New onset pemphigus foliaceus following AstraZeneca COVID-19 vaccination

Dear Editor,

Pemphigus vulgaris (PV) and pemphigus foliaceus (PF) are a rare group of immunobullous disorders that can lead to high morbidity and mortality. Analogous to the other autoimmune diseases, pemphigus is closely related to the immune response, which is impacted by several factors including; polymorphism of the genes, family history of other autoimmune disorders including pemphigus, gender, ethnicity, geographical area and environmental factors. Several triggers including different drugs and treatments, diseases, vaccines, nutrients, micronutrients, pregnancy and stress have all been implicated in the aetiology of the disease. Here, we present a case with PF whose disease developed after administration of the first dose of ChAdOx1 nCoV-19 (AstraZeneca) vaccination and was exacerbated following the second dose of this vaccine.

A 62-year-old woman in good health with no significant past medical history and no previous skin disease developed a generalized erythematous itchy rash 1 week after the first dose of AstraZeneca COVID-19 vaccination. Administration of the second dose of the same vaccine 12 weeks later resulted in significant skin worsening within 2 days, with extensive scaling and erythema. The patient was not taking any regular or new medications and did not have any family history of autoimmune conditions or skin problems. Physical examination demonstrated large erosive annular erythematous plaques on her face, trunk and limbs (Figure 1a). No mucosal involvement was present.

Histology demonstrated subcorneal acantholysis and a blister filled with neutrophils (Figure 1b). Direct immunofluorescence (DIF) revealed intercellular IgG in the epidermis in a chicken wire-like pattern (Figure 1c). A tissue biopsy for histology and DIF in combination with clinical findings remains the best diagnostic tool for pemphigus. Histology reveals acantholysis with suprabasal blistering in PV and subcorneal blistering in PF. DIF reveals intercellular IgG and C3 epidermal depositions in a chicken wire-like pattern. Although PV has the same DIF pattern and

FIGURE 1 (a) Large erosive annular erythematous plaques on the back. (b) Histology of the skin shows subcorneal acantholysis and blister filled with neutrophils. The epidermis is infiltrated by a large number of neutrophils. H and E stain ×100 magnification. (c) Direct immunofluorescent staining shows intercellular IgG in the epidermis in a chicken wire-like pattern. IMF stain ×200 magnification.
LETTER TO THE EDITOR

LETTER TO THE EDITOR

type of deposition, in PF the fluorescence signal is stronger in the upper epidermis. Based on the characteristic clinical and histological findings in this case, the diagnosis was confirmed as new onset Pemphigus Foliaceus following COVID-19 AstraZeneca vaccination. She was started on Prednisolone 0.75 mg/kg/day and mycophenolate mofetil as a steroid-sparing agent. This approach resulted in significant improvement of skin changes.

ChAdOx1 nCoV-19 (AstraZeneca) is an adenovirus-vectored vaccine, which has been proven to generate robust neutralising antibody and cellular immune responses against the SARS-CoV-2 spike glycoprotein. The main proposed mechanism for AstraZeneca vaccine-induced pemphigus could be a hyper-immune reaction in genetically predisposed individuals, with eventual formation of anti-desmoglein antibodies. An alternative hypothesis is that vaccine components could act as foreign antigens resulting in a cross-reaction with pemphigus antigens, as previously postulated with other immunizations.

There have been numerous reports of relapse or exacerbation of patients with a confirmed diagnosis of pemphigus following COVID-19 vaccination. Twelve cases of new emergence of pemphigus that are believed to be associated with the administration of various COVID-19 vaccines have also been reported in the literature (Table 1). The reported time interval between the vaccination and the occurrence of pemphigus is between 2 days to 1 month. Previous reports refer to only one dose of the vaccine, which includes either the first dose with no information on the second dose or the second dose only. This pattern of reaction with a low probability score might suggest that vaccination coincidentally occurs shortly before the pemphigus appears. This may, therefore, raise the question as to whether a true relationship between COVID-19 vaccines and the new occurrence of pemphigus actually exists. However, the case we present here, as well as the case reported by Solimani et al. help shed some light on this concern. We believe that the close association of COVID-19 vaccination with the acute onset of pemphigus in our patient, as well as the exacerbation following subsequent administration of the vaccine, is more than a coincidence. This is supported by a probability score of 8 (possible causal reaction) on the Naranjo Adverse Drug Reaction Probability Scale.

Although a direct pathological link between the COVID-19 vaccines and the onset of pemphigus is not yet identified, continued observation and documentation of true adverse events of the vaccination is essential.

ETHICS STATEMENT

The patients in this manuscript have given written informed consent to the publication of their case details.

FUNDING SOURCES

None.

CONFLICT OF INTEREST

None.

### TABLE 1

| Age/sex | Type of pemphigus | Type of vaccine                      | Time to onset                      | Reference                        |
|---------|-------------------|--------------------------------------|------------------------------------|----------------------------------|
| 38 F    | PV                | ChAdOx1 nCoV-19 (Astra Zeneca)        | 1 week after the first dose, No data on the second dose | Thongprasom K, Oral Dis, Sep 2021 |
| 40 F    | PV                | BNT162b2 (Pfizer-BioNTech)            | 5 days after the first dose, worsening 3 days after the second dose | Solimani et al.                 |
| 60 M    | PV                | mRNA-1273 (Moderna)                   | 1 week after the second dose        | Koutlas IG, Oral Dis, Nov 2021   |
| 34 M    | PV                | ChAdOx1 nCoV-19 (Astra Zeneca)        | Few days after the first dose, No data on the second dose | Hatami P, Dermatol Ther, Jan 2022 |
| 83 M    | PF                | BNT162b2 (Pfizer-BioNTech)            | 2 days after the second dose         | Lua A, Australas J Dermatol, Feb 2022 |
| 69 F    | PV                | Inactivated vaccine (CoronaVac)       | 1 week after the second dose        | Akoglu G, Dermatol Ther, Feb 2022 |
| 44 M    | PV                | ChAdOx1 nCoV-19 (Astra Zeneca)        | 1 week after the second dose        | Singh A, J Cosmet Dermatol, March 2022 |
| 76 F    | PV                | BBIBP-CorV (Sinopharm)                | 1 month after the second dose       | Saffarian Z, Dermatol Ther, March 2022 |
| 50 F    | PF                | BNT162b2 (Pfizer-BioNTech)            | 15 days after the second dose       | Hali F, Cureus, March 2022       |
| 58 F    | PV                | BNT162b2 (Pfizer-BioNTech)            | 1 month after the first dose, no data on the second dose | Calabria E, Pathol Res Pract, April 2022 |
| 60 F    | PV                | BNT162b2 (Pfizer-BioNTech)            | 1 week after the second dose         | Knechtl GV, J Eur Acad Dermatol Venereol, April 2022 |
| 89 M    | PV                | BNT162b2 (Pfizer-BioNTech)            | 1 month after the second dose       | Knechtl GV, J Eur Acad Dermatol Venereol, April 2022 |

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FUNDING SOURCES

None.

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None.
LETTER TO THE EDITOR

DATA AVAILABILITY STATEMENT
Data sharing not applicable – no new data generated, or the article describes entirely theoretical research.

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