Subacute thyroiditis after inactive SARS-CoV-2 vaccine

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SUMMARY
Viral infections have often been associated with subacute (De Quervain) thyroiditis. Rare cases of subacute thyroiditis have been reported after vaccines. Various vaccines have been developed with different techniques against SARS-CoV-2. This case report presents a rare case of subacute thyroiditis after the inactive SARS-CoV-2 vaccine, CoronaVac.

BACKGROUND
SARS-CoV-2 is a novel coronavirus that caused a global pandemic. Although its destructive results continue to increase, vaccination studies have created hope for treatment. Various vaccines have been developed for SARS-CoV-2. CoronaVac, an inactivated virus vaccine, is one of them. Two doses of CoronaVac were first administered to healthcare workers within Turkey’s framework of the vaccination programme. Subacute thyroiditis (SAT) is an inflammatory disease that occurs secondarily to viral or postviral responses and causes damage to the thyroid gland. Cases of SAT after various vaccines have been reported.

CASE PRESENTATION
A 38-year-old female physician was admitted to the endocrinology outpatient clinic with reports of swelling in the neck, pain, fatigue, loss of appetite and sweating in the evening 2 weeks after being administered with the second dose of CoronaVac vaccine. The CoronaVac vaccine was administered on the first day and the 28th day. The patient stated that a slight pain and tenderness occurred in the thyroid lodge after the first dose, but it passed within a few days, and she did not consult a doctor during this period. The patient had no prior illness, previous vaccination or drug use. Physical examination revealed stage 2 goitre, and there was pain in the right thyroid lobe when it was touched. In laboratory tests, thyrotropin (TSH): 0.008 uIU/mL (normal: 0.27–4.2), free T3: 12.88 pg/mL (normal: 2–4), free T4: 4.65 ng/dL (normal: 0.93–1.6), anti-Tpo: 9.49 IU/mL (normal: 0–34), anti-Tg: 81.58 IU/mL (normal: 0–115), C reactive protein: 8.76 mg/L (normal: 0–0.8), sedimentation: 78 mm/hour (normal: 0–20) were detected. CO2: 8.76 mmHg (normal: 26–43), CO: 48.76 mmHg (normal: 14–42) and PO2: 81.58% (normal: 95–100) were normal. Electrocardiography was performed and was normal. Thyroid ultrasonography revealed a partially healed nodule on the right thyroid lobe (white arrow in figure 1A). SAT diagnosis was established with a high acute phase, thyrotoxicosis and USG findings. Naproxen sodium 2×275 mg and propranolol 2×20 mg peroral treatment was initiated. Following treatment, the neck pain was alleviated. The patient stated that on the 14th day of the follow-up, her problems had mostly disappeared. The baseline and follow-up values of the patient are given in table 1.

OUTCOME AND FOLLOW-UP
On the 30th day, levothyroxine started to be administered at 25 μg/day due to a high TSH level (24.68 uIU/mL) and decreased free T4. As TSH was 20 uIU/mL on the 45th day, the levothyroxine dose was increased to 50 μg/day. On the 45th day, USG revealed partial recovery of the thyroid gland (figure 1B). Moreover, the patient had no reports other than the biochemical findings of hypothyroidism.

DISCUSSION
SAT is a self-limiting inflammatory thyroid disease characterised by neck pain, fever and thyroid dysfunction. There is usually a history of upper respiratory tract infection before SAT. Viral agents can cause SAT with an indirect immunological reaction, and there are some reports that viruses can cause SAT by direct tissue invasion. Many viruses have been reported as potential causative agents. However, an evident infectious agent can rarely be demonstrated in most patients. SAT cases have been reported following the SARS-CoV-2 infection. The general features of these are similar to SAT cases reported in other aetiologies. Most of them are seen in young women; pain in the thyroid region is usually the first presenting complaint, and an increase in inflammatory markers and thyrotoxicosis have been reported in other cases. In SAT cases, clinical and USG findings can provide a specific diagnosis, and a radioactive iodine uptake imaging revealed stage 2 goitre, and there was pain in the right thyroid lobe when it was touched. In laboratory tests, thyrotropin (TSH): 0.008 uIU/mL (normal: 0.27–4.2), free T3: 12.88 pg/mL (normal: 2–4), free T4: 4.65 ng/dL (normal: 0.93–1.6), anti-Tpo: 9.49 IU/mL (normal: 0–34), anti-Tg: 81.58 IU/mL (normal: 0–115), C reactive protein: 8.76 mg/L (normal: 0–0.8), sedimentation: 78 mm/hour (normal: 0–20) were detected. COVID-19 PCR testing was not performed as there were no COVID-related symptoms except SAT findings. Thyroid ultrasonography (USG) revealed an increased size of the right thyroid lobe, an irregularly demarcated hypoechoic area of approximately 3 cm in diameter compatible with thyroiditis starting from the capsule in the lateral and progressing into the lobe (white arrow in figure 1A). SAT diagnosis was established with a high acute phase, thyrotoxicosis and USG findings. Naproxen sodium 2×275 mg and propranolol 2×20 mg peroral treatment was initiated. Following treatment, the neck pain was alleviated. The patient stated that on the 14th day of the follow-up, her problems had mostly disappeared. The baseline and follow-up values of the patient are given in table 1.

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study is often not required, as was the case in this instance. The patient was a young woman diagnosed with typical symptoms and SAT USG findings.

Transmembrane protease serine-2 (TMPRSS-2) and ACE-2 receptors allow the SARS-CoV-2 to enter human cells. 7,8 ACE-2 and TMPRSS-2 are expressed in thyroid follicular cells, and their expression is higher than in lung cells, especially in women. 9 These situations may explain why SARS-CoV-2 causes SAT in women more frequently.

Inactive virus vaccines contain many proteins belonging to the pathogen virus and similar antigenic parts. 10 The increased affinity of the SARS-CoV-2 towards the thyroid suggests that this inactive vaccine may also affect the thyroid tissue. Few cases of SAT have been reported after various vaccinations in the literature. It is noteworthy that these cases are mostly seen after an inactivated virus vaccination. There are cases reported with SAT after an inactive virus vaccine: seasonal influenza vaccine, 2,11,12 inactivated virus vaccination. There are cases reported with SAT after an inactive virus vaccine: seasonal influenza vaccine, 2,11,12 H1N1 vaccine 13 and hepatitis B vaccine. 14

Besides viral proteins, adjuvant substances used in the vaccine to increase immunological reactions may also trigger autoimmune reactions. 15 The CoronaVac vaccine contains aluminium hydroxide as an adjuvant. A condition called the autoimmune/immune reactions. 15 The CoronaVac vaccine contains aluminium hydroxide as an adjuvant. A condition called the autoimmune/inflammatory syndrome induced by adjuvants (ASIA syndrome) was previously reported following various vaccines. 16 However, the CoronaVac vaccine has been related to causing the ASIA syndrome and SAT in a recent article. 17

In our case, the milder thyroiditis findings after the first dose of vaccine and typical SAT findings after the second dose are noteworthy. The aetiology of SAT remains unclear in many cases, but given the typical USG image and the temporal relationship between vaccination and the onset of thyrotoxicosis, as in this case, CoronaVac can probably be associated with SAT. Therefore, clinicians should be aware that thyroid symptoms can potentially be associated with the CoronaVac vaccine; however, such a side effect should never preclude vaccination.

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REFERENCES