COVID-19 vaccine-associated subacute thyroiditis: an unusual suspect for de Quervain’s thyroiditis

Mohammad Sadiq Jeeyavudeen,1 Alan W Patrick,1 Fraser W Gibb,1,2 Anna R Dover1,2

SUMMARY
Subacute thyroiditis following vaccination is an uncommon presentation of thyrotoxicosis. As the world undertakes its largest immunisation campaign to date in an attempt to protect the population from COVID-19 infections, an increasing number of rare post vaccine side effects are being observed. We report a case of a middle-aged woman who presented with painful thyroid swelling following the second dose of the COVID-19 mRNA vaccine BNT162b2 (Pfizer–BioNTech) with clinical, biochemical and imaging features consistent with destructive thyrotoxicosis. Symptomatic management only was required for the self-limiting episode. Thyroiditis typically has a mild and self-limiting course and thus this observation should not deter people from vaccination, as COVID-19 infection has a far greater morbidity and mortality risk than thyroiditis.

BACKGROUND
Subacute thyroiditis (SAT) post vaccination is an uncommon presentation of thyrotoxicosis. With the largest vaccination drive ongoing globally, rarer side effects are being noted. Here, we report an interesting case of thyrotoxicosis following COVID-19 vaccination, its clinical course, diagnostic challenge and management.

CASE PRESENTATION
A middle-aged woman presented with a history of rapid onset painful swelling of the thyroid gland and symptoms of thyrotoxicosis, including poor sleep, worsening night sweats, hyper defaecation and weight loss. The onset of symptoms was 2 weeks following her second dose of the COVID-19 mRNA vaccine BNT162b2 (Pfizer–BioNTech). There was no significant medical history and the patient was not on any regular prescribed medications. There was no history of prior use of lithium, amiodarone or interferon (which could potentially precipitate thyrotoxicosis).

INVESTIGATIONS
Initial biochemical investigations were consistent with thyrotoxicosis, with normal anti-thyroid peroxidase (TPO) and thyroid stimulating hormone (TSH) receptor antibody levels (TRAb; table 1). Technetium (Tc-99m) pertechnetate thyroid scintigraphy performed during the thyrotoxic phase demonstrated minimal isotope uptake, consistent with destructive thyroiditis (figure 1). She had no treatment other than a short course of a non-steroidal anti-inflammatory drug and her symptoms resolved over 6 weeks; her thyroid function normalised within 8 weeks (table 1).

OUTCOME AND FOLLOW-UP
Our patient was managed conservatively with a course of Non-steroidal anti-inflammatory drugs (NSAIDs) for her pain relief. As her thyrotoxicosis symptoms rapidly resolved, she did not require any antithyroid medication or beta-blocker. At her follow up review at 3 months, she continued to remain clinically and biochemically euthyroid.

DISCUSSION
SAT, also identified by various other names, is well recognised to cause follicular destruction and the rapid release of preformed thyroid hormones.1 If a likely precipitating event can be identified, this is most commonly a viral upper respiratory tract infection during the 2–3 weeks preceding the onset of thyroid symptoms.2 However, there have also been documented cases of SAT occurring shortly after influenza vaccination.3 The exact pathogenesis underlying vaccination-associated SAT is unclear. Some authors have attributed it to the adjuvant causing autoimmune/inflammatory syndrome induced by adjuvants (ASIA) but others have suggested that the vaccine core component might cause either direct injury or share the same epitope, inducing antibody cross-reaction between the antigen in the vaccine and thyroid follicular cells.4 Biopsy studies in these patients have demonstrated infiltration predominantly with multinuclear giant cell granulomas.5

SARS-CoV-2 virus gains entry into multiple endocrine tissues, including thyroid follicular cells, due to the abundance of angiotensin-converting enzyme-2 receptors expressed on their surface.6 Studies with human IgG1 monoclonal antibody against SAR-CoV-2 spike protein have shown cross-reactivity with TPO, thyroglobulin and other cellular components of the follicle.7 Some form of thyroid dysfunction is seen in nearly one-third of all COVID-19 infections in whom thyroid function has been measured.8 However, studies during the development of vaccines against the SARS-CoV-2 virus have not reported a significant incidence of thyroid complications. This may be due to patient numbers in these studies being too small to enable

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the identification of rare side effects. To our knowledge, there have been just two other similar reports of post-vaccination thyrotoxicosis so far, one following the Pfizer/BioNTech mRNA vaccine and the other following CoronaVac, which is an inactivated whole-cell vaccine. At this stage, it is very difficult to be certain whether it is the antigen component or the adjuvant which predominantly causes the immune/inflammatory response.

The diagnosis is essentially clinical, with elevated thyroid hormone and suppressed TSH concentrations. Low isotope uptake on thyroid scintigraphy and normal TPO and TSH receptor antibody levels are useful in excluding other common causes of hyperthyroidism. Where there is limited access to these investigations, a lower ratio of T3 to T4 favours destructive thyroiditis in comparison with Graves’ thyrotoxicosis. The treatment is symptomatic, with non-steroidal anti-inflammatory drugs and beta-blockers during the inflammatory thyrotoxic phase and subsequently levothyroxine if the patient goes on to develop symptomatic hypothyroidism, as with any form of SAT. Corticosteroids may be considered for the patient presenting with moderate-to-severe pain with or without thyrotoxicosis or who failed to respond to initial treatment with NSAIDs.

### Table 1 Laboratory results

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Trend in the results</th>
<th>Reference range</th>
</tr>
</thead>
<tbody>
<tr>
<td>TSH</td>
<td>Thyrotoxic phase</td>
<td>Euthyroid phase</td>
</tr>
<tr>
<td>TSH (&lt;0.010)</td>
<td>0.63</td>
<td>0.2–4.5 (mU/L)</td>
</tr>
<tr>
<td>Free T4</td>
<td>27</td>
<td>10</td>
</tr>
<tr>
<td>Total T3</td>
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<td>–</td>
</tr>
<tr>
<td>Anti-TPO</td>
<td>79.5</td>
<td>–</td>
</tr>
<tr>
<td>TRAb (&lt;1.2)</td>
<td>–</td>
<td>0–2.1 (IU/L)</td>
</tr>
<tr>
<td>Haemoglobin</td>
<td>124</td>
<td>130</td>
</tr>
<tr>
<td>White cell count</td>
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<td>5.1</td>
</tr>
<tr>
<td>Lymphocyte count</td>
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<td>1.38</td>
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<tr>
<td>Platelet count</td>
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<td>267</td>
</tr>
<tr>
<td>C reactive protein</td>
<td>23</td>
<td>&lt;1</td>
</tr>
</tbody>
</table>

TPO, thyroid peroxidase; TRAb, TSH receptor antibody; TSH, thyroid stimulating hormone.

### Learning points

- Subacute thyroiditis following COVID-19 vaccination is a rare complication.
- Taking a comprehensive history, including recent COVID-19 vaccination and/or infection, can help to identify rare vaccine-related complications.
- Thyroiditis is usually a mild self-limiting illness and the possible association with prior vaccination should not deter people from COVID-19 vaccination.

Vaccination on such a global scale will inevitably be associated with rare side effects, which were not evident in vaccine trials. However, SAT is generally relatively mild and transient, so it should not deter people from getting vaccinated, particularly given the much greater risks from contracting COVID-19 infection.

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### Contributors

MSJ and ARD were involved in the primary evaluation and management of the patient and conception and drafting of the case report. FG and AWP were involved in reviewing the existing literature and revising the report. ARD is the guarantor.

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**Figure 1** Technetium (Tc-99m) pertechnetate uptake scan showing only minimal uptake (indicating by red arrow) suggestive of destructive thyroiditis.
Case report

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REFERENCES