Vaccines for SARS-CoV-2 are effective in patients with systemic rheumatic disease (SRD) without exhibiting significant safety issues or causing disease flares, whereas two doses of mRNA vaccines lead to significantly better outcomes of breakthrough (ie, despite vaccination) COVID-19 compared with unvaccinated patients. Since a third (booster) dose is deemed necessary for better immunisation, we aimed to examine hospitalisation rates and mortality of breakthrough COVID-19 in patients with SRD who had received three doses of the vaccine (booster-vaccinated), compared with those who received two doses (fully vaccinated) or were unvaccinated. We also comparatively assessed breakthrough COVID-19 outcomes in booster-vaccinated individuals with or without SRDs.

We prospectively recorded date/type of vaccination, demographic, clinical and COVID-19-related features (date of infection, duration of self-reported symptomatology, hospitalisation, need for non-invasive ventilation and death), in consecutive SARS-CoV-2-infected patients followed up in our department from March 2020 (onset of pandemic) to February 2022. We also included as controls consecutive booster-vaccinated healthcare workers and patients' friends/relatives without SRD. Only individuals in whom breakthrough COVID-19 occurred ≥14 days after the second or third vaccination were enrolled in the study. Mean±SD was used for continuous variables and percentages (%) for categorical variables. Fisher’s exact test and Mann-Whitney test were used.

A total of 65 booster-vaccinated, 36 fully vaccinated and 60 unvaccinated patients with SRD, as well as 80 booster-vaccinated individuals without SRD, were enrolled. Demographic, clinical and treatment characteristics were similar across groups compared, except...
for lung disease, which was more common in the fully vaccinated patients compared with the other patient groups and the control group (online supplemental tables 1 and 2). COVID-19-related hospitalisations were less common in booster-vaccinated (2/65, 3.1%) than in fully vaccinated (5/36, 13.9%, p=0.09) or unvaccinated patients (18/60, 30.0%, p<0.0001). While 4/60 (6.7%) unvaccinated patients died, there were no deaths in the booster-vaccinated and fully vaccinated patient groups (figure 1). Moreover, clinical outcomes of breakthrough COVID-19 were comparable between booster-vaccinated patients with SRD and individuals without SRD (deaths: 0% for both groups, hospitalisations: 1.25% for individuals without SRD vs 3.1% for patients with SRD), except for duration of COVID-19 symptomatology, which was longer in patients with SRD than in controls (6.1±3.2 vs 4.9±3.1 days, p=0.01) (online supplemental table 2).

Therefore, in concert with studies examining data from the general population, we show that booster vaccination further reduces the frequency of COVID-19-related hospital admissions and deaths in people with SRDs. Notably, comparisons in small-scale, breakthrough infections following booster vaccination in individuals with and without SRDs show that outcomes are comparable between the two groups. This was true, despite the higher frequency of adverse prognostic factors for COVID-19, like age and lung disease, in SRD patients compared with the group of individuals without SRD. Our study has certain limitations. First, COVID-19 in this cohort occurred at different time points of the pandemic; thus, the possibility that different variants of SARS-CoV2 have infected our patients over the entire study period cannot be excluded. Of note, booster-vaccinated patients were enrolled during the same time period when both Delta and Omicron variants were prevalent. Second, antibody response, which might have been affected by immunosuppressive/immunomodulatory treatments, was not measured.

To conclude, these results suggest that booster COVID-19 vaccination has beneficial effects in patients with SRDs, on par with what has been shown for the general population. This, in combination with the reassuring results about the safety of vaccines, argues in favour of booster vaccination in patients with SRD.

Correction notice This article has been corrected since it was first published online. Maria G Tektonidou was incorrectly listed as Maria GG Tektonidou.

Contributors Study conception and design: PPS, MGT and GEF. Critical revision and approval of the manuscript: PPS. Data acquisition and drafting of the manuscript: all authors.

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests None declared.

Patient consent for publication Not applicable.

Provenance and peer review Not commissioned; externally peer reviewed.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/.

ORCID ids
George E Fragoulis http://orcid.org/0000-0003-4932-7023
Vasiliki-Kalliopi Bournia http://orcid.org/0000-0002-2162-1362
Gerasimos Evangelatou http://orcid.org/0000-0003-3822-3093
Antonis Fanourakis http://orcid.org/0000-0003-2969-031X
Evrydiki Kravvariti http://orcid.org/0000-0003-4330-3266
Maria G Tektonidou http://orcid.org/0000-0003-2238-0975

REFERENCES
Correction: Clinical outcomes of breakthrough COVID-19 after booster vaccination in patients with systemic rheumatic diseases


This article has been corrected since it was first published online. Maria G Tektonidou was incorrectly listed as Maria GG Tektonidou.

**Open access** This is an open access article distributed in accordance with the Creative Commons Attribution 4.0 Unported (CC BY 4.0) license, which permits others to copy, redistribute, remix, transform and build upon this work for any purpose, provided the original work is properly cited, a link to the licence is given, and indication of whether changes were made. See: https://creativecommons.org/licenses/by/4.0/.

© Author(s) (or their employer(s)) 2022. Re-use permitted under CC BY. Published by BMJ.

*RMD Open* 2022;8:e002279corr1. doi:10.1136/rmdopen-2022-002279corr1

[Check for updates]