | 23201 ACA - 0   | ACA - 0     |
|                   |                   | ACA>AAC        | AAC - 777   |
| D614G             | 23403A>G          | 23402 GAT - 1   | GAT - 1     |
|                   |                   | GAT>GGT        | GGT - 1803  |
| H655Y             | 23529C>T          | 23525 CAT - 3   | CAT - 3     |
|                   |                   | CAT>TAT        | TAT - 1639  |
| N679K             | 23597G>T          | 23597 AAT - 1   | AAT - 1     |
|                   |                   | AAT>AAG        | AAG - 682   |
| P681H             | 23604G>A          | 23603 CCT - 0   | CCT - 0     |
|                   |                   | CCT>CAT        | CAT - 536   |
| Q954H             | 24422A>T          | 24422 CAA - 1   | CAA - 1     |
|                   |                   | CAA>CAT        | CAT - 753   |
| N969K             | 24468G>T          | 24467 AAT - 0   | AAT - 0     |
|                   |                   | AAT>AAC        | AAA - 1682  |
| L981F             | 24503C>T          | 24503 CTT - 0   | CTT - 0     |
|                   |                   | CTT>TTT        | TTT - 1797  |

This table shows the amino acid change, the nucleotide position of the genome, codon change and the frequency of the codon on the assembled genome.

*Only deletions or insertion where the adjacent codon was preserved were counted. WT - Wild Type, i.e reads without the insertion.
## Extended Data Table 2 | Summary table of participants

|                                | All     | Vaccinated only | Infected and vaccinated |
|--------------------------------|---------|-----------------|------------------------|
| Number of Participants         | 19      | 6               | 13                     |
| Age (years)                    | 52 (39-67) | 54 (36-71)     | 51 (45-63)             |
| Days post-vaccination          | 26 (14-33) | 14.5 (8.5-37.5) | 28 (18-32)            |
| Days post-infection            |         |                 | 379 (127-468)         |
| Days post-infection to vaccination |       |                 | 353 (114-444)         |
| Date range of symptom onset    |         |                 | Jun 2020 – Jul 2021   |
| Male sex                       | 7       | 2               | 5                      |

All values are median (IQR) and inclusive of all samples used (early and late timepoints for 2 participants).
## Extended Data Table 3 | Participant information per sample

| Sample | Participant | Age | Sex | Days post 2\textsuperscript{nd} vaccination dose | Days diagnostic swab to sample | Date symptom onset or diagnostic test | Infecting virus* | FRNT\textsubscript{50} D614G | FRNT\textsubscript{50} Omicron |
|--------|-------------|-----|-----|-----------------------------------------------|--------------------------------|--------------------------------------|-----------------|-----------------|-----------------|
| 1      | 1           | 60-69 | F   | 10                                           | -                              | -                                    | -               | 196             | 10.8            |
| 2      | 2           | 70-79 | M   | 10                                           | -                              | -                                    | -               | 463             | 26.1            |
| 3      | 3           | 70-79 | M   | 45                                           | -                              | -                                    | -               | 205             | 14.6            |
| 4      | 4           | 30-39 | M   | 14                                           | -                              | -                                    | -               | 485             | 31.1            |
| 5      | 4           | 70-79 | F   | 10                                           | -                              | -                                    | -               | 199             | 15.4            |
| 6      | 4           | 70-79 | F   | 48                                           | -                              | -                                    | -               | 76.8            | 1.0             |
| 7      | 5           | 30-39 | F   | 10                                           | -                              | -                                    | -               | 1102            | 51.9            |
| 8      | 6           | 30-39 | F   | 33                                           | -                              | -                                    | -               | 151             | 4.6             |
| 9      | 7           | 40-49 | F   | 14                                           | 458                            | Jul-2020                             | Ancestral       | 10447           | 681             |
| 10     | 8           | 60-69 | F   | 63                                           | 468                            | Jul-2020                             | Ancestral       | 7468            | 414             |
| 11     | 9           | 20-29 | F   | 31                                           | 487                            | Aug-2020                             | Ancestral       | 2153            | 190             |
| 12     | 10          | 20-29 | M   | 37                                           | 493                            | Jul-2020                             | Ancestral       | 2697            | 121             |
| 13     | 11          | 60-69 | F   | 28                                           | 378                            | Jul-2020                             | Ancestral       | 54823           | 892             |
| 14     | 12          | 60-69 | M   | 26                                           | 379                            | Jul-2020                             | Ancestral       | 47023           | 1550            |
| 15     | 13          | 40-49 | F   | 32                                           | 479                            | Aug-2020                             | Ancestral       | 13517           | 955             |
| 16     | 14          | 50-59 | M   | 30                                           | 370                            | Sep-2020                             | Ancestral       | 11590           | 681             |
| 17     | 15          | 40-49 | F   | 22                                           | 456**                          | Jun-2020**                           | Ancestral/Delta | 564             | 5.0             |
| 18     | 16          | 40-49 | M   | 18                                           | 83                             | Jul-2021***                          | Delta           | 10511           | 749             |
| 19     | 17          | 70-79 | M   | 37                                           | 8                              | Jul-2021**                           | Delta           | 3074            | 138             |
| 20     | 18          | 50-59 | F   | 13                                           | 127                            | Jul-2021***                          | Delta           | 2205            | 385             |
| 21     | 19          | 60-69 | F   | 14                                           | 103                            | Jul-2021**                           | Delta           | 7160            | 174             |

* Determined by infection wave in South Africa. First infection wave (April-October 2020) consisted of ancestral strains with the D614G mutation. Third infection wave (April-October 2021) was dominated by the Delta variant. **Participant reinfected during Delta infection wave, sample is taken 3 months post-recovery of Delta infection. Asymptomatic during reinfection. ***Asymptomatic.
Reporting Summary

Nature Portfolio wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Portfolio policies, see our Editorial Policies and the Editorial Policy Checklist.

Statistics

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

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- The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
- A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
- The statistical test(s) used AND whether they are one- or two-sided
- Only common tests should be described solely by name; describe more complex techniques in the Methods section.
- A description of all covariates tested
- A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
- A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
- For null hypothesis testing, the test statistic (e.g. F, t, r) with confidence intervals, effect sizes, degrees of freedom and P value noted
- Give P values as exact values whenever suitable.
- For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
- For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
- Estimates of effect sizes (e.g. Cohen’s d, Pearson’s r), indicating how they were calculated

Our web collection on statistics for biologists contains articles on many of the points above.

Software and code

Policy information about availability of computer code

- Data collection: No code used
- Data analysis: All statistics and fitting were performed using MATLAB v.2019b.

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio guidelines for submitting code & software for further information.

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Sequence was deposited in GISAID, accession: EPI_ISL_7358094. All data are contained in the manuscript.
Field-specific reporting

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Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

| Sample size | Sample size was not pre-determined. We used all the samples we had available which met the inclusion/exclusion criteria. |
|-------------|-------------------------------------------------------------------------------------------------------------------|
| Data exclusions | We excluded samples from PfizerSNT162b2 vaccinated participants who were previously infected with the Beta variant since we wanted to compare to the Omicron to Beta virus neutralization. We excluded samples positive for SARS-CoV-2 nucleocapsid (i.e. previously infected) where we could not determine the infecting variant/strain by a time of infection. |
| Replication | Repeated in an independent experiment on a different day. Geometric mean of replicate samples was used. |
| Randomization | Groups were determined based on whether |
| Blinding | No blinding. |

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

| Materials & experimental systems | Methods |
|----------------------------------|---------|
| n/a | n/a |
| ☐ | ☑ Antibodies |
| ☐ | ☑ Eukaryotic cell lines |
| ☀ | ☑ Palaeontology and archaeology |
| ☑ | ☑ Animals and other organisms |
| ☑ | ☑ Human research participants |
| ☑ | ☑ Clinical data |
| ☑ | ☑ Dual use research of concern |
| ☑ | ☑ ChIP-seq |
| ☑ | ☑ Flow cytometry |
| ☑ | ☑ MRI-based neuroimaging |

Antibodies

- Foci were stained with a rabbit anti-spike monoclonal antibody (BS-R2812, GenScript A02058) at 0.5 μg/mL. Secondary goat anti-rabbit horseradish peroxidase (Abcam ab205718) antibody was added at 1 μg/mL.

Validation

- Information sheet for A02058 at [https://www.genscript.com/antibody/A02058-MonoRab_SARS_CoV_2_Spike_S1_Antibody_BS_R2812_mAb_Rabbit.html](https://www.genscript.com/antibody/A02058-MonoRab_SARS_CoV_2_Spike_S1_Antibody_BS_R2812_mAb_Rabbit.html).
- Information sheet for ab205718: [https://www.abcam.com/goat-rabbit-igg-hl-hrp-ab205718.html](https://www.abcam.com/goat-rabbit-igg-hl-hrp-ab205718.html)

Eukaryotic cell lines

Policy information about [cell lines](http://cell lines).

- Vero E6 cells (ATCC CRL-1586) obtained from Cellonex in South Africa. The H1299-E3 cell line was derived from H1299 (CRL-5803) as described in [2] and Figure S1. H1299 cells were a gift from M. Oren, Weizmann Institute of Science.

Authentication

- Cell lines have not been authenticated.

Mycoplasma contamination

- The cell lines have been tested for mycoplasma contamination and are mycoplasma negative.

Commonly misidentified lines

- None.
Human research participants

Policy information about studies involving human research participants

Population characteristics
Participant characteristics are summarized in Table S1 and listed per participant in Table S2.

Recruitment
Blood samples were obtained from hospitalized adults with PCR-confirmed SARS-CoV-2 infection and/or vaccinated individuals who were enrolled in a prospective cohort study approved by the Biomedical Research Ethics Committee at the University of KwaZulu-Natal.

Ethics oversight
Study approved by the Biomedical Research Ethics Committee at the University of KwaZulu–Natal (reference BREC/00001275/2020). Use of residual swab sample was approved by the University of the Witwatersrand Human Research Ethics Committee [HREC] (ref. M210752).

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Flow Cytometry

Plots

Confirm that:
- The axis labels state the marker and fluorochrome used (e.g., CD4-FITC).
- The axis scales are clearly visible. Include numbers along axes only for bottom left plot of group (a ‘group’ is an analysis of identical markers).
- All plots are contour plots with outliers or pseudocolor plots.
- A numerical value for number of cells or percentage (with statistics) is provided.

Methodology

Sample preparation
Plasma was separated from EDTA anticoagulated blood by centrifugation at 500 rcf for 10 min and stored at ~80°C. Aliquots of plasma samples were heat-inactivated at 56°C for 30 min and clarified by centrifugation at 10,000 rcf for 5 min.

Instrument
Plates were imaged in an ImmunoSpot Ultra-V S6-02-6140 Analyzer ELISPOT instrument with BioSpot Professional built-in image analysis (C.T.I).

Software
BioSpot Professional built-in image analysis (C.T.I).

Cell population abundance
H1299-E3 clone was previously generated and described. Abundance of infected cells with lentiviral infection was 30%/

Gating strategy
H1299-E3 clone was previously generated and described. Gating was based on FSC/SSC for live cells, then uninfected cells were used to determine mCherry positive gating.

- Tick this box to confirm that a figure exemplifying the gating strategy is provided in the Supplementary information.