CASE REPORT

Hypoparathyroidism: an uncommon adverse effect of treatment with durvalumab

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Summary

Immune checkpoint inhibitors (ICIs) are monoclonal antibodies approved for the treatment of numerous cancer types. Toxicities induced by ICIs may affect any organ system and manifest as endocrinopathy. The main side effects related to treatment are immune-related adverse events (irAEs), especially thyroid dysfunction and hypophysitis. Rare endocrine irAEs are diabetes insipidus, hypoparathyroidism, thyrotoxic crisis and hypogonadism. We report a case of hypoparathyroidism induced by ICI treatment with durvalumab, which has not previously been described.

Key words

- immune checkpoint inhibitors
- durvalumab
- immune-related adverse effects
- hypoparathyroidism

Learning points:

- Treatment with immune checkpoint inhibitors (ICIs) is associated with many endocrine side effects.
- It is recommended that patients treated with ICIs are observed by an endocrinologist.
- If side effects are treated accordingly, ICI therapy can continue.

Background

Immune checkpoints are small molecules (cytotoxic T lymphocyte antigen (CTLA-4), programmed cell death protein 1 (PD-1), programmed cell death ligand-1 (PD-L1)) expressed by immune cells involved in immune homeostasis. The targeted monoclonal antibodies directed against these regulatory checkpoint molecules (immune checkpoint inhibitors (ICIs)) are used to treat some types of cancer. PD-1 is an immune inhibitory receptor expressed on activated T cells, B cells, macrophages and natural killer cells. It has two binding ligands PDL-1 and PDL-2 expressed on normal cells. Binding of PD-1 to its ligand PD-L1 triggers an inhibitory signal, leading to reduced T-cell proliferation, inhibited T-cell activity and immune response and antitumor immunity. Physiologically, the PD-1/PD-L1 pathway emerges as a result of the need to control the degree of inflammation at locations expressing the antigen, in order to secure normal tissue from damage, prevent immune cell activation and kill normal cells. However, some malignant tumors take advantage of this mechanism. They overexpress a large number of PDL-1 on the surface to reduce T-cell activation and antigen-specific T-cell immune response and thereby bypass immune surveillance.

The mechanism of action of PD-1 and PD-L1 inhibitors is in their ability to block PD-L1 binding to PD-1, allowing T cells to then be able to kill tumor cells. ICI therapy has shown antitumor efficacy, and today it is a standard treatment for many tumor types (Deligiannis et al. 2021). The ICIs that are approved by the US Food and Drug Administration include anti-CTLA-4
Hypoparathyroidism is a rare complication induced by ICIs and encountered in treatment with nivolumab (Cubb et al. 2017), pembrolizumab (Piranavan et al. 2020), and pembrolizumab (Dadu et al. 2018). Hypoparathyroidism as a complication of inappropriate antidiuretic hormone secretion (SIADH) and hypocalcemia is associated with low-normal or low PTH levels. Antiparathyroid and calcium-sensing receptor-autoantibodies may be detectable in patients with ICI-related hypoparathyroidism mediated by these autoantibodies (George et al. 2021). Acute symptomatic hypoparathyroidism develops between 3 weeks and 7 months of treatment with ICIs (Umeguchi et al. 2018, El Kawgki et al. 2020). Hypoparathyroidism as a complication in checkpoint inhibitor therapy is largely irreversible, and long-term treatment with an activated form of vitamin D analog and calcium supplements is required.

Hypoparathyroidism is a rare complication induced by ICIs and encountered in treatment with nivolumab (Edd et al. 2018, Piranavan et al. 2019), ipilimumab with nivolumab (Cubbe et al. 2017, Win et al. 2017, Trinh et al. 2019, Dadu et al. 2020, El Kawgki et al. 2020) and pembrolizumab (Umeguchi et al. 2018, Lupi et al. 2020). In the three

<table>
<thead>
<tr>
<th>Test</th>
<th>Patient level</th>
<th>Normal range</th>
</tr>
</thead>
<tbody>
<tr>
<td>Calcium</td>
<td>1.41 mmol/L</td>
<td>2.1–2.55 mmol/L</td>
</tr>
<tr>
<td>Ionized calcium</td>
<td>0.65 mmol/L</td>
<td>1.13–1.32 mmol/L</td>
</tr>
<tr>
<td>Phosphate</td>
<td>2.11 mmol/L</td>
<td>0.74–1.52 mmol/L</td>
</tr>
<tr>
<td>Albumin</td>
<td>43 g/L</td>
<td>32.0–46.0 g/L</td>
</tr>
<tr>
<td>Magnesium</td>
<td>0.69 mmol/L</td>
<td>0.66–1.07 mmol/L</td>
</tr>
<tr>
<td>25-hydroxyvitamin D</td>
<td>38.6 nmol/L</td>
<td>50.0–125.0 nmol/L</td>
</tr>
<tr>
<td>PTH</td>
<td>&lt;0.58 pmol/L</td>
<td>1.58–6.03 pmol/L</td>
</tr>
</tbody>
</table>

PTH, parathyroid hormone.

Discussion

ICIs are approved for the treatment of some types of advanced cancer. Toxicities induced by ICIs are autoimmune and referred as immune-related adverse events (irAEs). The incidence of endocrinopathies during ICI therapy reaches 10% in meta-analysis of 38 studies, which involved a total of 7551 patients under ICIs (Deligiannis et al. 2021). Among the patients on anti-CTLA-4 agent monotherapy, the most frequent endocrine irAE is hypophysitis (5.6%), whereas among patients on anti-PD-1/PD-L1 agent monotherapy, the most frequent irAE is hypothyroidism (8.5%). Rare endocrinopathies during ICI therapy are diabetes mellitus, hypogonadotropic hypogonadism in the absence of hypophysitis, diabetes insipidus, syndrome of inappropriate antidiuretic hormone secretion (SIADH) and transient adrenocorticotropic hormone (ACTH)-dependent hypercortisolism (George et al. 2021).

Hypoparathyroidism is an extremely rare irAE. Hypoparathyroidism should be considered when hypocalcemia is associated with low-normal or low PTH levels. Antiparathyroid and calcium-sensing receptor-activating autoantibodies may be detectable in patients with ICI-related hypoparathyroidism mediated by these autoantibodies (George et al. 2021). Acute symptomatic hypoparathyroidism develops between 3 weeks and 7 months of treatment with ICIs (Umeguchi et al. 2018, El Kawgki et al. 2020). Hypoparathyroidism as a complication in checkpoint inhibitor therapy is largely irreversible, and long-term treatment with an activated form of vitamin D analog and calcium supplements is required.

Hypoparathyroidism is a rare complication induced by ICIs and encountered in treatment with nivolumab (Edd et al. 2018, Piranavan et al. 2019), ipilimumab with nivolumab (Cubbe et al. 2017, Win et al. 2017, Trinh et al. 2019, Dadu et al. 2020, El Kawgki et al. 2020) and pembrolizumab (Umeguchi et al. 2018, Lupi et al. 2020). In the three
reported cases, other concomitant endocrinopathies were confirmed: autoimmune thyroiditis with a thyrotoxic phase (Win et al. 2017, Dadu et al. 2020) and hypophysitis with adrenal insufficiency (El Kawgki et al. 2020). Despite similarities in clinical manifestation, emerging data suggest immunological differences between irAEs and ‘traditional’ autoimmune diseases, as conventional autoantibodies are often not detectable when irAEs are diagnosed (Khan & Monzon et al. 2020). Mechanisms which generate irAEs have been suggested: preexisting susceptibility to autoimmunity, aberrant presentation of ‘self’ by the tumor, loss of tolerance driven by the tissue or tumor microenvironment (Burke et al. 2020).

All reported cases of ICI-induced hypoparathyroidism in the Embase and Medline databases are listed in Table 2.

Because of the lack of associated causes of hypoparathyroidism (anterior neck surgery, radiotherapy of the neck, absence of other autoimmune syndromes, negative antiparathyroid antibodies, hypomagnesemia, severe vitamin D deficiency and hypoalbuminemia), we believe that the patient developed hypoparathyroidism as a very rare complication associated with durvalumab therapy.

### Conclusion

ICIs are effective therapeutic modalities for many cancers, but their use also produces many adverse side effects in the endocrine system. Supervision by an endocrinologist is therefore recommended for optimal care. Patients on ICIs should be routinely monitored for the development of endocrinopathies before immunotherapy: fasting venous glycemia (if anti-PD1/PD-L1), natremia, thyroid-stimulating hormone (TSH), fT4, 08:00 h cortisol (without corticosteroid intake), +/-ACTH (depending on 08:00 h cortisol level), luteinizing hormone (LH), follicle-stimulating hormone (FSH), testosterone in males, LH, FSH, estradiol in females and FSH in menopausal females. During immunotherapy, patients should be monitored in each course of treatment for 6 months and every two courses for the following 6 months. Patients should also be monitored in cases of clinical alert signs: fasting venous glycemia (if anti-PD1/PD-L1), natremia, TSH, fT4, 08:00 h cortisol and testosterone in males (Castinetti et al. 2019). In contrast to other irAEs, patients even with high grades of endocrine irAEs may continue their ICI therapy, provided the hormone replacement therapy is adequate and the symptoms are controlled (George et al. 2021).

### Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of this case report.

### Funding

This work did not receive any specific grant from any funding agency in the public, commercial or not-for-profit sector.

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**Table 2**  All reported cases of ICI-induced hypoparathyroidism.

<table>
<thead>
<tr>
<th>Authors</th>
<th>Drug diagnosis</th>
<th>Onset hypoparathyroidism after treatment ICI</th>
<th>Duration hypoparathyroidism</th>
<th>Antibodies</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cubb et al.</td>
<td>I + N Melanoma</td>
<td>4 weeks</td>
<td>Parathyroid function has not recovered since diagnosis</td>
<td>Negative anti-PTH antibodies</td>
</tr>
<tr>
<td>Dadu et al.</td>
<td>I + N Melanoma</td>
<td>4 weeks</td>
<td>3 years and 3 months</td>
<td>Negative CaSR antibodies</td>
</tr>
<tr>
<td>Edd et al.</td>
<td>N NSCLC</td>
<td>5 months</td>
<td>Passing hospice shortly after moving</td>
<td>Not measured</td>
</tr>
<tr>
<td>El Kawgki et al.</td>
<td>I + N Melanoma</td>
<td>7 months</td>
<td>77 days</td>
<td>Negative anti-PD-L1 antibodies</td>
</tr>
<tr>
<td>Lupi et al.</td>
<td>P Lung adenocarcinoma N SCLC</td>
<td>15 weeks</td>
<td>9 months</td>
<td>Positive CaSR antibodies</td>
</tr>
<tr>
<td>Piranavan et al.</td>
<td>P Lung adenocarcinoma N SCLC</td>
<td>5 months</td>
<td>1 year</td>
<td>Positive CaSR antibodies</td>
</tr>
<tr>
<td>Trinh et al.</td>
<td>I + N Melanoma</td>
<td>4 weeks</td>
<td>Improvement after 8 months</td>
<td>Detectable CaSR antibodies</td>
</tr>
<tr>
<td>Umeguchi et al.</td>
<td>P NSCLC</td>
<td>3 weeks</td>
<td>5 months</td>
<td>Not measured</td>
</tr>
<tr>
<td>Win et al.</td>
<td>I + N Melanoma</td>
<td>1,5 months</td>
<td>4 months</td>
<td>Not measured</td>
</tr>
</tbody>
</table>

CaSR, calcium-sensing receptor; I, ipilimumab; ICI, immune checkpoint inhibitor; N, nivolumab; NSCLC, non-small cell lung cancer; P, pembrolizumab; PTH, parathyroid; SCLC, small cell lung cancer.
Patient consent
Written informed consent for publication of their clinical details was obtained from the patient.

Author contribution statement
A K conceived the study, A K and M H did the literature search, analyzed data and wrote the manuscript, T B was in-patient treating doctor, K K contributed to biochemistry, immunology and hormonal analysis and interpretations. All contributors approved the final accepted version of the manuscript.

References


Cubb T, Patel S, et al. 2017 Primary hypoparathyroidism: a new endocrine immune related adverse event (irAEs) secondary to combination treatment with PD-1 and CLA-4 checkpoint inhibitors. Endocrine Reviews 38 (Supplement 1). (https://doi.org/10.1093/edrv/38.supp.1)


Deligiannis NG, Sosa S, Danilowicz K & Rizzo LFL 2021 Endocrine dysfunction induced by immune checkpoint inhibitors. Medicine 81 269–278.


Khan OF & Monzon J 2020 Diagnosis, monitoring, and management of adverse events from immune checkpoint inhibitor therapy. Current Oncology 27 543–550. (https://doi.org/10.3747/co.27.5111)


