

Immunogenic dynamics and SARS-CoV-2 variant neutralisation of the heterologous ChAdOx1-S/BNT162b2 vaccination: Secondary analysis of the randomised CombiVacS study

Javier García-Pérez,^{a,b,1} María González-Pérez,^{c,1} María Castillo de la Osa,^d Alberto M. Borobia,^e Luis Castaño,^f María Jesús Bertrán,^g Magdalena Campins,^h Antonio Portolés,^{i,j,s} David Lora,^{k,l,s,u} Mercedes Bermejo,^a Patricia Conde,^c Lourdes Hernández-Gutiérrez,^d Antonio Carcas,^e Eunete Arana-Arri,^f Marta Tortajada,^m Inmaculada Fuentes,ⁿ Ana Ascaso,ⁱ María Teresa García-Morales,^{k,s} Humberto Erick de la Torre-Tarazona,^a José-Ramón Arribas,^{b,o} Natale Imaz-Ayo,^f Eugènia Mellado-Pau,^g Antonia Agustí,^{p,q} Carla Pérez-Ingudua,ⁱ Agustín Gómez de la Cámara,^{k,l,r,s} Jordi Ochando,^c Cristóbal Belda-Iniesta,^t Jesús Frías,^e José Alcami,^{a,b,2*} and Mayte Pérez-Olmeda^{b,d,2**}, on behalf of the CombiVacS study Group³

^aUnidad de Inmunopatología del SIDA, Centro Nacional de Microbiología, Instituto de Salud Carlos III (ISCIII), Madrid, Spain

^bCentro de Investigación Biomédica en Red de Enfermedades Infecciosas (CIBERINFEC), Instituto de Salud Carlos III (ISCIII), Madrid, Spain

^cLaboratorio de Referencia en Inmunología, Centro Nacional de Microbiología, Instituto de Salud Carlos III (ISCIII), Madrid, Spain

^dLaboratorio de Serología, Centro Nacional de Microbiología, Instituto de Salud Carlos III (ISCIII), Madrid, Spain

^eServicio de Farmacología Clínica, Departamento de Farmacología y Terapéutica, Facultad de Medicina, Hospital Universitario La Paz, IdiPAZ, Universidad Autónoma de Madrid, Madrid, Spain

^fBiocruces Bizkaia, Hospital Universitario Cruces, CIBERDEM, CIBERER, Endo-ERN, UPV-EHU, Barakaldo, Spain

^gServicio de Medicina Preventiva y Epidemiología, Hospital Clínic de Barcelona, Barcelona, Spain

^hServicio de Medicina Preventiva y Epidemiología, Hospital Universitari Vall d'Hebron, Universitat Autònoma de Barcelona, Barcelona, Spain

ⁱServicio de Farmacología Clínica, Hospital Clínico San Carlos, IdISSC, Madrid, Spain

^jDepartamento de Farmacología y Toxicología, Facultad de Medicina, Universidad Complutense de Madrid (UCM), Madrid, Spain

^kInstituto de Investigación Sanitaria Hospital 12 de octubre (imas12), Facultad de Medicina, Universidad Complutense de Madrid (UCM)

^lCentro de Investigación Biomédica en Red de Epidemiología y Salud Pública (CIBERESP), Instituto de Salud Carlos III (ISCIII), Madrid, Spain

^mServicio de Prevención de Riesgos Laborales, Salud Laboral, Hospital Clínic de Barcelona, Barcelona, Spain

ⁿUnidad de Soporte a la Investigación Clínica, Vall d'Hebron Institut de Recerca, Barcelona, Spain

^oServicio de Medicina Interna, Departamento de Medicina, Facultad de Medicina, Hospital Universitario La Paz, IdiPAZ, Universidad Autónoma de Madrid, Madrid, Spain

^pServicio de Farmacología Clínica, Hospital Universitari Vall d'Hebron, Barcelona, Spain

^qDepartament de Farmacologia, Terapèutica i Toxicologia, Universitat Autònoma de Barcelona, Bellaterra, Barcelona, Spain

^rFacultad de Medicina, Universidad Complutense de Madrid (UCM), Madrid, Spain

^sSpanish Clinical Research Network – SCReN – ISCIII, Madrid, Spain

^tDirectorate Instituto de Salud Carlos III, Madrid, Spain

^uFacultad de Estudios Estadísticos, Universidad Complutense de Madrid (UCM), Madrid, Spain

Summary

Background The CombiVacS study was designed to assess immunogenicity and reactogenicity of the heterologous ChAdOx1-S/BNT162b2 combination, and 14-day results showed a strong immune response. The present secondary analysis addresses the evolution of humoral and cellular response up to day 180.

eClinicalMedicine
2022;50: 101529

Published online 1 July
2022

<https://doi.org/10.1016/j.eclim.2022.101529>

*Corresponding author at: Unidad de Inmunopatología del SIDA Instituto de Salud Carlos III, 28220 Majadahonda, Spain.

**Corresponding author at: Laboratorio de Serología, Instituto de salud Carlos III, 28220 Majadahonda, Spain.

E-mail addresses: ppalcami@isciii.es (J. Alcami), maytepererez@isciii.es (M. Pérez-Olmeda).

¹ Both authors contributed equally and should be considered as first authors.

² Both authors contributed equally and should be considered joint senior authors.

³ Study group members are listed in Appendix 1.

Methods Between April 24 and 30, 2021, 676 adults primed with ChAdOx1-S were enrolled in five hospitals in Spain, and randomised to receive BNT162b2 as second dose (interventional group [IG]) or no vaccine (control group [CG]). Individuals from CG received BNT162b2 as second dose and also on day 28, as planned based on favourable results on day 14. Humoral immunogenicity, measured by immunoassay for SARS-CoV-2 receptor binding domain (RBD), antibody functionality using pseudovirus neutralisation assays for the reference (G614), Alpha, Beta, Delta, and Omicron variants, as well as cellular immune response using interferon- γ and IL-2 immunoassays were assessed at day 28 after BNT162b2 in both groups, at day 90 (planned only in the interventional group) and at day 180 (laboratory data cut-off on Nov 19, 2021). This study was registered with EudraCT (2021-001978-37) and ClinicalTrials.gov (NCT04860739).

Findings In this secondary analysis, 664 individuals (441 from IG and 223 from CG) were included. At day 28 post vaccine, geometric mean titres (GMT) of RBD antibodies were 5616.91 BAU/mL (95% CI 5296.49–5956.71) in the IG and 7298.22 BAU/mL (6739.41–7903.37) in the CG ($p < 0.0001$). RBD antibodies titres decreased at day 180 (1142.0 BAU/mL [1048.69–1243.62] and 1836.4 BAU/mL [1621.62–2079.62] in the IG and CG, respectively; $p < 0.0001$). Neutralising antibodies also waned from day 28 to day 180 in both the IG (1429.01 [1220.37–1673.33] and 198.72 [161.54–244.47], respectively) and the CG (1503.28 [1210.71–1866.54] and 295.57 [209.84–416.33], respectively). The lowest variant-specific response was observed against Omicron and Beta variants, with low proportion of individuals exhibiting specific neutralising antibody titres (NT50) $>1:100$ at day 180 (19% and 22%, respectively).

Interpretation Titres of RBD antibodies decay over time, similar to homologous regimes. Our findings suggested that delaying administration of the second dose did not have a detrimental effect after vaccination and may have improved the response obtained. Lower neutralisation was observed against Omicron and Beta variants at day 180.

Funding Funded by Instituto de Salud Carlos III (ISCIII).

Copyright © 2022 The Authors. Published by Elsevier Ltd. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>)

Keywords: SARS-CoV-2; Heterologous vaccination; Neutralisation; Variants; Antibodies

Research in context

Evidence before this study

We did not perform a systematic search of the literature because of the rapidly developing situation. In summer 2021, the Delta variant of SARS-CoV-2 emerged and replaced the circulating Alpha strain. In November 2021 the Omicron variant was first described, and became predominant by March 2022 worldwide. In parallel, in summer 2021 different works pointed to a decrease in vaccine protection after six months of immunisation, in particular against new variants, due to waning of neutralising antibody activity. Together, these observations open the debate on a third dose booster that was adopted in many countries over the final months of 2021.

Added value of this study

The present results provide additional evidence on late immunogenicity – up to 6 months – of the heterologous ChAdOx1-S/BNT162b2 vaccination regime. Further, SARS-CoV-2 variant-specific elicited immunity is also reported here.

Implications of all the available evidence

Antibody titres decay over time, but delay in second dose administration has no deleterious effect on the immune response; on the contrary it resulted in better humoral responses. Functionally, all individuals exhibited neutralising titres $>1:100$ against G614 reference strain at day 28 after vaccine and a relevant proportion (76%) did so at day 180. However, variant-specific neutralisation was variable, with the lowest activity observed at day 180 against Omicron variant followed by Beta and Delta (19%, 22% and 56% with NT50 $>1:100$, respectively). These findings support the use of heterologous regimes, which is consistent with that arising from homologous schemes, and a third dose strategy in patients previously immunised with a combination of adenovirus- and mRNA-based vaccines.

Introduction

Early after the SARS-CoV-2 outbreak started – rapidly evolving to a worldwide pandemic – active immunization emerged as the key priority of global healthcare policies. Most scientific efforts have focused in vaccines development, which successfully resulted in three

homologous two-dose vaccines and one single-dose vaccine available for use in the European Union between late 2020 and early 2021. Notwithstanding, rare severe thrombotic with thrombocytopenia events related to ChAdOx1-S vaccine (Vaxzevria, AstraZeneca) and shortage in supplies had an impact on European vaccination plans and drove an interest in heterologous regimes. The combination of ChAdOx1-S and the mRNA vaccine BNT162b2 (Comirnaty, BioNTech) has been the first heterologous scheme studied. Results on 14-day¹ and 28-day² immunogenicity showed that robust humoral and cellular immune responses were elicited. Accordingly, and as planned in the protocol, participants included in the control group were offered to receive BNT162b2 as a second dose.

Concurrently, public health plans to control the pandemic faced a recurrent issue, namely the periodic outbreak of new SARS-CoV-2 variants. To date, numerous variants have been identified, of which Beta (B.1.351), Gamma (P.1), and Delta (B.1.617.2) are currently considered as variants of concern (VoC), the latter beginning in summer 2021 and soon becoming dominant.³ More recently a new variant – Omicron (B.1.1.529) – was first described in Botswana and South Africa. Because of its high transmissibility, Omicron is displacing Delta as the dominant variant in most world countries in less than 1 month, representing a new challenge in the control of pandemic.⁴ Not surprisingly, dynamics over time of variant-specific immune response induced by vaccination is a hot matter of research. Several works have reported a deterioration of vaccine-induced antibody response and a waning of protection against infection with either homologous regimes.^{5,6} By variants, a 3.5-fold to 14-fold reduction of serum neutralisation titres against Beta variant from vaccinated individuals has been reported.^{7–9} Further, ChAdOx1-S vaccine results evidenced undetectable neutralisation activity against Beta variant in 60% of vaccinated individuals and decay by a factor of up to 31.5% in the remaining 40%.¹⁰ Preliminary data from Omicron variant were even more worrying as a 14 to 30-fold reduction in neutralisation susceptibility elicited by immunisation was reported.^{11,12}

Notwithstanding this, the impact of waning neutralising antibodies on clinical efficacy is not clear. Some studies found that protection against hospitalization or death persisted at a robust level,⁶ while others showed that efficacy notably decreased, or even failed, against Beta^{10,13} and Delta⁶ variants. A consequent question is whether this reduction makes the variant resistant to vaccination. Some results on Beta variant comparing sera from naturally infected and vaccinated individuals have suggested that those vaccinated retain protective levels of humoral immunity,⁷ while others evidenced no efficacy in mild and moderate disease.¹⁰ It must be mentioned that effectiveness may be influenced by the interval between doses, with longer time associated to an enhanced antibody response¹⁴ and higher

effectiveness, as well as by cellular immunity, given the relevant role of CD4+ and CD8+ responses found in COVID-19 patients and cross-reactivity observed in unexposed individuals.^{15,16} Also, as suggested for Omicron variant, a decreased virulence could contribute to lower the rates of hospital admissions and death.^{17,18}

In addition, decay of variant-specific antibody titres is shifting focus towards the need of a third dose, especially in those with weakened immune systems such as older adults¹⁹ and immunosuppressed patients.^{20,21} Considering the available evidence, the European Medicines Agency (EMA) has issued recommendations for an extra dose with mRNA vaccines and is completing conclusions on booster doses for people with a normal immune system.²²

Increasing evidence on immunity dynamics aiming to answer these questions derives mostly from homologous vaccines, while heterologous regimes are still less studied. Here we present additional results of the CombiVacS study¹ addressing a) total and neutralising antibody dynamics of heterologous ChAdOx1-S/BNT162b2 vaccines combination, and b) immune response against different SARS-CoV-2 variants, including Delta and Omicron variants. This analysis also aims to provide valuable data to the debate on extra booster doses.

Methods

Study design and participants

Data from the 12-month, phase 2, open-label, randomised, controlled CombiVacS study are included in this secondary analysis. Full descriptions of the methods as well as safety and initial immunogenicity analyses have been previously published in detail.¹ Full study protocol is provided in Appendix 1 (p 26).

Healthy, or clinically stable, adults from 18 to 59 years old with no history of SARS-CoV-2 infection who had been vaccinated with a single dose of ChAdOx1-S between 8 and 12 weeks before screening were enrolled in the CombiVacS study to evaluate immunogenicity and reactogenicity of a second dose of the mRNA COVID-19 vaccine BNT162b2.

All participants provided written informed consent before enrolment. The trial complies with the principles of the Declaration of Helsinki and Good Clinical Practice. This study was approved by the Spanish Agency of Medicines and Healthcare Products (AEMPS) and by the Ethics Committee at University Hospital La Paz.

Randomisation and masking

Briefly, participants were randomly assigned (2:1) to receive one intramuscular injection of BNT162b2 (interventional group, IG) or maintain observation (control group, CG). Since the main immunogenicity objective was met, and reactogenicity was acceptable,¹

participants included in the control group were offered to receive BNT162b2 as a second dose at day 28, as planned in the protocol. A systematic randomisation stratified by study site, gender and age (18-49 years, and 50-59 years) was used. The randomization list was centrally generated with the SAS software for Windows (version 9.4; SAS Institute Inc., Cary, NC, USA), and imported into the secure Research Electronic Data Capture platform (REDCap version 8.7.4; Vanderbilt University, Nashville, TN, USA) used for the study electronic case report form (eCRF).

Procedures

Study procedures have been described in full previously.¹ In brief, at randomization clinical assessments were performed and blood samples for safety and immunology collected from all participants. Concurrently, participants in the interventional group were administered 0.3 mL BNT162b2 dose as a single intramuscular injection (day 0), whilst individuals from control group were vaccinated on day 28 of study. Planned follow-up visits for safety and immunologic purposes were scheduled on days 7, 14, 28, 90, 180 and 360. All vaccinated participants were on-site monitored for safety for at least 15 minutes. Safety procedures included both direct report from individuals during the post-vaccine observation period and online report using an electronic diary throughout the study follow-up period.

To the present analysis, the commercial electrochemiluminescence immunoassay (ECLIA) Elecsys® Anti-SARS-CoV-2 S assay (Roche Diagnostics GmbH, Mannheim, Germany) was used to detect antibodies (including IgG) specific to the SARS-CoV-2 spike protein receptor binding domain (RBD-S protein) on the Cobas e411 module (Roche Diagnostics GmbH, Mannheim, Germany),²³ with a measuring range from 0.4 to 250 U/mL (up to 2,500 U/mL with onboard 1:10 dilution and up to 12,500 with onboard 1:50 dilution). Values higher than 0.8 BAU/mL were considered positive.

Measurement of neutralising antibodies titres in a predefined subset of 198 participants was carried out by preincubation of diluted plasma samples with titrated pseudoviruses (10 ng p24Gag per well) generated by co-transfection of pNL4-3ΔenvRen and an expression vector for the different viral spikes (pcDNA3.1-S-CoV2Δ19-G614, -Alpha, -Beta, -Delta, or -Omicron) and added to Vero E6 cells in 96-well plates. Viral infectivity 48 hours post infection was assessed by measuring luciferase activity (Renilla Luciferase Assay, Promega, Madison, WI, USA) using a 96-well plate luminometer LB 960 Centro XS³ (Berthold Technologies, Oak Ridge, TN, USA). The titre of neutralising antibodies was calculated as 50% inhibitory dose (neutralising titre 50, NT₅₀), expressed as reciprocal of four-fold serial dilution of heat-inactivated sera (range 1:32–1:131,072) resulting in a 50% reduction of pseudovirus infection

compared to control without serum. Samples below the detection threshold (1:32 serum dilution) were given 1:16 value. Positive and negative controls were included in the assays and non-specific neutralisation was assessed using a related pseudovirus expressing the vesicular stomatitis virus envelope (VSV-G). Cellular immune response was measured in participants from two pre-selected sites by quantification of interferon- γ (IFN- γ) and interleukin-2 (IL-2) present in plasma on overnight stimulation of whole blood cultured with pools of SARS-CoV-2 spike peptides (2 μ g/mL) or dimethyl sulfoxide control. Cytokines were quantified using the next-generation enzyme-linked immunosorbent assay (ELISA) tool, Ella (ProteinSimple, San Jose, CA, USA). Full details on the pseudo-virus neutralisation assays and cellular immunity quantification are provided in the Appendix 1 (pp 18-19).

Outcomes

Outcomes included in the present secondary analysis were humoral immune response to vaccination as per antibodies titres and neutralising antibody titres at days 28, 90 and 180 after the BNT162b2 dose. Of note, in the control group outcomes at day 28 post-vaccine correspond to day 56 of study, and outcomes at day 180 correspond to day 152 post vaccine. In the control group, no outcomes were planned at day 90, according to the protocol. Alpha-, Beta-, Delta-, and Omicron-specific neutralising antibody titres at days 28 and 180 post-dose have been analysed in both study groups. Cellular response defined as inflammatory IFN- γ and IL-2 cytokines production against SARS-CoV-2 spike peptide pools at days 28 and 180 post-BNT162b2 dose were also assessed.

Statistical analysis

To the present analysis, the immunogenicity population included all the participants who were randomly assigned, completed all applicable visits, and for whom serological samples were available at the baseline visit and on days 28, 56 (only applicable to control group), 90 (only applicable to interventional group) and 180. Laboratory data cut-off was on November 19, 2021. Day-28 variables (i.e. humoral immunogenicity by RBD-specific IgG analysis, neutralising activity of SARS-CoV-2-specific antibodies, and cellular immunity) were analysed in 658, 194 and 114 individuals, respectively. Missing values from later visits were not imputed (Appendix 1 p 17).

Data were presented as geometric mean and 95% confidence interval (CI) or, for categorical variables, number, and percentage, unless otherwise stated. For serological measurements, difference at each time point was evaluated using ratio of geometric means. Since the outcome variable, i.e. antibodies against SARS-CoV-2 spike protein RBD on day 28 post-dose, was restricted

by technical limitations, a truncated regression model was used. The model incorporated right censoring values, raw data response with distribution lognormal data, and treatment effect (interventional group versus control group) adjusted by sex, age, and time between vaccine doses. Additionally, reverse cumulative distribution curve (RCDC) was plotted. A stratified analysis by sex, age and interval between vaccine doses was done for the humoral and cellular immunity endpoints. Laboratory parameters with values below detection limit were replaced by a value equal to the lowest limit divided by two. All analyses were carried out using the statistical software SAS, version 9.4. Sample size was calculated for primary efficacy endpoint – i.e. antibody titres at day 14 –, and was also considered appropriate to evaluate most of secondary endpoints (Appendix 1 p 14). Sample size calculation methods as well as independent data monitoring committee procedures have been described previously.¹

Composition of the independent data monitoring committee is provided in the Appendix 1 (p 23). This study was registered with EudraCT (2021-001978-37) and ClinicalTrials.gov (NCT04860739).

Role of the funding source

The funder – Institute of Health Carlos III (ISCIII), a public research organization – designed the trial in cooperation with the Spanish Clinical Trials Platform (SCReN), a public network of clinical trial units at the Spanish National Health System funded by the ISCIII. Trial coordination, participant recruitment and data analysis were performed by SCReN. All immunological procedures were performed at ISCIII. All authors reviewed and approved the original draft. All authors had full access to the full data in the study and accept responsibility to submit for publication.

Results

Of 676 participants enrolled and randomised between April 24th and 30th, 2021, 450 individuals were assigned to the interventional group, receiving BNT162b2 as second dose, and 226 were to the control group, maintaining observation. Participant's flow up to day 14 of study has been fully described previously.¹ 441 participants from the interventional group completed day 28 of study and 418 completed day 180. 223 individuals from control group received BNT162b2 vaccination at day 28; 212 and 199 of them completed day 56 of study (day 28 post vaccination) and day 180 (day 152 post vaccine), respectively (Figure 1).

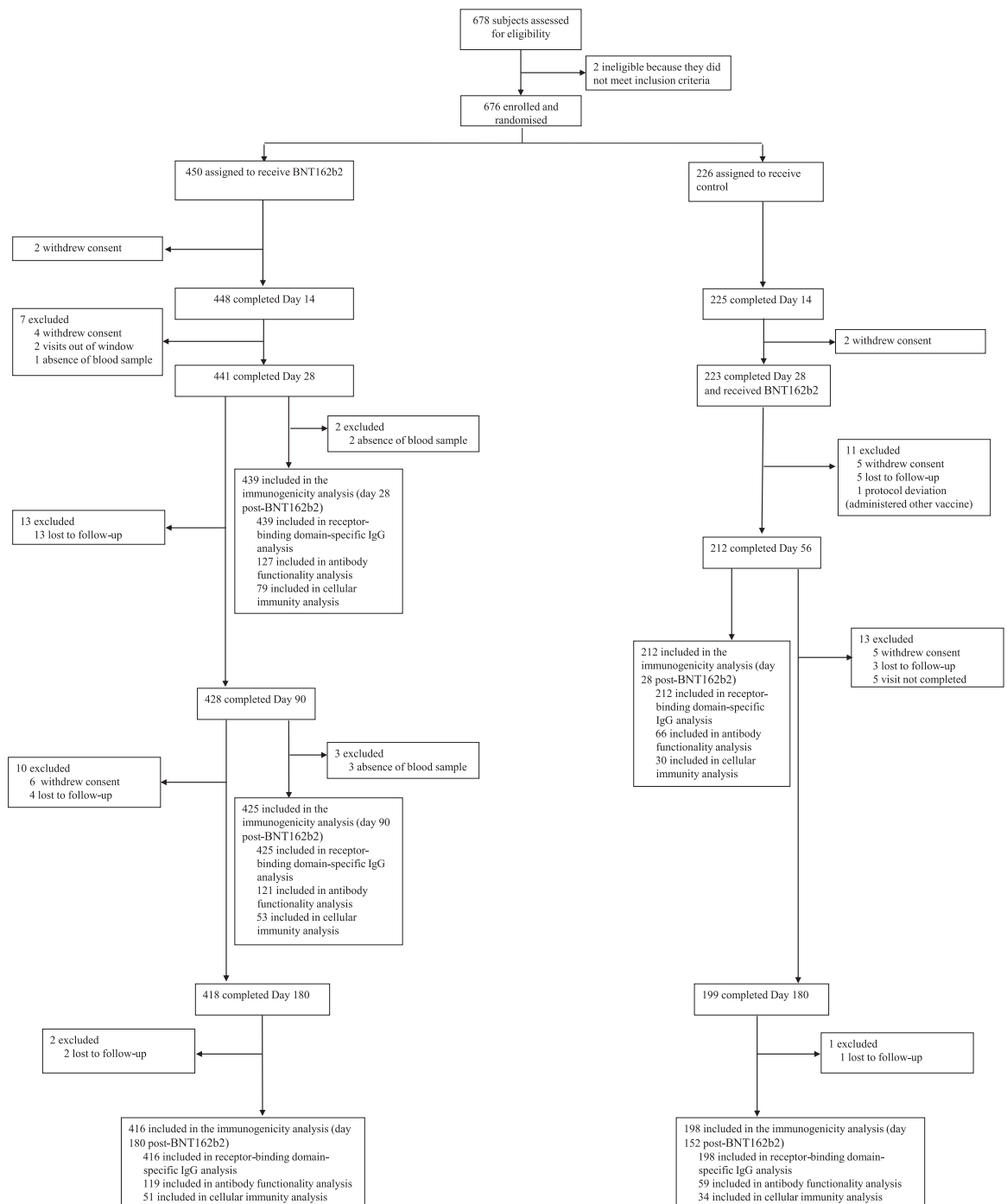
Baseline and demographic characteristics of the population are summarised in Table 1. 378 (57%) individuals were women and 289 (43%) were men. 431 (65%) participants were aged 18–49 years, and the mean age of both groups was 44.03 years (SD 8.82). In the control

group, mean (SD) interval between ChAdOx1-S and BNT162b2 administration was 89.03 days (5.92). Neither differences between groups were found in demographic characteristics at day 180 (Appendix 1 p 15).

Median time in collection of day-28 post- BNT162b2 dose blood sample was similar in both study groups (28 days [interventional] vs. 27 [control]), however variability was higher in individuals from the control group (range 21–38 days [interventional] vs. 16–42 [control]) (Appendix 1 p 2).

Results on immunogenicity dynamics in both groups show a decay in titres of antibodies specific to the SARS-CoV-2 S-protein RBD over time (Appendix 1 pp 3–4). In the interventional group, geometric mean titres (GMT) of S-RBD antibodies decreased from 7739.21 BAU/mL (95% CI 7371.53–8161.96) at day 14 after BNT162b2 second dose (fully reported earlier)¹ to 5616.91 BAU/mL (95% CI 5296.49–5956.71) at day 28, 2303.28 BAU/mL (95% CI 2141.66–2477.1) at day 90 and 1142.0 BAU/mL (1048.69–1243.62) at day 180. Of note, waning was slower from third to six month (mean lognormal difference -0.303 [95% CI -0.324–(-) 0.283]) than from first to third (-0.389 [95% CI -0.405–(-) 0.374]). The regression model for outcome variables to 180 days by treatment (interventional group versus control group) and adjusted by covariables is showed in the Appendix 1 p 16. Interestingly, immunogenic response in the control group was significantly stronger at day 28 after second dose (7298.22 BAU/mL [95%CI 6739.41–7903.37]) than in the interventional group ($p < 0.0001$). Likewise, antibody levels remained higher at day 180 in the control compared to interventional group (1836.4 BAU/mL [95%CI 1621.62–2079.62]; $p < 0.0001$) (Figure 2; Appendix 1 p 3). This effect was also observed in stratified analyses by sex and age (Appendix 1 pp 5–6). Adjusted differences in day-28 and day-180 lognormal RBD values in the interventional vs. control group resulted from the regression model were -0.0881 (95% CI -0.1239–(-)0.0523 [$p < 0.0001$]) and -0.1760 (95% CI -0.2277–(-)0.1242 [$p < 0.0001$]), respectively. In addition, the effect of delayed vaccination in the control group is linear over time, resulting in a difference of S-RBD antibody levels in the test vs. control group of -0.1329 (95% CI -0.1831–(-)0.0826) over the follow-up period.

Consistently with waning of antibody titres observed in the interventional group, a decay in neutralising antibodies was also evidenced. At day 14, GMT of neutralising antibodies was 1905.69 (95%CI 1625.65–2233.98) in the interventional group, which decreased to 1429.01 (95%CI 1220.37–1673.33) at day 28, to 480.68 (95%CI 398.27–580.13) at day 90 and to 198.72 (95%CI 161.54–244.47) at day 180. In the control group, neutralising antibody titres were similar to the interventional group both 28 days after second dose (1503.28 [95%CI 1210.71–1866.54]) and at day 180 (295.57 [95%CI 209.84–416.33]) (Figure 3; Appendix 1 p 7). RCDC for



neutralising antibodies in both study groups are shown in Appendix 1 (p 8). All patients from both interventional and control group exhibited high neutralising activity (NT₅₀ >1:100) against the reference variant G614 28 days post-BNT162b2 dose; a threshold that has been recently described²⁴ as associated with vaccine

efficacy. Yet decreased, a relevant proportion of individuals (76%) exhibited NT₅₀ >1:100 at day 180 of study. (Appendix 1 p 9).

Differences between variants and original G614 strain at different time points were analysed (Appendix 1, p 10). By SARS-CoV-2 variants, the poorest

	Interventional group (n = 450)	Control group (n = 226) ^d	Overall (n = 676)
Sex, n (%)			
Male	193 (43%)	101 (45%)	294 (43%)
Female	257 (57%)	125 (55%)	382 (57%)
Age (years), mean (SD)	43.93 (8.88)	44.10 (8.82)	43.98 (8.85)
Age group, n (%)			
18-49 years	293/450 (65%)	144/226 (64%)	437/676 (65%)
Male	123/293 (42%)	65/144 (45%)	188/437 (43%)
Female	170/293 (58%)	79/144 (55%)	249/437 (57%)
50-59 years	157/450 (35%)	82/226 (36%)	239/676 (35%)
Male	70/157 (45%)	36/82 (44%)	106/239 (44%)
Female	87/157 (55%)	46/82 (56%)	133/239 (56%)
Days between vaccines, mean (SD)	61.16 (5.73)	89.03 (5.92)	70.33 (14.32)

Table 1: Baseline characteristics of the population.

^d Based on 223 subjects (3 subjects withdrew consent before being immunized).

neutralisation capability at day 28 post-vaccination was found with Omicron variant in both interventional and control groups (GMT 144.84 [95%CI 116.65–179.85] and 204.84 [95%CI 151.99–276.06]). A decrease in NT50 was also observed for Beta variant in both interventional and control groups (GMT 293.21 [95%CI 234.8–366.15] and 483.89 [95%CI 352.53–664.2]). Of note, titre of Omicron and Beta-neutralising antibodies at day 28 in the control group was higher than in the interventional group ($p = 0.0641$ and $p = 0.0102$, respectively), whilst no differences between groups

were evidenced for Alpha- and Delta-neutralising antibody titres (Figure 4; Appendix 1 p 11). NT50 against Delta variant on day 28 after second dose was 717.13 (95%CI 587.27–876.13) in the interventional group and 837.14 (95%CI 609.7–1149.41) in the control group. Overall, day-28 NT50 was above 1:100 in 94% to 100% patients against all variants excepting Beta (88% patients) and Omicron (69%) (Appendix 1 p 12).

At day 180 Omicron-neutralising antibody titres decayed to 34.46 (95%CI 27.72–42.85) and 61.52 (95%CI 43.66–86.71) in the interventional and control

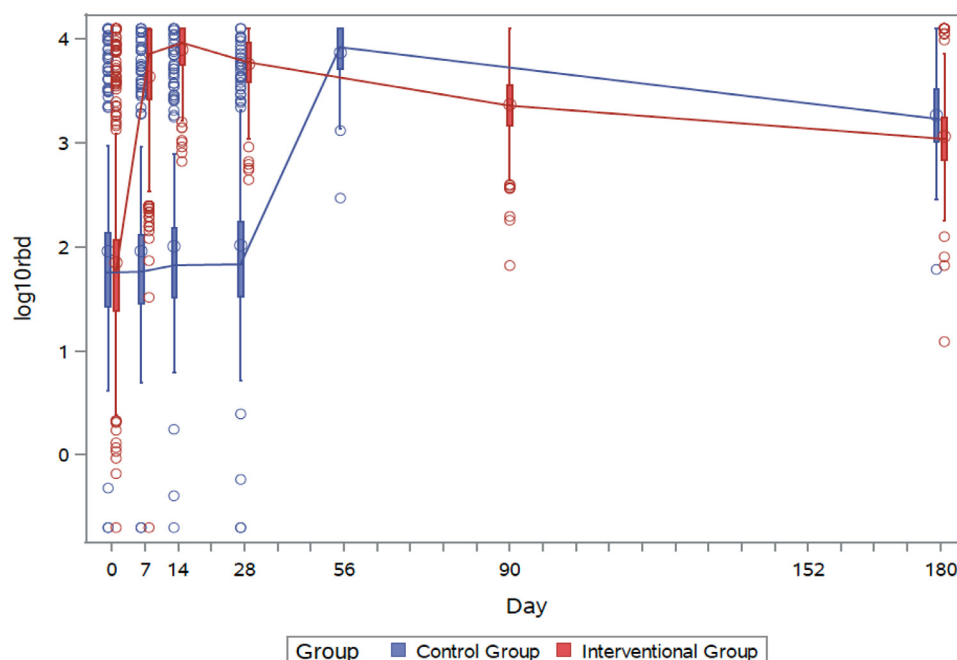


Figure 2. RBD (anti-spike) antibody titres measured in both interventional (red) and control (blue) groups over time. Interventional group was immunised at day 0 and control group at day 28. Accordingly day 180 corresponds to day 152 post vaccine in the control group.

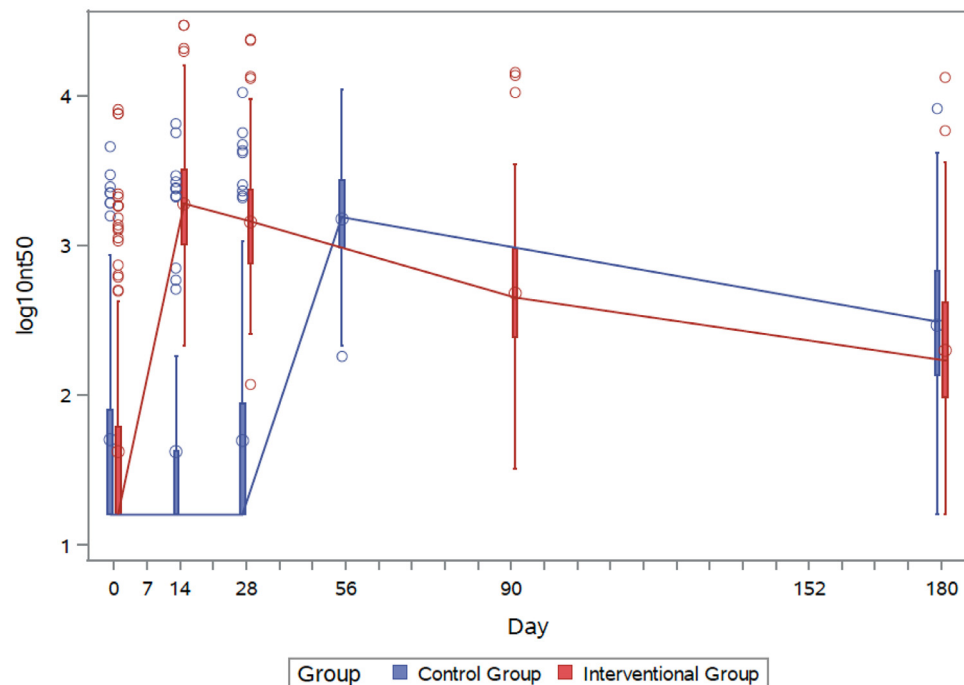


Figure 3. Neutralising antibodies titres (NT50) measured in both interventional (red) and control (blue) groups over time. Interventional group was immunised at day 0 and control group at day 28. Accordingly, day 180 corresponds to day 152 post vaccine in the control group.

groups, respectively ($p = 0.0038$). Beta-neutralising antibody titres decayed to 37.08 (95%CI 29.76–46.2) and 76.24 (95%CI 53.99–107.67) in the interventional and control groups, respectively ($p = 0.0004$). Delta NT50 decayed to 94.57 (95%CI 71.83–124.52) and 192.89 (95%CI 126.93–293.12) at day 180 in the interventional and control groups, respectively ($p = 0.0043$) (Figure 4; Appendix 1 p 11).

Regarding dynamics of functional spike-specific T-cell response, an increase in levels of both IFN- γ and IL-2 after vaccination is followed by a progressive waning over time. In the interventional group, maximum IFN- γ production was observed at day 14 post-dose. Levels decreased to 380.93 pg/mL (95% CI 309.07–469.5) at day 28 and 223.8 pg/mL (166.25–301.28) at day 180. In the control group, IFN- γ levels 28 days after BNT162b2 dose were 485.32 pg/mL (343.51–685.68), and decreased to 171.23 pg/mL (120.15–244.02) at day 180. Similarly, IL-2 concentrations were maximum at day 28 post-vaccination in both the interventional and control groups (244.07 pg/mL [95% CI 204.89–290.74] and 299.2 [217.81–411.01], respectively) and progressively decayed until day 180 (171.54 [133.08–221.12] and 170.25 [122.24–237.11], respectively). Of note, day-180 levels of both IFN- γ and IL-2 were higher than those present at baseline (Figure 5; Appendix 1 p 13).

Discussion

Our results provide evidence that humoral immune response of patients vaccinated with the heterologous ChAdOx1-S/BNT162b2 regime decays over time after peaking at day 14 post- BNT162b2 dose. A decline ranging 25%–27% in total RBD and neutralising antibody levels was observed after 28 days, which increased up to 70%–75% on day 90 and 86%–90% on day 180. This waning of immunogenicity was expected consistently with previous reports from COVID-19 mRNA vaccines^{5,25,26} that reported antibody half-life of 28–33 days.²⁶ A similar decline of about 80% in antibody levels was found with homologous BNT162b2 regime at 90 days after the second dose⁵ as well as with homologous mRNA-1273 regime, albeit the latter to a lesser extent (decline around 60%).²⁵ With homologous vaccination with ChAdOx1, antibodies are induced at lower levels than with homologous RNA regimens or heterologous vaccination of ChAdOx1 with BNT162b2²⁷ or mRNA-1273,²⁸ although it has been reported that antibodies decay with a slower kinetics.²⁹ Furthermore, these results are consistent with known kinetics of humoral immune response against acute viral infections, in which extrafollicular short-lived plasmablasts contribute to early antibody production – IgM, IgG, IgA –, while a secondary increasing contribution of germinal centre-derived plasma cells – with longer lifespan and larger secretory capacity – leads to secretion

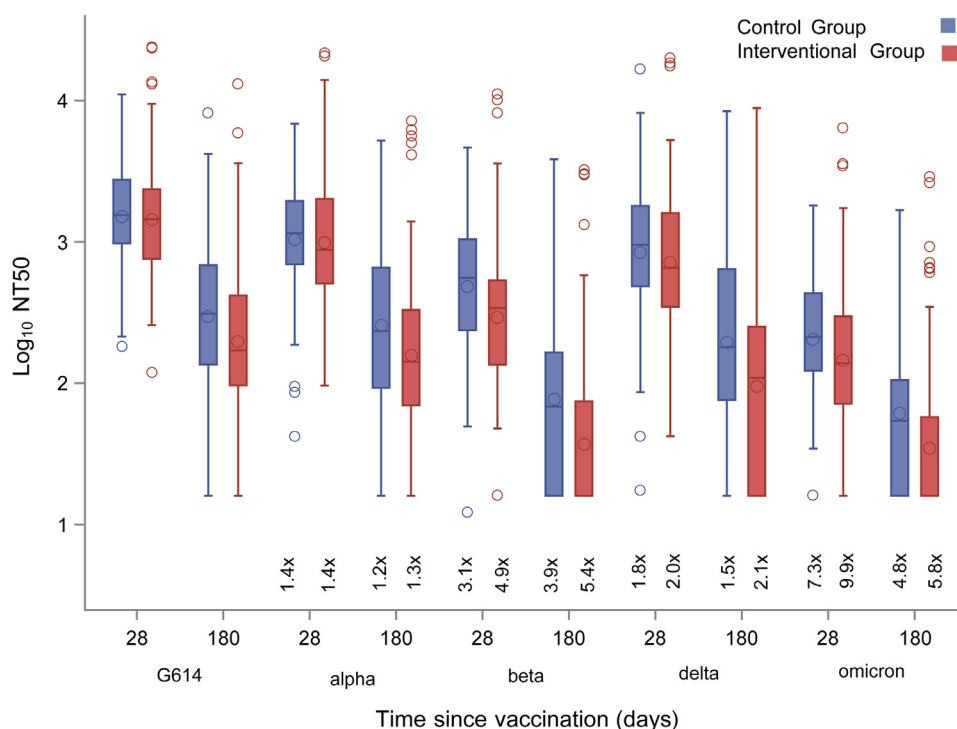


Figure 4. Neutralising antibodies titres (NT50) against SARS-CoV-2 variants measured in both interventional (red) and control (blue) groups at days 28 and 180 after BNT162b2 administration. Interventional group was immunised at day 0 and control group at day 28. Accordingly, day 180 corresponds to day 152 post vaccine in the control group.

Dashes and circles inside boxes indicated the median and arithmetic mean, respectively. Box limits indicate the interquartile range (IQR). Whiskers are adjusted to maximal and minimal values if lower than 1.5 times the IQR. Further outliers are indicated as circles.

of class-switched antibodies, mainly IgG. Considering that half-life of IgM is substantially shorter than IgG, a decay in antibody titres is common once the extrafollicular response is resolved. However they will rapidly rise if memory B cells are re-exposed to viral antigens in the future.³⁰ In this regard, we observed a slowing-down in SARS-CoV-2 antibody decay from month 3 to month 6, consistent with previous reports.²⁶

BNT162b2 administration to the control group 28 days later than the interventional one did not result in worse or weaker antibody responses 28 days after immunization. Actually, S-RBD antibodies and all variant-specific neutralising titres were higher – S-RBD and Beta-specific titres significantly higher – in the control group four weeks after vaccination suggesting a benefit of second dose delay. However, lack of antibody determination 14 days after immunization – when top levels of antibodies are reached – does not allow to perform a parallel kinetics of S-RBD and neutralizing antibodies between CG and IG to fully demonstrate that delayed administration of BNT162b2 results in better antibody responses.

Importantly, our results suggest that high levels of protection against Delta-variant persisted in both IG

and CG at day 28 after the second dose of the heterologous ChAdOx1-S/BNT162b2 scheme.

Regarding differences found at day 180 between control and intervention group it must be noted that this measurement carried over the 28-day delay in BNT162b2 administration to the control group (measured at day 152 – instead of 180 – after dose). A second explanation for these differences could be related with vaccination delay itself, supporting an apparent benefit for longer intervals between doses as found previously.^{14,31} Indeed, results from regression models pointed in this direction. Such delay could favour the maturation process of memory B cells from germinal centres (GC), over which B cells accumulate somatic mutations in their variable region leading to selection of those with higher affinity for a given viral antigen.³⁰ A recent study has demonstrated that antigen-driven activation of memory B cells persisted and matured up to 6 months after SARS-CoV-2 infection.³² The observation that at day 28 neutralisation activity in the control group against Beta was higher – and trended to higher against Omicron – while no differences were observed between groups for G614 reference strain, Alpha and Delta variants suggests that the 4-week delay in vaccination

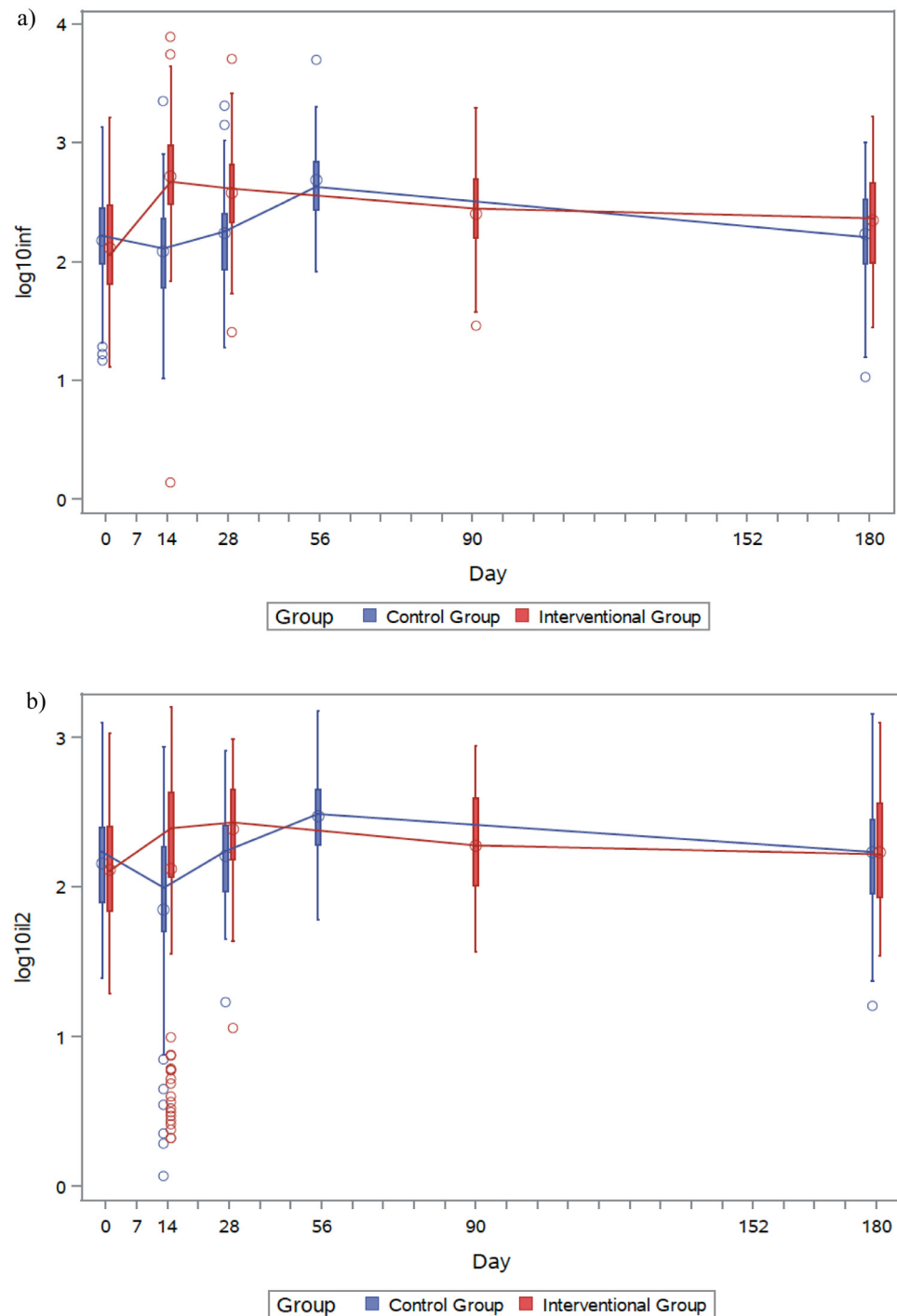


Figure 5. IFN- γ (a) and IL-2 (b) measured in both interventional (red) and control (blue) groups over time. Interventional group was immunised at day 0 and control group at day 28. Accordingly, day 180 corresponds to day 152 post vaccine in the control group.

window for the control group in our study might have contributed to a better affinity maturation against 'difficult' variants such as Beta and Omicron. Regarding cellular responses, similar decay of IFN- γ and IL-2 were

found in both groups at different time points which is consistent with the generation of memory T lymphocytes in which maturation process and selection of Tc receptor affinity is not dependent on somatic mutation.

Notwithstanding this, the sharp decay in RBD antibodies and neutralisation titres observed at day 180 as compared to day 14 and 28 support the use of a third immunization to reach higher protection levels, particularly considering the high infectivity potential of the Omicron variant.³³ Actually, despite the persistence of immune memory, antibody decay increases the risk of SARS-CoV-2 infection and a third dose becomes necessary to achieve protection against asymptomatic and symptomatic infections, particularly in aged groups above 60 and patients with risk factors for developing severe COVID-19 as immune suppression.^{19,20} It has been described that boosting with a third dose of mRNA vaccines generate potent neutralization of Omicron, shortening the difference in neutralization titres with other variants.³⁴ Thus, increased protection of a booster dose would be related not only to higher levels of neutralizing antibodies but to antibody maturation leading to the generation of antibodies with increased affinity to their targets. These qualitative changes are particularly important against escape variants as Omicron and represent an added value for a third dose. Unfortunately, very recently it has been described a waning effectiveness of a third dose of BNT162b2 against hospital admission after 3 months due to the Omicron variant.³⁵

As previously described,¹ the main limitation of the CombiVacS study is the absence of a control group completing the homologous ChAdOx1-S scheme to compare with the heterologous ChAdOx1-S/BNT162b2 regimen. This arm would have been very useful to compare antibody waning and neutralization activity in individuals vaccinated with homologous or heterologous vaccine regimens. Besides, the abovementioned 4-week delay between both study groups in BNT162b2 administration led to capture 5-month, rather than 6-month, post dose data in the control group. Although this limitation may have influenced some differences observed at day 180, overall results are consistent between groups. As mentioned, lack of antibody determination at day 14 in the CG limits the interpretation of the results. Also, we have found a low proportion of high-responder outliers, in particular before immunization with the second vaccine dose. We cannot rule out asymptomatic SARS-CoV-2 infection between both vaccine doses leading to a “booster-like” response after first immunization.

In conclusion, follow-up of individuals included in the CombiVacS trial that were immunised with heterologous ChAdOx1-S/BNT162b2 confirm waning of humoral and cellular responses over time, nevertheless a relevant proportion of individuals exhibited neutralising activity > 1:100 six months after full vaccination excepting against Beta and Omicron variants. These results support the use of a third dose six months after regular vaccination to enhance immune response, particularly against new VoCs, as Omicron. Further studies addressing immunogenicity using different heterologous vaccination schemes are warranted.

Contributors

Trial conceptualisation was done by C.B.-I., J.A., M.P.-O., A.M.B., A.J.C., J.F., J.R.A. and M.C. A.J.C., J.F., and A.Ag. developed the study methods. J.A., M.P.-O., A.M.B., J.F., L.C., M.J.B., J.G.-P., M.C., A.P., M.G.-P., E.A.-A., M.T., A.As., N.I.-A., E.M., C.P.-I., J.O., M.C.-O., M.B., P.C., L.H.-G., I.F., H.E.D.T. and J.R.A. were study investigators. M.T.G.M., D.L., J.G.-P. and A.G.C. ensured data verification. J.G.-P., D.L., J.A., M.P.-O., M.T.G.M., J.O., A.M.B. and A.J.C. were responsible for the present secondary statistical analysis. C.B.-I., J.A., M.P.-O., J.G.-P., M.G.-P., A.M.B., L.C., M.C., M.J.B., A.P., J.O., J.F., and J.R.A. supervised the study. C.B.-I. was responsible for funding acquisition. J.A., M.P.-O., J.O., J.G.-P. and A.M.B. wrote the original draft of this Article. All authors reviewed and edited the manuscript, and approved the manuscript for submission. All authors reviewed and approved the original draft. All authors had full access to the full data in the study and accept responsibility to submit for publication.

Data sharing statement

Individual participant data will be made available when the trial is complete, on request to the corresponding authors. After approval of a proposal, data will be shared through a secure online platform.

Declaration of interests

JA has received fees for educational programs from Gilead, MSD, GSK and Janssen outside of the submitted work. MC has participated in advisory boards and has received research funding from GSK, Sanofi Pasteur, Pfizer, Novavax and Janssen. CB-I is the deputy general manager of the Instituto de Salud Carlos III. JRA has received fees from Janssen, outside of the submitted work. AMB is principal investigator of clinical trials sponsored by GlaxoSmithKline, Daiichi-Sankyo, Janssen, and Farmalider, outside of the submitted work. All other authors declare no competing interests.

Acknowledgements

Funded by Instituto de Salud Carlos III (ISCIII). AMB, AJC, JO, and JF are members of the VACCCELERATE (European Corona Vaccine Trial Accelerator Platform) Network, which aims to facilitate and accelerate the design and implementation of COVID-19 phase 2 and 3 vaccine trials. JO is a member of the INsTRuCT (Innovative Training in Myeloid Regulatory Cell Therapy) Consortium, a network of European scientists from academia and industry focused on developing innovative immunotherapies. This work is funded by Instituto de Salud Carlos III, a Spanish public body assigned to the Ministry of Science and Innovation that manages and promotes public clinical research related to public

health. The Spanish Clinical Trials Platform is a public network funded by the Instituto de Salud Carlos III (grant numbers PTC20/00018 and PT17/0017), the State Plan for Research, Development, and Innovation 2013–16, the State Plan for Scientific and Technical Research and Innovation 2017–20, and the Subdirector General for Evaluation and Promotion of Research, Instituto de Salud Carlos III, cofinanced with FEDER funds. CombiVacS was designed under the umbrella of the VACCELERATE project. VACCELERATE and INSTRUCT received funding from the EU's Horizon 2020 Research and Innovation Programme (grant agreement numbers 101037867 and 860003). The Instituto de Salud Carlos III is the Spanish partner in the VACCELERATE project. This work is partially funded by Institute of Health Carlos III (Instituto de Salud Carlos III – ISCIII –), (grants PI19CIII/00004 to JA and PI21CIII/00025 to MPO and JGP), and COVID-19 FUND (grants COV20/00679 and COV20/00072 to MPO and JA) and CIBERINFEC, co-financed by the European Regional Development Fund (FEDER) “A way to make Europe”. The authors thank all trial participants, the international data safety monitoring board (Appendix 1 p 23), and the trial steering committee (Appendix 1 pp 24–25). The authors thank Esther Prieto for editorial assistance and writing support (employed by Hospital Universitario La Paz; funded by the Instituto de Salud Carlos III, grant number PCT20/00018) and María Castillo-de la Osa (PEJ2018-004557-A) for excellent technical assistance.

Supplementary materials

Supplementary material associated with this article can be found in the online version at doi:[10.1016/j.eclim.2022.101529](https://doi.org/10.1016/j.eclim.2022.101529).

Appendix 1. CombiVacS Vaccine Trial Group

FIST NAME	SURNAME
Isabel	Jado
Concepción	Perea
Giovanni	Fedele
Javier	Hernández
María Jesús	Llamas
Francisco	Díez-Fuertes
Esther	Calonge
Almudena	Cascajero
Paloma	Jiménez-Santana
Jana	Baranda
Isabel	Cervera
Marcos	Berges-Buxeda
Pilar	Portolés

(Continued)

FIST NAME	SURNAME
Lucía	Martínez de Soto
Amelia	Rodríguez Mariblanca
Lucía	Díaz García
Elena	Ramírez García
Enrique	Seco Meseguer
Stefan Mark	Stewart Balbás
Alicia	Marín Candón
Irene	García García
Mikel	Urroz Elizalde
Jaime	Montserrat Villatoro
Paula	de la Rosa
Marta	Sanz García
Cristina	López Crespo
Vega	Mauleón Martínez
Raquel	de Madariaga Castell
Laura	Vitón Vara
Julio	García Rodríguez
Antonio	Buño
Eduardo	López Granados
Carmen	Cámara
Esther	Rey Cuevas
Pilar	Ayllon García
María	Jiménez González
Victoria	Hernández Rubio
Paloma	Moraga Alapont
Amparo	Sánchez
Rocío	Prieto
Silvia	Llorente Gómez
Cristina	Miragall Roig
Marina	Aparicio Marlasca
Fernando	de la Calle
Marta	Arsuaga
Blanca	Duque
Susana	Meijide
Aitor	García de Vicuña
Ana	San Expósito
Mikel	Gallego
Dolores	García-Vázquez
Ana Belén	de la Hoz
Gustavo	Pérez-Nanclares
Olaia	Velasco
Josu	Aurrekoetxea
María Angeles	Lázaro
Alejandro	García Castaño
Inés	Urrutia
Rosa	Martínez
Begoña	Calvo
Laura	Saso
Sara	Gómez
Marta	Aldea
Lourdes E	Barón-Mira
M ^a Ángeles	Marcos

(Continued)

FIST NAME	SURNAME
Laura	Granés
Sulayman	Lazaar
Sara	Herranz
Montserrat	Malet
Sebastiana	Quesada
Anna	Vilella
Anna	Llupia
Victoria	Olivé
Antoni	Trilla
Begoña	Gómez
Elisenda	González
Sheila	Romero
Francisco Javier	Gámez
Cristina	Casals
Laura	Burunat
Juan José	Castelló
Patricia	Fernández
Josep Lluís	Bedini
Jordi	Vila
Xavier	Martínez-Gómez
Susana	Otero-Romero
Blanca	Borras-Bermejo
Oleguer	Parés-Badell
Cesar	Llorente
José Angel	Rodrigo-Pendás
Lluís	Armada
Sonia	Uriona
Judit	Riera-Arnau
José	Santos
Carla	Sans-Pola
Lina	Camacho-Arteaga
Aitana	Plaza
Carla	Aguilar
Laia	Pinos
Gisela	Gili
Carmen	Altadill
Gloria	Torres
Esther	Palacio
Margarita	Torrens
Ana	Feliu
Julia	Calonge
Elena Ballarín	Ballarín Alins
Eulàlia	Pérez-Esquiro
Lourdes	Vendrell Bosch
Leonor	Laredo
Manuel	Sanchez-Craviotto
Jose Antonio	Gil-Marín
Esperanza	Gonzalez-Rojano
Natalia	Rodríguez-Galán
Ana Belén	Rivas-Paterna
Teresa	Iglesias
Ouhao	Zhu-Huang

(Continued)

FIST NAME	SURNAME
Angel	Hernández-Bartolomé
Daniel	Lozano-Martín
Verónica	Alvarez-Morales
Oliver	Astasio
Natalia	Pérez-Macías
María Aránzazu	Urrutia-de-la-Plaza
Carmen	Sanz
Agustín	Molina

References

- Borobia AM, Carcas AJ, Pérez-Olmeda M, et al. Immunogenicity and reactogenicity of BNT162b2 booster in ChAdOx1-S-primed participants (CombiVacS): a multicentre, open-label, randomised, controlled, phase 2 trial. *Lancet*. 2021;398:121–130.
- Liu X, Shaw RH, Stuart ASV, et al. Safety and immunogenicity of heterologous versus homologous prime-boost schedules with an adenoviral vectored and mRNA COVID-19 vaccine (Com-COV): a single-blind, randomised, non-inferiority trial. *Lancet*. 2021;398:856–869.
- European Centre for Disease Prevention and Control. SARS-CoV-2 variants of concern as of 7 October 2021. <https://www.ecdc.europa.eu/en/covid-19/variants-concern>. Accessed 14 October 2021.
- Del Rio C, Omer SB, Malani PN. Winter of omicron-the evolving COVID-19 pandemic. *JAMA*. 2021;327:319–320. <https://doi.org/10.1001/jama.2021.24315>.
- Naaber P, Tserel L, Kangro K, et al. Dynamics of antibody response to BNT162b2 vaccine after six months: a longitudinal prospective study. *Lancet Reg Health Eur*. 2021;10:100208. <https://doi.org/10.1016/j.lanepe.2021.100208>.
- Andrews N, Tessier E, Stowe J, et al. Duration of protection against mild and severe disease by Covid-19 vaccines. *N Engl J Med*. 2022;386:340–350.
- Edara VV, Norwood C, Floyd K, et al. Infection- and vaccine-induced antibody binding and neutralization of the B.1.351 SARS-CoV-2 variant. *Cell Host Microbe*. 2021;29:516–521.e3.
- Dejnirattisai W, Zhou D, Supasa P, et al. Antibody evasion by the P.1 strain of SARS-CoV-2. *Cell*. 2021;184:2939–2954.e9.
- Shen X, Tang H, Pajon R, et al. Neutralization of SARS-CoV-2 variants B.1.429 and B.1.351. *N Engl J Med*. 2021;384(24):2352–2354. [NEJMc2103740](https://doi.org/10.1056/NEJMc2103740).
- Madhi SA, Baillie V, Cutland CL, et al. Efficacy of the ChAdOx1 nCoV-19 Covid-19 vaccine against the B.1.351 variant. *N Engl J Med*. 2021;384:1885–1898.
- Dejnirattisai W, Shaw RH, Supasa P, et al. Reduced neutralisation of SARS-CoV-2 omicron B.1.1.529 variant by post-immunisation serum. *Lancet*. 2021;399(10321):234–236. [S0140-6736\(21\)02844-0](https://doi.org/10.1016/S0140-6736(21)02844-0).
- Carreño JM, Alshammari H, Tcheou J, et al. Activity of convalescent and vaccine serum against SARS-CoV-2 Omicron. *Nature*. 2021;602(7898):682–688. <https://doi.org/10.1038/s41586-022-04399-5>.
- Shinde V, Bhikha S, Hoosain Z, et al. Efficacy of NVX-CoV2373 Covid-19 vaccine against the B.1.351 variant. *N Engl J Med*. 2021;384:1899–1909.
- Flaxman A, Marchevsky NG, Jenkin D, et al. Reactogenicity and immunogenicity after a late second dose or a third dose of ChAdOx1 nCoV-19 in the UK: a substudy of two randomised controlled trials (COV001 and COV002). *Lancet*. 2021;398:981–990.
- Sattler A, Angermair S, Stockmann H, et al. SARS-CoV-2-specific T cell responses and correlations with COVID-19 patient predisposition. *J Clin Invest*. 2020;130:6477–6489.
- Grifoni A, Weiskopf D, Ramirez SI, et al. Targets of T cell responses to SARS-CoV-2 coronavirus in humans with COVID-19 disease and unexposed individuals. *Cell*. 2020;181:1489–1501.e15.
- Christie B. Covid-19: early studies give hope omicron is milder than other variants. *BMJ*. 2021;375:n3144.

- 18 Ledford H. How severe are Omicron infections? *Nature*. 2021;600:577–578.
- 19 Bar-On YM, Goldberg Y, Mandel M, et al. Protection of BNT162b2 vaccine booster against Covid-19 in Israel. *N Engl J Med*. 2021;385(15):1393–1400. <https://doi.org/10.1056/NEJMoa2114255>.
- 20 Kamar N, Abravanel F, Marion O, Couat C, Izopet J, Del Bello A. Three doses of an mRNA Covid-19 vaccine in solid-organ transplant recipients. *N Engl J Med*. 2021;385:661–662.
- 21 Hall VG, Ferreira VH, Ku T, et al. Randomized trial of a third dose of mRNA-1273 vaccine in transplant recipients. *N Engl J Med*. 2021;385:1244–1246.
- 22 European Medicines Agency (EMA). *Comirnaty and Spikevax: EMA Recommendations on Extra Doses and Boosters*. 2021. <https://www.ema.europa.eu/en/news/comirnaty-spikevax-ema-recommendations-extra-doses-boosters>. Accessed 15 November 2021.
- 23 Meyer B, Torriani G, Yerly S, et al. Validation of a commercially available SARS-CoV-2 serological immunoassay. *Clin Microbiol Infect*. 2020;26:1386–1394.
- 24 Gilbert PB, Montefiori DC, McDermott AB, et al. Immune correlates analysis of the mRNA-1273 COVID-19 vaccine efficacy clinical trial. *Science*. 2022;375:43–50.
- 25 Widge AT, Roupael NG, Jackson LA, et al. Durability of responses after SARS-CoV-2 mRNA-1273 vaccination. *N Engl J Med*. 2021;384:80–82.
- 26 Goel RR, Painter MM, Apostolidis SA, et al. mRNA vaccines induce durable immune memory to SARS-CoV-2 and variants of concern. *Science*. 2021;374(6572):eabm0829.
- 27 Liu X, Shaw RH, Stuart ASV, et al. Safety and immunogenicity of heterologous versus homologous prime-boost schedules with an adenoviral vectored and mRNA COVID-19 vaccine (Com-COV): a single-blind, randomised, non-inferiority trial. *Lancet*. 2021;398:856–869.
- 28 Normark J, Vikström L, Gwon YD, et al. Heterologous ChAdOx1 nCoV-19 and mRNA-1273 vaccination. *N Engl J Med*. 2021;385:1049–1051.
- 29 Stirrup O, Krutikov M, Tut G, et al. SARS-CoV-2 anti-spike antibody levels following second dose of ChAdOx1 nCoV-19 or BNT162b2 in residents of long-term care facilities in England (VIVALDI). *J Infect Dis*. 2022;jiac146. <https://doi.org/10.1093/infdis/jiac146>.
- 30 Baumgarth N, Nikolich-Zugich J, Lee FE-H, Bhattacharya D. Antibody responses to SARS-CoV-2: let's stick to known knowns. *J Immunol*. 2020;205:2342–2350.
- 31 Voysey M, Costa Clemens SA, Madhi SA, et al. Single-dose administration and the influence of the timing of the booster dose on immunogenicity and efficacy of ChAdOx1 nCoV-19 (AZD1222) vaccine: a pooled analysis of four randomised trials. *Lancet*. 2021;397:881–891.
- 32 Sokal A, Chappert P, Barba-Spaeth G, et al. Maturation and persistence of the anti-SARS-CoV-2 memory B cell response. *Cell*. 2021;184:1201–1213.e14.
- 33 Viana R, Moyo S, Amoako DG, et al. Rapid epidemic expansion of the SARS-CoV-2 Omicron variant in southern Africa. *Nature*. 2022;603:679–686.
- 34 Garcia-Beltran WF, St Denis KJ, Hoelzemer A, et al. mRNA-based COVID-19 vaccine boosters induce neutralizing immunity against SARS-CoV-2 Omicron variant. *Cell*. 2022;185:457–466.
- 35 Tartof SY, Slezak JM, Puzniak L, et al. Durability of BNT162b2 vaccine against hospital and emergency department admissions due to the omicron and delta variants in a large health system in the USA: a test-negative case-control study. *Lancet Respir Med*. 2022;S2213-2600(22):1–11. [https://doi.org/10.1016/S2213-2600\(22\)00101-1](https://doi.org/10.1016/S2213-2600(22)00101-1).