

Omicron extensively but incompletely escapes Pfizer BNT162b2 neutralization

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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.

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 Africa², followed shortly afterwards by confirmed transmission in Hong Kong³. 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 immunity^{4,5}. 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

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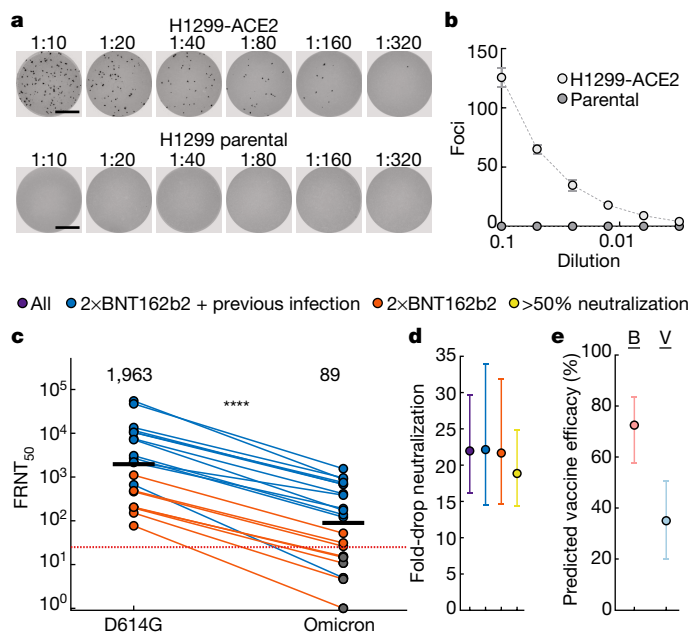


Fig. 1 | ACE2 dependence and neutralization of the Omicron variant by Pfizer BNT162b2-elicited immunity. **a**, Representative images showing infection foci in wells of a multi-well plate with titration of live SARS-CoV-2 Omicron virus on H1299-ACE2 and H1299 parental cells. Numbers above well images denote viral stock dilution. Scale bars, 2 mm. **b**, Number of foci as a function of Omicron virus stock dilution. Data are mean \pm 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 (FRNT₅₀) 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. $P = 4.8 \times 10^{-5}$, 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^{17,18}. Predictions are for vaccinated and boosted (B, red) or vaccinated-only (V, blue) individuals.

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 (FRNT₅₀, the inverse of the plasma dilution required for a 50% reduction in infection focus number).

Consistent with previous studies⁷⁻⁹, we observed that individuals who were vaccinated and had previously been infected exhibited higher neutralization capacity for ancestral virus relative to those who were vaccinated only (Fig. 1c). For all participants, the ability to neutralize Omicron was lower than for ancestral virus (Fig. 1c). The geometric mean titre (GMT) FRNT₅₀ for all participants was 1,963 for D614G and 89 for Omicron, a 22-fold difference (95% confidence interval 16–30) (Fig. 1d); the fold drop was the same for individuals who were vaccinated and had previously been infected (95% confidence interval 16–34) and in the vaccinated-only group (95% confidence interval 15–32) (Fig. 1d). Six of the samples showed fitted values for 50% Omicron neutralization that corresponded to a plasma concentration higher than the most concentrated plasma tested (a 1:25 dilution). This included the two samples collected at a later time point after vaccination, one of which showed a complete knockout of neutralization activity with Omicron (Fig. 1c, Extended Data Table 3). Excluding these 6 values from the analysis reduced the difference in GMT FRNT₅₀ between D614G and Omicron to 19-fold (95% confidence interval 14–25), well within the 95% confidence intervals of the fold differences for the raw values (Fig. 1d). Of note, Omicron virus neutralization by samples from individuals who were previously infected and vaccinated was similar to D614G neutralization by samples from participants vaccinated with two doses of BNT162b2 but not previously infected (Fig. 1c). GMT FRNT₅₀ for Omicron in the previously infected and vaccinated group was 305 (95% confidence interval 134–695), whereas GMT FRNT₅₀ for ancestral virus in the vaccinated-only group was 263 (95% confidence interval 147–472).

We compared these results with neutralization of the Beta variant^{6,10-16} using Beta and D614G virus infection of H1299-ACE2 (Extended Data Fig. 3a) and Vero E6 (Extended Data Fig. 3b) cells. The fold difference relative to the ancestral D614G virus was 4.3 for H1299-ACE2 cells and 5.0 for Vero E6 cells. Thus, results from these two cell lines indicated that Omicron exhibited approximately fourfold greater escape relative to Beta in our assays.

This study was not designed to reliably evaluate vaccine efficacy or protection from severe disease. However, a prediction of vaccine efficacy after a 22-fold drop in neutralization can be made in individuals vaccinated with BNT162b2 and individuals who were both vaccinated and boosted on the basis of data from randomized control trials using a model relating neutralization level to vaccine efficacy^{17,18}. Using this model and the differences in neutralization between Omicron and other SARS-CoV-2 strains (Methods), we predict a vaccine efficacy for preventing symptomatic infection by Omicron of 73% (95% confidence interval 58–83%) for vaccinated and boosted individuals, and 35% (95% confidence interval 20–50%) for vaccinated-only individuals; this suggests that Omicron compromises the ability of the vaccine to protect against infection in individuals in the vaccinated-only group but not in vaccinated and boosted individuals (Fig. 1e). We note that these predictions are similar to reports of actual vaccine efficacy in the UK¹⁹.

Shortly after we released these findings, several other groups have reported similar results^{3,20-23} including Pfizer–BioNTech (<https://www.businesswire.com/news/home/20211208005542/en/>). These results mirror ours, with large reductions in neutralization of Omicron compared with ancestral virus by vaccine-elicited immunity, neutralizing monoclonal antibodies and plasma from convalescent individuals previously infected with other variants. Notably, the Pfizer–BioNTech study reports that boosting seems to increase neutralization breadth, which reduces the escape by Omicron relative to ancestral virus; these results have been validated independently²¹. Unlike what was reported for boosting, we did not observe a lower fold drop in previously vaccinated and infected relative to the vaccinated-only participants in this study.

Limitations of this study include the presence of an R346K substitution in our virus stock. This putative escape mutation²⁴, which may confer moderate antibody resistance (https://jbloomlab.github.io/SARS2_RBD_Ab_escape_maps/escape-calc/), is not found in the majority

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^{25,26}.

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 ventilation²⁷. Although there may be other unidentified contributing factors that lower pathogenicity²⁸, 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 Africa²⁸. 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^{17,18}. 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 Africa²⁹ 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>.

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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 v.3 protocol (<https://www.protocols.io/view/covid-19-artic-v3-illumina-library-construction-an-bibtckann>). 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.zibraproject.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 1.126 (<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:T91, M:D3G, M:Q19E, M:A63T, N:P13L, N:R203K, N:G204R, ORF1a:K856R, ORF1a:L2084I, ORF1a:A2710T, ORF1a:T3255I, ORF1a:P3395H, ORF1a:I3758V, ORF1b:P314L, ORF1b:I1566V, ORF9b:P10S, S:A67V, S:T95I, S:Y145D, S:L212I, S:G339D, S:R346K, S:S371L, S:S373P, S:S375F, S:K417N, S:N444OK, S:G446S, S:S477N, S:T478K, S:E484A, S:Q493R, S:G496S, S:Q498R, S:N501Y, S:Y505H, S:T547K, S:D614G, S:H655Y, S:N679K, S:P681H, S:N764K, S:D796Y, S:N856K, S:Q954H, S:N969K and S:L981F. Deletions: N:E31, N:R32, N:S33, ORF1a:S2083, ORF1a:L3674, ORF1a:S3675, ORF1a:G3676, ORF9b:E27, ORF9b:N28, ORF9b:A29, S:H69, S:V70, S:G142, S:V143, S:Y144 and S:N211. The sequence was deposited at GISAID under accession EPI_ISL_7358094.

SARS-CoV-2 nucleocapsid enzyme-linked immunosorbent assay (ELISA)

Nucleocapsid protein ($2 \mu\text{g ml}^{-1}$) (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 a 1:100 dilution in a blocking buffer and 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 $1 \text{ M H}_2\text{SO}_4$, 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-5803) as described in previous work⁶ 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 gene³⁰) were then subcloned 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 line, 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-E3. 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 4.5×10^5 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 2×10^5 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 2×10^5 cells per ml, 20 ml total, 18–20 h earlier in a T75 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 500g 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% CO_2 . Cells were infected with 100 μl of the virus-antibody mixtures for 1 h, then 100 μl of a $1 \times \text{RPMI 1640}$ (Sigma-Aldrich, R6504), 1.5% carboxymethylcellulose (Sigma-Aldrich, C4888) overlay was added without removing the inoculum. Cells were

Article

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 $0.5 \mu\text{g ml}^{-1}$ 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 goat anti-rabbit HRP conjugated antibody (Abcam ab205718) was added at $1 \mu\text{g ml}^{-1}$ and incubated for 2 h at room temperature with shaking. TrueBlue peroxidase substrate (SeraCare 5510-0030) was then added at $50 \mu\text{l}$ 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:

$$Tx = \frac{1}{1 + (D/ID_{50})}$$

Here Tx 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-2¹⁸, and more recently with variants of concern¹⁷ 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 I/II trials (μ) using the equation:

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

Here, N is the probability density function of a normal distribution with mean $\mu - f$ and standard deviation σ , and k and x_{50} 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 $x_{50} = \log_{10} 0.2$ for symptomatic infection¹⁸. 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^{17,18}.

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_7358094. 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/nextstrain/ncov>). Image analysis and curve fitting scripts in MATLAB v.2019b are available on GitHub (<https://github.com/sigallab/NatureMarch2021>).

30. Sigal, A. et al. Variability and memory of protein levels in human cells. *Nature* **444**, 643–646 (2006).

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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.B., K.K. and Y.G. set up and managed the cohort and cohort data. S.C., L.J., K.K., T.M.-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. prepared 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.

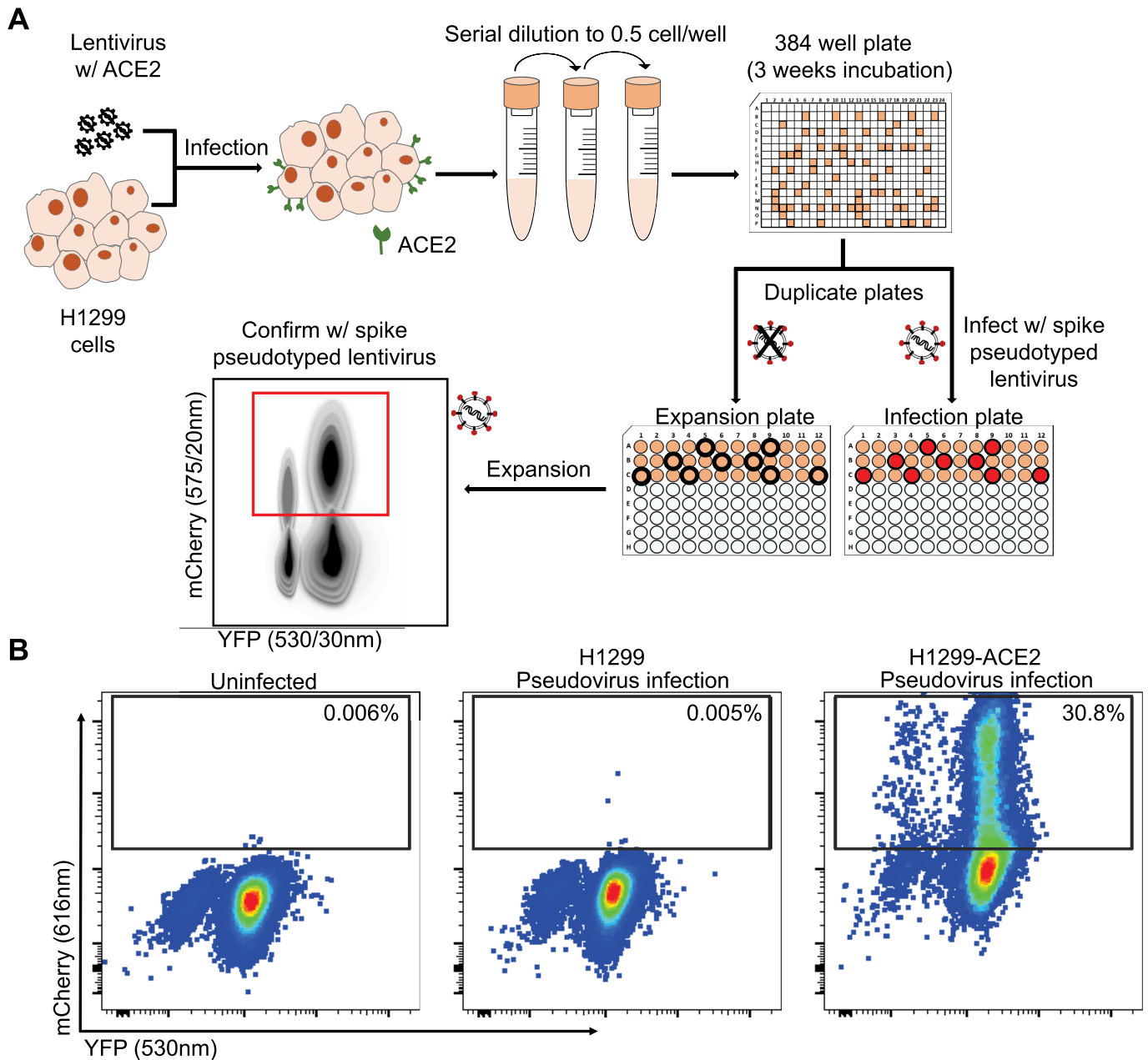
Additional information

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.

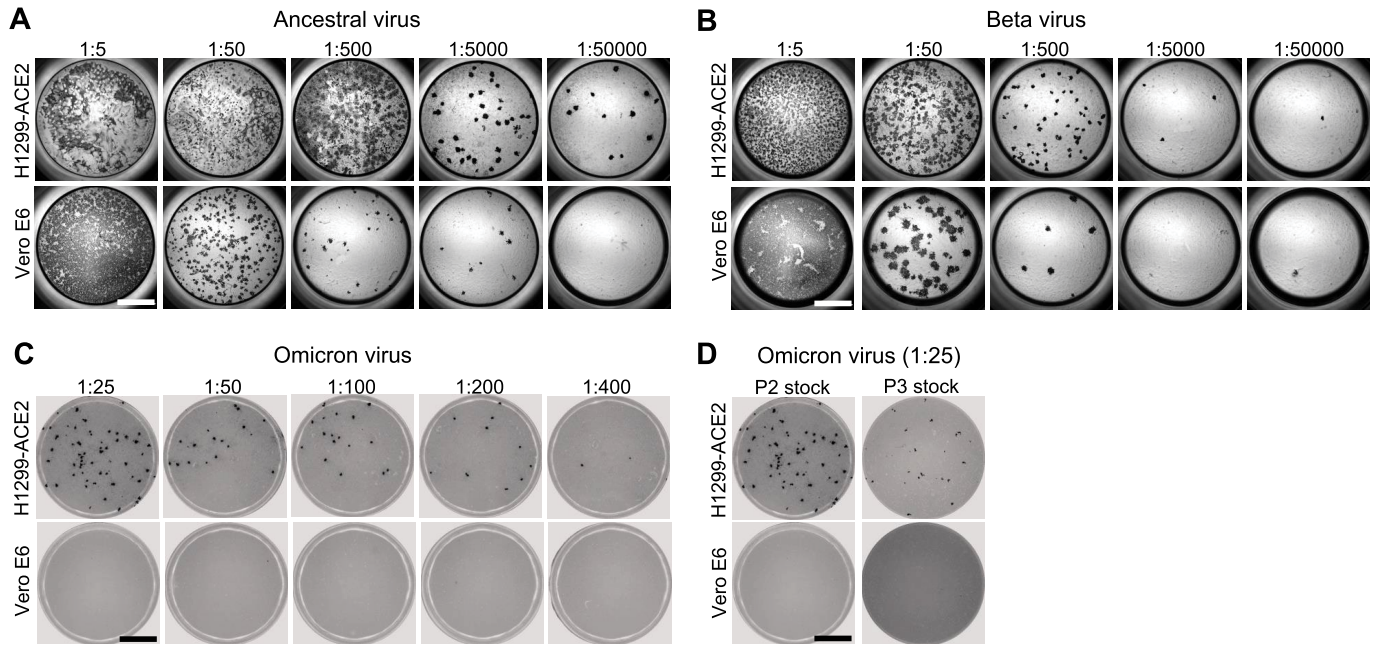
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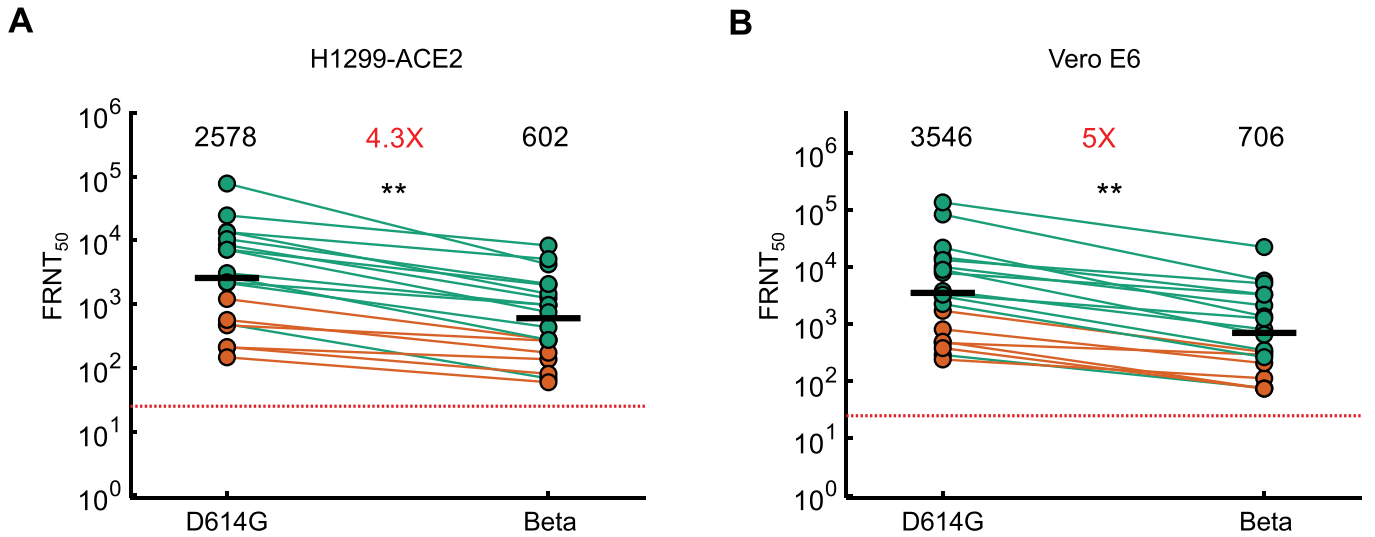
Extended Data Fig. 1 | Generation of H1299-ACE2 clonal cell line. (A) The H1299 human non-small cell lung carcinoma cell line with YFP labelled histone H2AZ was spinfected with the pHAGE2-EF1a-Int-ACE2 lentivector. Cells were single cell cloned by limiting dilution in a 384-well plate. Clones were expanded 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 2×10^5 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 (FRNT₅₀) 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.

Article

Extended Data Table 1 | Codon frequency table

Amino Acid Change	Nucleotide Change	Codon(s) Change	K032623_N67
A67V	21762C>T	21761 GCT>GTT	GCT - 0 GTT - 133
*H69_V70del	21766_21771delACATGT	21766_21771ACATGT >del	ACATGT - 0 del - 123
T95I	21846C>T	21845 ACT>ATT	ACT - 0 ATT - 164
G142D	21987_21989delGTG	21987_21989GTG >del	GTG - 0 del - 432
V143_Y145del	21990_21995delTTTATT	21990_21995TTTATT >del	TTTATT - 0 del - 432
L212I	22194_22196delATT	22194_22196ATT >del	ATT - 0 del - 146
R214_D215	22204_22205insGAGCCAGAA	22204_22205GAGCCAGAA >ins	WT - 37 insGAGCCAGAA - 74
G339D	22578G>A	22577 GGT>GAT	GGT - 0 GAT - 255
R346K	22599G>A	22598 AGA>AAA	AGA - 1 AAA - 250
S371L	22674C>T	22674 TCC>CTC	TCC - 0 CTC - 152
S373P	22679T>C	22679 TCA>CCA	TCA - 3 CCA - 166
S375F	22686C>T	22685 TCC>TTC	TCC - 0 TTC - 160
K417T	22813G>T	22811 AAG>AAT	AAG - 3 AAT - 934
N440K	22882T>G	22880 AAT>AAG	AAT - 3 AAG - 791
G446S	22898G>A	22898 GGT>AGT	GGT - 30 AGT - 870
T478K	22995C>A	22994 ACA>AAA	ACA - 0 AAA - 59
E484A	23013A>C	23012 GAA>GCA	GAA - 0 GCA - 110
Q493R	23040A>G	23039 CAA>CGA	CAA - 0 CGA - 128
G496S	23048G>A	23048 GGT>AGT	GGT - 0 AGT - 150
Q498R	23055A>G	23054 CAA>CGA	CAA - 1 CGA - 144
N501Y	23063A>T	23063 AAT>TAT	AAT - 0 TAT - 209
Y505H	23075T>C	23075 TAC>CAC	TAC - 1 CAC - 261
T547K	23202C>A	23201 ACA>AAA	ACA - 0 AAA - 777
D614G	23403A>G	23402 GAT>GGT	GAT - 1 GGT - 1803
H655Y	23525C>T	23525 CAT>TAT	CAT - 3 TAT - 1639
N679K	23599T>G	23597 AAT>AAG	AAT - 1 AAG - 682
P681H	23604C>A	23603 CCT>CAT	CCT - 0 CAT - 535
Q954H	24424A>T	24422 CAA>CAT	CAA - 1 CAT - 753
N969K	24469T>A	24467 AAT>AAA	AAT - 0 AAA - 1692
L981F	24503C>T	24503 CTT>TTT	CTT - 0 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).

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Extended Data Table 3 | Participant information per sample

Sample	Participant	Age	Sex	Days post 2 nd vaccination dose	Days diagnostic swab to sample	Date symptom onset or diagnostic test	Infecting virus*	FRNT ₅₀ D614G	FRNT ₅₀ Omicron
1	1	60-69	F	10	-	-	-	196	10.8
2	2	70-79	M	10	-	-	-	463	26.1
3	2	70-79	M	45	-	-	-	205	14.6
4	3	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	664	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 reinfecting during Delta infection wave, sample is taken 3 months post-recovery of Delta infection. Asymptomatic during reinfection. ***Asymptomatic.

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- 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

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Life sciences study design

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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 PfizerBNT162b2 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 (ie 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.

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n/a	Involved in the study
<input type="checkbox"/>	<input checked="" type="checkbox"/> Antibodies
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<input checked="" type="checkbox"/>	<input type="checkbox"/> Palaeontology and archaeology
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Methods

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<input checked="" type="checkbox"/>	<input type="checkbox"/> ChIP-seq
<input type="checkbox"/>	<input checked="" type="checkbox"/> Flow cytometry
<input checked="" type="checkbox"/>	<input type="checkbox"/> MRI-based neuroimaging

Antibodies

Antibodies used	Foci were stained with a rabbit anti-spike monoclonal antibody (BS-R2B12, 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_R2B12_mAb_Rabbit.html . Information sheet for ab205718: https://www.abcam.com/goat-rabbit-igg-hl-hrp-ab205718.html

Eukaryotic cell lines

Policy information about [cell lines](#)

Cell line source(s)	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 (See ICLAC register)	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:

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