



HAL
open science

Sotrovimab retains activity against SARS-CoV-2 omicron variant BQ.1.1 in a non-human primate model

Cécile Hérate, Romain Marlin, Franck Touret, Nathalie Dereuddre-Bosquet, Flora Donati, Francis Relouzat, Laura Junges, Mathilde Galhaut, Océane Dehan, Quentin Sconosciuti, et al.

► To cite this version:

Cécile Hérate, Romain Marlin, Franck Touret, Nathalie Dereuddre-Bosquet, Flora Donati, et al.. Sotrovimab retains activity against SARS-CoV-2 omicron variant BQ.1.1 in a non-human primate model. *Heliyon*, 2023, 9 (6), pp.e16664. 10.1016/j.heliyon.2023.e16664 . hal-04584291v2

HAL Id: hal-04584291

<https://hal.science/hal-04584291v2>

Submitted on 24 May 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License



Sotrovimab retains activity against SARS-CoV-2 omicron variant BQ.1.1 in a non-human primate model

Cécile Hératé^{a,1}, Romain Marlin^{a,1}, Franck Touret^b, Nathalie Dereuddre-Bosquet^a, Flora Donati^{c,d}, Francis Relouzat^a, Laura Junges^a, Mathilde Galhaut^a, Océane Dehan^{c,d}, Quentin Sconosciuti^a, Antoine Nougairède^b, Xavier de Lamballerie^b, Sylvie van der Werf^{c,d}, Roger Le Grand^{a,*}

^a Université Paris-Saclay, Inserm, CEA, Center for Immunology of Viral, Auto-immune, Hematological and Bacterial Diseases (IMVA-HB/IDMIT), 18 route du Panorama, 92265, Fontenay-aux-Roses, France

^b Unité des Virus Émergents (UVE), Aix Marseille Université, IRD 190, INSERM 1207, 27 Bd Jean Moulin, 13005, Marseille, France

^c Institut Pasteur, Université Paris Cité, CNRS UMR 3569, Molecular Genetics of RNA Viruses Unit, 25-28 Rue du Dr Roux, 75015, Paris, France

^d Institut Pasteur, Université Paris Cité, National Reference Center for Respiratory Viruses, 25-28 Rue du Dr Roux, 75015, Paris, France

ARTICLE INFO

Keywords:

SARS-CoV2

Sotrovimab

Non human primate

BQ.1.1

ABSTRACT

The SARS-CoV2 Omicron variants have acquired new Spike mutations leading to escape from the most of the currently available monoclonal antibody treatments reducing the options for patients suffering from severe Covid-19. Recently, both *in vitro* and *in vivo* data have suggested that Sotrovimab could retain partial activity against recent omicron sub-lineage such as BA.5 variants, including BQ.1.1. Here we report full efficacy of Sotrovimab against BQ.1.1 viral replication as measure by RT-qPCR in a non-human primate challenge model.

1. Results

Circulating SARS-CoV-2 Omicron variants have acquired mutations in the receptor-binding domain (RBD) resulting in higher ACE2-binding affinity. These changes are also associated with increasing transmission efficiency and escape to pre-existing neutralizing antibodies. In Europe, the major circulating variant BQ.1.1, derived from BA.5, is escaping most of the available *anti*-SARS-CoV-2 monoclonal antibody treatments. Recent *in vitro* data are controversial with some suggesting that Sotrovimab binds Omicron sub-variants, promotes Fc dependent effector functions, and still has the capacity to neutralize them [1]. While most of the *in vitro* studies conducted on cell lines, mainly VeroE6 cells, report a drastic decrease of Sotrovimab inhibition effect on recent omicron subvariant including BQ.1.1 [2–4].

Moreover, treatment of S309 (Sotrovimab parent antibody; 10 or 30 mg/kg) in mice or with Sotrovimab (7 or 14 mg/kg) in hamsters provided protection from a BQ.1.1 challenge [1,5]. However, further data are needed to confirm Sotrovimab *in vivo* activity and pharmacokinetics and to determine whether it should remain, in addition to Nirmatrelvir-Ritonavir in the list of available treatments for patients at risk [6].

* Corresponding author. Université Paris-Saclay, Inserm, CEA, Fontenay-aux-Roses, France, 18 route du Panorama, 92265, Fontenay-aux-Roses, France.

E-mail address: roger.le-grand@cea.fr (R. Le Grand).

¹ equally contributed to this work.

<https://doi.org/10.1016/j.heliyon.2023.e16664>

Received 13 March 2023; Received in revised form 9 May 2023; Accepted 24 May 2023

Available online 30 May 2023

2405-8440/© 2023 Université Paris-Saclay. Published by Elsevier Ltd. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

In this controversial context and to offer new possibilities to clinicians, we conducted an efficacy study in a well characterized SARS-CoV2 NHP model [7–10]. Here we report Sotrovimab efficacy in a non-human primate (NHP) model of SARS-CoV-2 infection. Three female cynomolgus macaques (*Macaca fascicularis*) aged 14–15 years and weighing between 4.6 and 7.3 kg were treated with 10 mg/kg of Sotrovimab (Xevudy) intravenously 96 h prior to viral challenge. Treated animals and two additional controls were challenged with 1×10^5 pfu of SARS-CoV-2 BQ.1.1 (hCoV-19/France/IDF-IPP50823/2022 - EPI_ISL_15195982) via combined intranasal and intratracheal routes using an experimental protocol previously reported for other variants [10].

The treatment was carried out without any adverse effects being recorded. Sotrovimab was measured in the serum of NHP at days 1, 4, 8 and 11 post-treatment, showing a similar exposure profile (Fig. 1A) as observed in humans during the COMET-ICE trial [11]. Lymphopenia was reported at 2 days post-challenge (d.p.c) in non-treated animals and in 2 out of 3 treated animals as expected in this model [10]. In this study, to empower statistical analysis, we included 4 historical controls challenged with the same batch of BQ.1.1 SARS-CoV2 virus stock at a similar dose to the 2 concomitant control NHPs. Efficacy was monitored by genomic viral RNA (gRNA) quantification using RT-qPCR (Fig. 1B). In untreated animals, gRNA was detected in tracheal fluids collected with swabs, with a median peak viral loads at 2–3 d.p.c of $6.11 \log_{10}$ copies/mL. Viral gRNA was also detected at 3 d.p.c in the broncho-alveolar lavages (BAL) with a median value of $5.55 \log_{10}$ copies/mL. At 10 d.p.c, virus was still detectable in BAL in 5 control NHPs out of 6. By contrast, the three treated animals had viral gRNA below the limit of detection both in trachea (Fig. 1B) and BAL (Fig. 1C). Comparisons of the area under the curve (D0-D14) for tracheal viral load kinetics as well as BAL viral load at day 3 reveal a significant difference ($p = 0.0238$) between controls and Sotrovimab treated animals.

Sotrovimab had previously been withdrawn from the therapeutic panel due to its initially proposed poor *in vitro* efficacy against Omicron variants, in particular BA.2 [12]. Here, we demonstrate that Sotrovimab inhibits viral replication of BQ.1.1 in NHPs both in upper and lower respiratory tracts when given for prophylaxis. Our results support the ability of Sotrovimab to prevent infection with SARS-CoV-2 (BQ.1.1 variant) following inoculation therefore it may have potential activity against this variant. Further studies of use of Sotrovimab as a treatment for variant BQ.1.1 are required. Our results thus support the use of Sotrovimab in humans against BQ.1.1, in case of ineligibility to Nirmatrelvir-Ritonavir.

Author contribution statement

Cécile Hérate: Conceived and designed the experiments; Analyzed and interpreted the data; Wrote the paper.
 Romain Marlin: Conceived and designed the experiments; Wrote the paper.
 Franck Touret; Laura Junges; Mathilde Galhaut; Quentin Sconosciutti: Performed the experiments.
 Nathalie Dereuddre-Bosquet: Conceived and designed the experiments; Analyzed and interpreted the data.
 Flora Donati; Océane Dehan; Antoine Nougairède: Analyzed and interpreted the data.
 Francis Relouzat: Conceived and designed the experiments; Analyzed and interpreted the data.
 Xavier de Lamballerie; Sylvie van der Werf; Roger Le Grand, P.h.D, D.V.M: Conceived and designed the experiments.

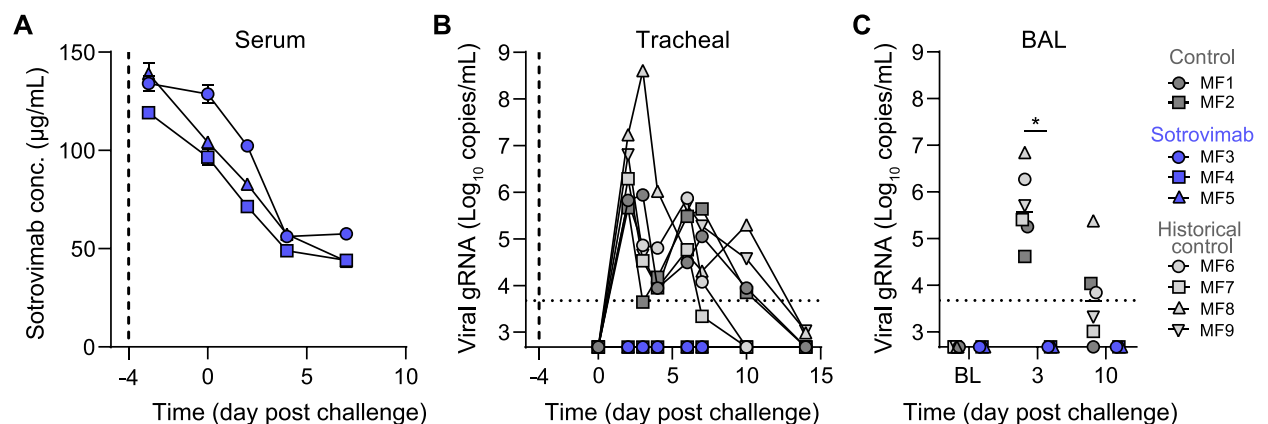


Fig. 1. Pharmacokinetics of Sotrovimab in serum and viral loads in the respiratory tract of BQ.1.1 SARS-CoV-2 exposed cynomolgus macaques treated with Sotrovimab. Animals MF3, MF4 and MF5 were treated 4 days before challenge (blue) while MF1 and MF2 were not treated (dark grey). MF6, MF7, MF8 and MF9 were added as non-treated historical controls (light grey) A. Pharmacokinetics of Sotrovimab in serum. NHPs were challenged 4 days after Sotrovimab injection. B. Genomic viral RNA (gRNA) was measured in tracheal fluids collected with swabs during the acute phase of infection. C. BAL gRNA was analyzed at baseline, day 3 and day 10 post challenge. Horizontal dotted lines represent the limit of quantification ($2.7 \log_{10}$ copies/mL). Vertical dotted lines indicate the day of Sotrovimab treatment. Two-tailed non-parametric Mann-Whitney tests were used for comparisons. * $p < 0.05$. (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

Data availability statement

Data will be made available on request.

Additional information

No additional information is available for this paper.

Declaration of competing interest

The authors have declared that no conflict of interest exists.

Acknowledgment

The Infectious Disease Models and Innovative Therapies (IDMIT) research infrastructure is supported by the “Programme Investissements d’Avenir” managed by ANR under reference ANR-11-INBS-0008. The NHP experiment was part of BIOVAR and PRODEVA programs funded by the ANRS-MIE project EMERGEN (ANRS0151).

Appendix

Materials and methods

Ethics and biosafety statement

Five female cynomolgus macaques (*M. fascicularis*), aged 14–15 years and originating from Mauritian AAALAC-certified breeding centres were included to this study. Four female cynomolgus macaques from Mauritian AAALAC-certified breeding centres, aged 14–15 years were also added as historical controls. All animals were housed in IDMIT facilities (CEA, Fontenay-aux-roses), under BSL-3 containment (Animal facility authorization #D92-032-02, Préfecture des Hauts de Seine, France) and in compliance with European Directive 2010/63/EU, the French regulations and the Standards for Human Care and Use of Laboratory Animals, of the Office for Laboratory Animal Welfare (OLAW, assurance number #A5826-01, US). The protocols were approved by the institutional ethical committee “Comité d’Ethique en Expérimentation Animale du Commissariat à l’Energie Atomique et aux Energies Alternatives” (CETE #44) under statement number A20-066 and A22-006. The study was authorized by the “Research, Innovation and Education Ministry” under registration number respectively APAFIS#29191–2021011811505374 v1 and APAFIS# 36939–2022042217237124 v1.

Antibody treatment

Sotrovimab (Xevudy 500 mg solution to dilute for perfusion) is a commercial antibody also named VIR-7831 and GSK4182136. Recommendations for humans consists in 500 mg by intravenous route, equivalent to 10 mg/kg for a patient of 50 kg. For NHPs, the treatment was administrated 96 h prior challenge by intravenous perfusion in 30 min. Sotrovimab was diluted at 1.434 mg/mL in NaCl 0.9% and injected at a dose of 10 mg/kg, which represents a volume between 34 and 50 mL.

Animals were monitored for heart rate, respiratory rate and oximetry every 10 min from treatment initiation until 30 min after the end of injection.

Quantification of Sotrovimab monoclonal antibody

Sotrovimab exposure was measured using the commercial enzyme-linked immunosorbent assay (ELISA) *anti*-SARS-CoV-2 Quantivac (IgG) kit (Euroimmun) which is directed against the S1 domain of the spike protein. Results were expressed in binding antibody units per mL (BAU/mL) following manufacturer instructions and converted to µg/mL using blank plasma from untreated/infected animals spiked with known quantities of Sotrovimab.

SARS-CoV-2 challenge

Treated and control animals were challenged with SARS-CoV-2 BQ.1.1 (hCoV-19/France/IDF-IPP50823/2022 - EPI_ISL_15195982), provided by the National Reference Center for respiratory viruses at Institut Pasteur, via the combination of intranasal (1/10) and intratracheal (9/10) routes (day 0), using atropine (0.04 mg/kg) for pre-medication and ketamine (5 mg/kg) with medetomidine (0.05 mg/kg) for anesthesia. Tracheal fluids and blood samples were regularly collected following challenge. Broncho alveolar lavages (BAL) were performed at 3 and 10 days post challenge. NHPs were followed for behavior assessment and clinical score during the 7 days of the infection. Blood cell counts, hemoglobin and hematocrit were determined from EDTA blood using a DXH800 analyzer (Beckman Coulter).

Viral quantification

Viral genomic RNA (gRNA) was quantified in swabs and BAL samples by RT-qPCR with a plasmid standard concentration range. The protocols describing the procedure for the detection of SARS-CoV-2 is available on the WHO website (https://www.who.int/docs/default-source/coronaviruse/real-time-rt-pcr-assays-for-the-detection-of-sars-cov-2-institut-pasteur-paris.pdf?sfvrsn=3662fcb6_2).

Statistical analysis

Data were collected using classical Excel files (Microsoft Excel 2016). Differences between unmatched groups were compared using the Mann–Whitney *U* test (Graphpad Prism 8.0).

References

- [1] A. Addetia, L. Piccoli, J.B. Case, Y.-J. Park, M. Beltramello, B. Guarino, et al., Therapeutic and vaccine-induced cross-reactive antibodies with effector function against emerging Omicron variants, *bioRxiv* (2023), 2023.01.17.523798.
- [2] D. Planas, T. Bruel, I. Staropoli, F. Guivel-Benhassine, F. Porrot, P. Maes, et al., Resistance of Omicron subvariants BA.2.75.2, BA.4.6, and BQ.1.1 to neutralizing antibodies, *Nat. Commun.* 14 (1) (2023) 824.
- [3] F. Touret, E. Giraud, J. Bourret, F. Donati, J. Tran-Rajau, J. Chiaravalli, et al., Enhanced neutralization escape to therapeutic monoclonal antibodies by SARS-CoV-2 omicron sub-lineages, *iScience* 26 (4) (2023), 106413.
- [4] P. Arora, A. Kempf, I. Nehlmeier, S.R. Schulz, H.M. Jäck, S. Pöhlmann, et al., Omicron sublineage BQ.1.1 resistance to monoclonal antibodies, *Lancet Infect. Dis.* 23 (1) (2023) 22–23.
- [5] J.-S. Driouich, O. Bernardin, F. Touret, X. de Lamballerie, A. Nougairède, *bioRxiv* (2023).
- [6] M. Cox, T.P. Peacock, W.T. Harvey, J. Hughes, D.W. Wright, B.J. Willett, et al., SARS-CoV-2 variant evasion of monoclonal antibodies based on in vitro studies, *Nat. Rev. Microbiol.* 21 (2) (2023) 112–124.
- [7] P. Maisonnasse, Y. Aldon, A. Marc, R. Marlin, N. Dereuddre-Bosquet, N.A. Kuzmina, et al., COVA1-18 neutralizing antibody protects against SARS-CoV-2 in three preclinical models, *Nat. Commun.* 12 (1) (2021) 6097.
- [8] P. Maisonnasse, J. Guedj, V. Contreras, S. Behillil, C. Solas, R. Marlin, et al., Hydroxychloroquine use against SARS-CoV-2 infection in non-human primates, *Nature* 585 (7826) (2020) 584–587.
- [9] N. Zabaleta, W. Dai, U. Bhatt, C. Hérate, P. Maisonnasse, J.A. Chichester, et al., An AAV-based, room-temperature-stable, single-dose COVID-19 vaccine provides durable immunogenicity and protection in non-human primates, *Cell Host Microbe* 29 (9) (2021), 1437–53.e8.
- [10] C. Fenwick, P. Turelli, D. Ni, L. Perez, K. Lau, C. Hérate, et al., Patient-derived monoclonal antibody neutralizes SARS-CoV-2 Omicron variants and confers full protection in monkeys, *Nat Microbiol* 7 (9) (2022) 1376–1389.
- [11] Agency EM (2021).
- [12] T. Bruel, K. Stéfic, Y. Nguyen, D. Toniutti, I. Staropoli, F. Porrot, et al., Longitudinal analysis of serum neutralization of SARS-CoV-2 Omicron BA.2, BA.4, and BA.5 in patients receiving monoclonal antibodies, *Cell Rep Med* 3 (12) (2022), 100850.