Effects of casirivimab/imdevimab on systemic and mucosal immunity against SARS-CoV-2 in B-cell depleted patients with autoimmune rheumatic diseases refractory to vaccination

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B-cell depleting therapy is a mainstay to treat autoimmune rheumatic diseases characterised by autoantibody production. While B cell depletion effectively turns down autoimmune inflammation, it also hampers the development of protective immunity after infections and vaccination. The latter has become of particular concern, as treatment with anti-CD20 antibody rituximab is associated with more severe courses of COVID-19 and abolished or severely impaired humoral responses to SARS-CoV-2 vaccination.

While newly initiated B-cell depletion is usually preceded by SARS-CoV-2 vaccination, several situations leave patients without protective immunity during treatment. These include (1) pre-existing B-cell depletion before vaccination was available, (2) vaccination refusal and (3) loss of vaccination response during treatment. In addition, B-cell depletion is often long-lasting and needs to be repeated to prevent potentially life-threatening flares. Therefore, strategies need to be developed providing anti-SARS-CoV-2 immunity in such patients. Several neutralising monoclonal antibodies (mAbs) against spike S1 protein of SARS-CoV-2 have been developed. Casirivimab and imdevimab are two recombinant IgG1 mAbs that bind the receptor-binding domain of the SARS-CoV-2 spike protein and prevent its binding to the angiotensin-converting enzyme 2 at low concentrations (31pM). Use of casirivimab/imdevimab combination has been shown to reduce hospitalisations and deaths from COVID-19.

We analysed 22 patients with autoimmune rheumatic diseases who received pre-exposure prophylaxis (PrEP) with a single subcutaneous injection of casirivimab/imdevimab (600mg/600mg) in December 2021 due to lack of anti-SARS-CoV-2 IgG at least 21 days after three SARS-CoV-2 vaccinations. Patients’ characteristics are shown in figure 1. We obtained 92 serum and 75 saliva samples at 4 consecutive timepoints. Serum and salivary anti-SARS-CoV-2 Spike IgG were quantified by ELISA (Euroimmun, Lübeck, Germany) at baseline and 1, 14 and 30 days after administration. IgG levels are given as antibody ratios by dividing the optical density of the sample by that of the calibrator. Serum anti-SARS-CoV-2 IgG, which were absent at baseline (mean ratio±SD 0.2±0.1; cut-off for positive>1.1), reached maximal concentrations on day 1 (8.4±0.8) after administration and remained stable on day 14 (7.5±1.3) and 30 (8.6±0.8). Anti-SARS-CoV-2 IgG serum levels 30 days after PrEP were significantly lower (p<0.001) than after vaccination of healthy individuals (11.1±1.4) or after infection and vaccination (12.4±0.9). Furthermore, anti-SARS-CoV-2 IgG were also found in saliva on day 1 (1.2±0.7), with further increases on day 14 (2.8±1.4) and 30 (3.9±2.2). Salivary anti-SARS-CoV-2 IgG at day 30 were significantly higher (p<0.05) than in vaccinated (1.4±1.4) or infected and vaccinated controls (1.9±1.7). No side effects were reported. Five patients (19.2%) had a close contact with a person infected with SARS-CoV-2, after which all but one remained PCR negative. The fifth patient turned PCR positive and developed mild fever and cough.

These data show that PrEP with casirivimab/imdevimab is safe and provides a fast induction of anti-SARS-CoV-2 humoral immunity in B-cell depleted patients failing previous vaccination. Furthermore, substantial levels of casirivimab/imdevimab were found in saliva, exceeding the levels of anti-SARS-CoV-2-specific IgG observed in healthy controls after infection.
Notably, these data were obtained during the wave of the SARS-CoV-2 Delta variant in Germany, for which casirivimab/imdevimab is highly effective. For the Omicron variants casirivimab and imdevimab are less effective, although imdevimab retained some in vitro efficacy against the BA.2 subvariant.5 Recently, the long-acting tixagevimab/cilgavimab has been approved as PrEP for all Omicron variants.6 Whether tixagevimab/cilgavimab reaches similar serum and saliva concentrations in B-cell depleted patients with autoimmune rheumatic diseases remains to be determined.

Contributors FF, KS, AK, DS, GS and TH were involved in study design. Sample collection was done by DB, LV-M, FH, KT, KM, BM, AK, DS and GS. Experiments and data analysis were performed by FF, KS, KT and TH. FF, KS, GS and TH were responsible for the figure. Data interpretation was done by FF, KS, DB, LV-M, KT, AK, DS, GS and TH. Writing of the manuscript was done by FF, KS, AK, DS and TH. GS and TH contributed equally and share senior coauthorship.

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