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**Title:** The first dose of COVID-19 vaccine may trigger pemphigus and bullous pemphigoid flares: is the second dose therefore contraindicated?

**Running head:** COVID-19 vaccine and autoimmune blistering diseases

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*Dear Editor,*

during the COVID-19 pandemics the management of immunosuppressed patients, such as those affected by autoimmune bullous diseases (AIBDs), became a matter of particular concern for healthcare systems; dissemination of sound and updated information became mandatory to orient physicians and, more generally healthcare policy [1,2]. Furthermore, AIBDs patients were encouraged take advantage from teledermatological rather than in-person consultations; also, they were encouraged to undergo COVID-19 vaccination, preferably during remission periods or at least while under low immunosuppression [3,4]. To date, data regarding the effect of Covid-vaccines on AIBDs are scanty. We present 5 cases (from 4 different centers) of bullous pemphigoid (BP) - respectively of pemphigus vulgaris (PV) -, all confirmed by clinical, immunopathologic/serological findings, where the patients had undergone COVID-vaccination during a period of remission of their bullous disease and has subsequently experienced a disease flare.

**Case 1.** A 63-years-old female with bullous pemphigoid (BP) in clinical remission since 6 months, after treatment oral prednisone was administered the first dose of Moderna mRNA-1273 vaccine and 3 days later she experienced a flare of her disease: the latter started with a generalized erythema 1 day after the vaccination and on the third day several blisters appeared on the trunk. The diagnosis was made in a telemedical consultation and oral prednisone was prescribed. 28 days later, the patient was given the second dose of the vaccine, no disease flares and no injection site reactions took place.

**Case 2.** A 40-years-old male, with pemphigus vulgaris (PV) in remission since one year after rituximab therapy, came to our attention 3 days after the first dose of the Moderna mRNA-1273 vaccine because of a PV flare. He presented with several blisters on the back and on the upper limbs (Figure 1) and started a therapy with mycophenolate mofetil and oral prednisone. After 28 days he was given the second vaccine dose: he only experienced pain on the injection site for 2 days.

**Case 3.** A 84-years-old male with BP in remission since 4 years after oral prednisone and azathioprine treatment was administered the first dose of the Moderna mRNA-1273 vaccine. After 2 weeks he started showing mild blistering lesions on the trunk, which were deemed not to require systemic immunosuppressive treatment. 28 days later the patient was given the second dose; a

worsening of the lesions which became more widespread, involving also the oral cavity, took place. A treatment with oral prednisone was started.

- **Case 4.** A 82-years-old female with BP in remission since 3 years after oral prednisone and mycophenolate mofetil treatment was given the first dose of the Pfizer mRNA BNT162b2 vaccine. 3 days later, the patient experienced a moderate BP flare in the form of small blisters on the arms and legs. The patient lived in a nursing home where only teledermatological consultations were possible. Oral prednisone was prescribed, and the second dose was administered 21 days later. No further BP flares and no injection site reactions occurred.
- **Case 5.** A 80-years-old male with a history of PV remitted 1 year before after oral prednisone and mycophenolate mofetil treatment experienced a flare 3 days after the first dose of the Pfizer mRNA BNT162b2 vaccine. He was seen in a teledermatological consultation because of severe blisters on the back and he was treated with oral prednisone. After 21 days, the patient was able to complete the Pfizer mRNA BNT162b2 vaccination with the second dose. No further BP flares and no injection site reactions occurred.

Despite the ongoing treatment with immunosuppressants, all patients developed IgG antibodies against the SARS-CoV-2 S1- Receptor Binding Domain (RBD) (>150 UI) 1 month after the second dose, which supports the assumption that the vaccination was effective in all cases presented.

These anecdotal data suggest that both mRNA vaccines and adenoviral vector vaccines may trigger relapses in AIBD patients. Vaccines are important in AIBDs patients, since the latter are characterized by a vulnerability against infections in the mucocutaneous barrier [5]: completion of vaccination is advisable, and patients should be treated for flares when needed. However, data on the effect on immunosuppressants on IgG antibodies titers in short and long term are needed. IgG S1-RBD persistence in AIBD patients is also a matter of concern; this should encourage dermatologists to monitor anti-COVID-19 immunity.

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## References

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## Figures Legend:

**Fig. 1** – Patient number 2 showing a flare of PV after the 1<sup>st</sup> dose of Moderna mRNA-1273 vaccine

## Tables Legend:

**Table 1** – Data of AIBDs affected patients

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	<b>Sex/Age</b>	<b>AIBD</b>	<b>Previous medications for AIBDs</b>	<b>Vaccine (2 doses)</b>	<b>New medications for flare up</b>	<b>Anti SARS- CoV-2 S1-RBD IgG &gt;50 UI/L</b>
<b>1</b>	F/75	BP	Oral prednisone	Moderna	Oral prednisone	Yes
<b>2</b>	M/40	PV	Rituximab	Moderna	Oral prednisone Mycophenolate mofetil	Yes
<b>3</b>	M/84	BP	Oral prednisone Azathioprine	Moderna	Oral prednisone*	Yes
<b>4</b>	F/82	BP	Oral prednisone Mycophenolate mofetil	Pfizer	Oral prednisone	Yes
<b>5</b>	M/80	PV	Oral prednisone Mycophenolate mofetil	Pfizer	Oral prednisone	Yes

\*treatment started only after the 2<sup>nd</sup> dose



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