Ipilimumab-induced autoimmune hypophysitis: a differential for sellar mass lesions

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Summary

Autoimmune hypophysitis (AH) has been previously described in a typical demographic population, primarily women in the reproductive age group and perinatal period. The era of immune modulation using anti-cytotoxic T-lymphocyte-associated antigen 4 biological therapy (ipilimumab) against advanced cancers like metastatic melanomas has now resulted in a new form of hypophysitis being increasingly recognised under a spectrum of immune-related adverse events. Drug-related AH often presents with subtle symptoms and a pituitary mass, with the potential for fatality necessitating wide awareness and a high index of clinical suspicion given that it is usually treatable. We describe below two cases of AH within the last three months at our centre, which were treated with different regimens and produced good endocrine outcomes.

Learning points:

- AH is a new and defined clinical entity occurring as a side effect of ipilimumab, which enhances immune-mediated destruction of metastatic melanoma.
- It can present insidiously and have life-threatening complications related to hypocortisolism, hence a high index of clinical suspicion must be exerted by treating physicians, and seems to result in resolution of pituitary masses and variable improvements of pituitary function.
- Clinical improvement, radiological resolution of pituitary masses and variable normalisation of pituitary function are possible with early treatment with high-dose oral or i.v. steroids and hormone replacement therapy, although duration and dosing protocols are unclear at this stage.
- Ipilimumab should continue to be prescribed as treatment for metastatic melanoma; however, close clinical observation of patient’s progress must be maintained while they are on this drug.
- Predictive factors for onset of AH remain unclear and it is imperative that AH is distinguished from pituitary metastases.
- Further studies are required to determine the safety of continuing therapy with ipilimumab in patients who have developed AH while on treatment.

Background

This case series accentuates the importance of a high index of clinical suspicion among cancer patients on ipilimumab therapy. Although not uncommon, autoimmune hypophysitis (AH) is a relatively new clinical entity which, if left untreated, can have potentially life-threatening complications. Treatment of this condition is available and successful, establishing resolution of clinical symptomology, highlighting the importance of early detection. This case series describes accurate diagnosis
and treatment options for patients who present with ipilimumab-induced hypophysitis.

Case presentation

Case 1

A 64-year-old male presented with a large destructive lesion of the T9 vertebra, due to metastatic melanoma. He was treated with palliative radiotherapy and commenced on 3 mg/kg dose regimen of ipilimumab, usually administered 3–4 weeks apart in up to four doses. However, following the third dose, he reported extreme fatigue, somnolence, emotional lability, retro-orbital headaches, constipation, anorexia and nausea. Biochemical findings confirmed central hypocortisolism and hypothyroidism (Table 1). Magnetic resonance imaging (MRI) of the pituitary revealed a 9 mm T1 hypointense contrast enhancing anterior pituitary mass with marginally increased stalk thickness of 3.3 mm (Fig. 1A). He did not have evidence of diabetes insipidus.

Given the acute presentation and imaging features suggesting a high likelihood of AH, he was commenced on 75 mg oral prednisolone once daily for 2 weeks, and thyroxine (T4) replacement at 75 mg daily. He made remarkable symptomatic improvement and the steroids were tapered to 50 mg prednisone for 2 weeks, and then at weekly intervals to 25, 10 and 7.5 mg. Eight weeks later, he was clinically well with normal pituitary function, with significant improvement on a progress MRI with reduction of the pituitary gland to 5.5 mm, and normal pituitary stalk thickness at 2.1 mm (Fig. 1B). He is now of T4 and low-dose maintenance hydrocortisone in divided doses (14 mg/day) with a plan to taper further as tolerated. His melanoma disease burden remains stable thus far.

Case 2

A 65-year-old woman presented with metastatic melanoma in the liver and peritoneum for which she was commenced on 3 mg/kg of ipilimumab. She initially tolerated the treatment well, but further developed increasing fatigue, weight gain of 6 kg and bloating, along with elevation in blood glucose levels to 8–13 mmol/l (144–234 mg/dl) following her third dose. Her blood tests revealed secondary hypothyroidism, normal prolactin levels and central hypogonadism (Table 1). MRI detected a new hypointense heterogeneous mass of the pituitary gland, measuring 12 mm craniocaudally, and 13 mm anteroposteriorly, causing slight distortion of the optic chiasm suggesting ipilimumab-induced hypophysitis (Fig. 2A). She was commenced on 3 days of i.v. methylprednisolone at 80 mg once daily as an in-patient, followed by a tapering regimen of high-dose oral prednisolone for 4 weeks starting at 80 mg daily, in addition to 50 μg of daily T4.

Four weeks later, she demonstrated marked clinical improvement, with normal thyroid hormone levels (Table 1) and resolution of the pituitary mass on MRI, which now showed a homogeneous enhancement measuring 1×0.3×0.3 cm with a central stalk completely clear of the optic chiasm (Fig. 2B). Her reproductive hormones were not assayed due to her post-menopausal state. Oral prednisolone was tapered to 10 mg for 2 weeks, followed by 5 mg until further review. Unfortunately, the disease did not respond to ipilimumab and progressed rapidly. She was transferred to the palliative care unit where she ultimately succumbed to her disease. She, however, needed maintenance T4 and prednisone over the period.


downloadable in PDF format Table 1 Table outlining hormone levels in the patients’ pre- and post-high-dose steroid therapy. Values and normal references in SI units and parentheses indicate conventional units.

|                  | Case 1 | Case 2 |
|------------------|--------|--------|
|                  | Pre    | Post   | Pre    | Post   |
| ACTH – 10–50 ng/l (10–50 pg/ml) | 10 (10) | 21 (21) | 73 (73) | 10 (10) |
| Cortisol – 50–650 nmol/l (5–25 μg/dl) | 62 (2.25) | 269 (9.75) | 669 (24.25) | 813 (29.5) |
| TSH – 0.5–5 mIU/l | 0.08 | 0.6 | 0.04 | 0.8 |
| FT3 – 2.5–6.0 pmol/l (162–390 pg/dl) | 2.8 (156) | 4.8 (312) | 2.3 (149) | 2.9 (188) |
| FT4 – 10–20 pmol/l (0.9–2.3 ng/dl) | 6.1 (0.47) | 11.0 (0.85) | 10.2 (0.79) | 12.9 (1.0) |
| Prolactin <500 mIU/l (4–23 μg/l) | 174 (4.0) | 266 (6.12) | 167 (3.84) | 338 (7.77) |
| LH | 1.8 | 2.2 | 2 | 1 |
| FSH | 6.6 | 1.5 | 6 | 6 |
| Testosterone | 11 | 15 | 61 | <37 |

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Treatment

Treatment provided has been described previously within case description.

Outcome and follow-up

Case 1 was commenced on high-dose oral prednisolone daily for 2 weeks, and ongoing T4 replacement, on which he improved clinically, allowing for a steroid tapering regimen to be commenced. He clinically resolved over-time with normal pituitary function and resolution of the pituitary mass. Currently, he continues on low-dose oral hydrocortisone at maintenance dosing and is stable from an oncological perspective.

Alternatively, our second patient was commenced on three days of i.v. methylprednisolone daily, followed by a tapering regimen of high-dose oral prednisolone for four weeks. She was also placed on T4 replacement therapy. Four weeks later, she improved clinically and radiological imaging demonstrated resolution of the pituitary tumour, hence an attempt to wean her steroid regimen was made. Her melanoma disease burden, however, progressed during this period and she ultimately lost her cancer battle within the following month.

It is important to note that our patients demonstrated symptomatic improvement at scheduled follow-up appointments at different times (i.e. 2 weeks vs 4 weeks) post-commencement of treatment, indicating that response to hormone replacement therapy may vary among patients.

Discussion

Ipilimumab is a MAB to cytotoxic T-lymphocyte-associated antigen 4 that inhibits tumour progression by enhancing immune-mediated destruction of malignant cells (1) (2) (3). While this novel class of drugs hold promise in the treatment of various cancers, they also have the potential to unleash an autoimmune flare in susceptible patients, involving the gastrointestinal, endocrine and neurological systems (4). Recent studies have drawn attention to a group of endocrine immune-related adverse events, including AH, thyroid disease, adrenal insufficiency and SIADH (2).

Observational studies suggest an association among dose and duration of therapy, clinical response and severity of the side effect profile. Multiple trials have used varying doses of ipilimumab ranging from 0.3–10 mg/kg, with optimal clinical response shown to occur with 3 mg/kg (5). While affecting <5% of all ipilimumab-treated patients, AH can cause life-threatening complications, particularly hypocortisolism, and tends to clinically manifest between administration of the third and fourth dose of infusions (6).

Fortunately, treatment involving high-dose steroids along with hormone replacement has shown promising results in terms of clinical, biochemical and radiographic resolution of the condition. Current studies report patient response to i.v. administration of high-dose steroids (5); however, our first case was treated successfully as an outpatient with oral prednisone. Further studies are needed to compare response to i.v. steroids vs oral steroids. Furthermore, MRI findings among AH patients have known to be quite varied, but typically involves diffuse enlargement of the pituitary gland, more often with hypointensity on T1-weighted imaging, heterogeneous enhancement and accompanied by thickening of the pituitary stalk (6) (7). Both cases demonstrated pituitary masses with hypointensity and heterogeneous changes, which responded very well to high-dose steroids indicating the inflammatory pathology.

A pituitary biopsy remains the gold standard in establishing the diagnosis and distinguishing hypophysitis from pituitary metastases. However, due to technical limitations...
difficulties involved, this is not usually feasible. In such circumstances, clinical markers such as the acute presentation with hypocortisolism or hypopituitarism, normal prolactin levels, the hypointense lesion on imaging and absence of diabetes insipidus point towards hypophysitis rather than pituitary metastases from the melanoma. The therapeutic response to steroids confirms the diagnosis. Corsello et al. suggest that history of prior radiotherapy may have a protective role in preventing AH among patients treated with ipilimumab (8). This was not the case in Case 1, who was alone in receiving prior oncology treatment upon being commenced on ipilimumab. At this stage, predictive factors to identify those at risk of AH prior to therapy remain unclear, with neither of our patients having a known history of other autoimmune conditions, infections or allergies that could have potentially predisposed them to this immune-mediated reaction. Efficacy of ipilimumab recommencement post-treatment for AH is yet to be determined, with some studies predicting the same clinical response of AH to treatment in patients recommenced on the drug after hormone replacement therapy.

Most described cases seem to require ongoing steroid hormone replacement (8). In our cases, the first patient has been off T4 and on a maintenance dose of hydrocortisone, whereas, with our second case, we were unable to trial a taper of therapy due to deterioration in her clinical condition from the metastatic melanoma.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Patient consent
This is to confirm that patient consent was obtained from our study participant. Consent was not obtained from the second as she is deceased.

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