Histologically confirmed isolated IgG4-related hypophysitis: two case reports in young women

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Summary

IgG4-related hypophysitis is a recently described entity belonging to the group of IgG4-related diseases. Many other organs can also be affected, and it is more common in older men. To date, 32 cases of IgG4-related hypophysitis have been reported in the literature, 11 of which included confirmatory tissue biopsy and the majority affecting multiple organs. The aim of this report is to present two cases of biopsy-proven IgG4-related hypophysitis occurring in two young female patients with no evidence of involvement of other organs at the time of diagnosis.

Learning points:

- IgG4-related hypophysitis belongs to the group of IgG4-related diseases, and is a fibro-inflammatory condition characterized by dense lymphoplasmacytic infiltrates rich in IgG4-positive plasma cells and storiform fibrosis.
- It is more common in older men, but young women may also present this type of hypophysitis.
- Although involvement of other organs is frequent, isolated pituitary disease is possible.
- Frequent clinical manifestations include anterior hypopituitarism and/or diabetes insipidus.
- The diagnosis may be confirmed with any of the following criteria: a pituitary biopsy with lymphoplasmacytic infiltrates, with more than ten IgG4-positive cells; a sellar mass and/or thickened pituitary stalk and a biopsy-proven involvement of another organ; a sellar mass and/or thickened pituitary stalk and IgG4 serum levels > 140 mg/dl and sellar mass reduction and symptom improvement after corticosteroid treatment.
- Glucocorticoids are recommended as first-line therapy.

Background

Hypophysitis is a rare entity characterised by chronic inflammation of the pituitary, and is classified as primary and secondary based on its aetiology. Secondary hypophysitis includes cases resulting from other sellar lesions (Rathke cleft cyst, craniopharyngioma, germinoma, and pituitary adenomas); from therapeutic use of immunomodulatory drugs (CTLA4 blocking antibody, interferon α) or as a consequence of systemic diseases such as Wegener’s granulomatosis, tuberculosis, sarcoidosis or syphilis (1).

The primary forms comprise five histopathological subtypes: lymphocytic or autoimmune, granulomatous, xanthomatous, necrotising, and IgG4-positive plasma cell-rich hypophysitis (1). Lymphocytic hypophysitis is the most common subtype, showing extensive pituitary infiltration by lymphocytes, predominantly affecting women between 30 and 40 years of age, often occurring
during pregnancy or in the postpartum period, and frequently associated with other autoimmune disorders (2). IgG4-related hypophysitis is a recently described entity belonging to the group of IgG4-related diseases first described in 2001 by Hamano et al. (3) in a patient with sclerosing pancreatitis. We now know that many other organs can also be affected and that it is more common in older men (4).

To date, 32 cases of IgG4-related hypophysitis have been reported in the literature, 11 of which included confirmatory tissue biopsy (5) and the majority affecting multiple organs. The aim of this report is to present two new cases of biopsy-proven IgG4-related hypophysitis occurring in two young female patients, with no evidence of involvement of other organs at the time of diagnosis.

Case reports

Case 1

Case presentation A 25-year-old woman presented to our clinic with a history of secondary amenorrhoea for the past eight months. Her relevant past history revealed chronic anaemia due to thalassaemia minor, menarche at age 13 years and no prior pregnancies. She was found to present normal vital signs and her physical examination was unremarkable. She was not taking any prescribed medication.

Investigation Initial laboratory tests revealed hypo-gonadotrophic hypogonadism without hyperprolactinaemia and possible but not proven secondary adrenal failure (Table 1). Brain magnetic resonance imaging (MRI) revealed a sellar tumour with suprasellar extension (Fig. 1A and B). Computerised visual field (CVF) assessment was normal.

Treatment Based on lesion size and image characteristics, the presumptive diagnosis was a macroadenoma,
so glucocorticoid replacement therapy was started and the tumour later resected through a transsphenoidal approach with no complications. After surgery, amenorrhoea persisted and secondary hypothyroidism was detected. Oral contraceptives and thyroxine (50 μg daily) were prescribed.

**Outcome and follow-up** Three months later a new MRI of the pituitary gland showed a persistent suprasellar tumour (Fig. 1C and D) in spite of lack of symptoms. Histopathology revealed adenohypophyseal tissue with dense inflammatory lymphoplasmacytic infiltrates (Fig. 2A). IgG4-positive immunostaining was observed in more than ten plasma cells per high power field (40×), confirming IgG4-related hypophysitis diagnosis (Fig. 2B). The new MRI findings and histologically confirmed diagnosis prompted a change in medication from hydrocortisone 15 mg daily to high-dose meprednisone; however, the patient did not accept this treatment although she finally agreed to take meprednisone 8 mg daily. MRI of the chest, abdomen, and pelvis did not reveal any pathology.

An MRI repeated after four months showed significant reduction in the pituitary mass lesion, demonstrating a good response to the low-dose steroids (Fig. 1E and F). After 20 months of follow-up there was no evidence of relapse of the pituitary mass or other organ involvement (not shown). She continued to persist with oral contraceptives and levothyroxine requirements.

**Case 2**

**Case presentation** A 37-year-old woman came for consultation because of severe headache, secondary amenorrhoea, polydipsia, and polyuria (urine output 10 l/day), weight loss and fatigue. Her past medical history noted a menarche at 12 years of age, no pregnancies and gluten-sensitive enteropathy that had improved shortly after starting a gluten-free diet. She was normotensive, and she had no abnormality detected on physical examination.

**Investigation** The brain MRI showed a pituitary lesion extending into the suprasellar region with signs of intratumoural bleeding and compression of the optic chiasm (Fig. 3A and B). Baseline laboratory tests indicated panhypopituitarism and mild hyperprolactinaemia, interpreted as disconnection hyperprolactinaemia (Table 2). CVF showed left superior temporal quadrantrinopia.

**Treatment** Replacement treatment was started with hydrocortisone 15 mg/day, oral desmopressin 0.05 mg every 8 h, as well as the oral contraceptive. The patient underwent transsphenoidal surgery (TSS) with the presumptive diagnosis of pituitary macroadenoma with intratumoural haemorrhage.

Histopathology results reported IgG4-related hypophysitis (Fig. 4A and B) and treatment was changed to meprednisone 16 mg daily. Computed tomography scans of the chest, abdomen, and pelvis were normal.

**Figure 2**
(A) Pituitary gland pathology report: dense inflammatory lymphoplasmacytic infiltrates. (B) IgG4-positive immunostaining in more than ten plasma cells per high power field (arrows, 40×).

**Figure 3**
(A and B) Preoperative nonenhanced MRI: an extensive intrasellar lesion expanding into suprasellar region was found, with signs of bleeding, compression and optic chiasm displacement. (C and D) Postoperative nonenhanced MRI under meprednisone treatment showed significant reduction in sellar mass size.
Outcome and follow-up
The patient’s general health improved over subsequent follow-up. A repeat MRI performed 30 days after surgery showed a substantial reduction in sellar lesion size (Fig. 3C and D) and her CVF test improved. The patient continued clinical follow-up and treatment in her home town for eight months from diagnosis, and has not experienced a relapse of the hypophysitis nor any extrapituitary involvement. She remains with anterior hypopituitarism and diabetes insipidus.

Discussion
IgG4-related disease is a fibro-inflammatory condition characterised by lesions with dense lymphoplasmacytic infiltrates, rich in IgG4-positive plasma cells, storiform fibrosis and often, though not always, high plasma IgG4 concentrations (6). Various disorders such as Riedel’s thyroiditis, Mikulicz’s disease, retroperitoneal fibrosis, tubulo-interstitial nephritis, and sclerosing cholangitis, among others, are now considered part of this disease spectrum (5).

IgG4-related hypophysitis was first described in 2004 based on clinical data (7); however, since 2007, the diagnosis normally requires histopathological confirmation (8). To date, 32 cases have been reported (5) (9).

In 2011, Leporati et al. established five criteria for IgG4-related hypophysitis diagnosis: i) the presence of mononuclear cell infiltrates within the pituitary gland rich in plasma cells and lymphocytes, with more than ten IgG4-positive cells; ii) a sellar mass and/or thickened pituitary stalk; iii) biopsy-proven involvement of other organs, with IgG4-positive immunostaining; iv) increased IgG4 serum levels (>140 mg/dl); and v) reduction in sellar mass size and symptom improvement after corticosteroid treatment. The authors propose that IgG4-related hypophysitis diagnosis may be confirmed when any of the following criteria combinations are met: criterion i alone, criteria ii or iii, or ii + iv + v (1).

Although the definitive diagnosis of hypophysitis is histopathological, imaging characteristics such as a symmetrical enlargement of the pituitary gland, rapid intense and uniform gadolinium enhancement, lack of erosion or depression of the sellar floor (2) and thickened pituitary stalk (9) should increase the diagnostic suspicion.

In our study, none of the patient presented characteristic pituitary changes detected by MRI for hypophysitis; instead, the images were heterogeneous and contrast enhancement was irregular. Moreover, in the first case there was considerable invasion of the sphenoid sinus and, in the second, signs of bleeding were present. Because these images did not initially suggest hypophysitis, both patients were submitted to TSS at which time definitive diagnosis of IgG4-related hypophysitis was established under the first Leporati’s criterion, as also occurred in 11 of the 32 cases published to date (5) (9). Baseline IgG4 levels were not assayed since we did not initially suspect IgG4-related disease.

Of the 32 cases reported in the literature, 29 were men aged between 40 and 81 years, and 28 patients had other IgG4-related diseases, mostly pulmonary lesions, retroperitoneal fibrosis, autoimmune pancreatitis, lymphadenopathies, sialoadenitis, pachymeningitis, and kidney

Table 2 Pre-operative laboratory results of case 2.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Results</th>
</tr>
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<tbody>
<tr>
<td>Haematocrit (37–47%)</td>
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<tr>
<td>Haemoglobin (11.5–16 g/dl)</td>
<td>13.6 g/dl</td>
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<tr>
<td>White cell blood count</td>
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<td>Glycaemia (70–110 mg/dl)</td>
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<tr>
<td>Creatinine (0.5–1.2 ng/dl)</td>
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<td>Sodium (135–145 mmol/l)³</td>
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<td>Potassium (3.5–5 mmol/l)</td>
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<tr>
<td>Total bilirubin (0.1–1.4 ng/dl)</td>
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</tr>
<tr>
<td>Alkaline phosphatase (31–100 U/l)</td>
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<tr>
<td>Aspartate aminotransferase</td>
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<tr>
<td>Alanine aminotransferase</td>
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<tr>
<td>Albumin (3.2–5 g/dl)</td>
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</tr>
<tr>
<td>Luteinizing hormone (1–18 mU/ml)</td>
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<td>Follicle-stimulating hormone</td>
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<tr>
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</tr>
<tr>
<td>Thyrotrophin (0.47–4.64 μU/ml)</td>
<td>&lt;0.01 μU/ml</td>
</tr>
<tr>
<td>Free thyroxine (0.7–1.8 ng/dl)</td>
<td>0.83 ng/dl</td>
</tr>
<tr>
<td>TSH (0.34–5.6 mU/l)</td>
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</tr>
<tr>
<td>Cortisol (5–25 μg/dl)</td>
<td>4.4 μg/dl</td>
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<tr>
<td>Adreno corticotropin (10–46 pg/ml)</td>
<td>&lt;5</td>
</tr>
<tr>
<td>Prolactin (5–25 ng/ml)</td>
<td>125 ng/ml</td>
</tr>
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</table>

NV, normal value; IGF1, insulin growth factor 1.

*aUnder desmopressin treatment.

http://www.edmcasereports.com

![Figure 4](https://example.com/figure4.png)

(A) Pituitary gland histopathology report: dense lymphoplasmacytic inflammatory infiltrates. (B) IgG4-positive immunostaining in more than ten plasma cells per high power field (arrows).
lesions (3). In contrast, both patients in our study were young women with no signs of clinical or radiological involvement of other organs at time of diagnosis.

With respect to clinical manifestations, both patients presented hypogonadotrophic hypogonadism and one had diabetes insipidus, in agreement with findings in the literature, where 25 of 32 cases presented varying degrees of hypopituitarism, 21 had diabetes insipidus and 16 patients presented both conditions (5) (9).

Concerning treatment, glucocorticoids are recommended as first-line therapy. More than half of the published cases received prednisolone (0.5–1 mg/kg per day), although some patients respond well to hydrocortisone replacement therapy (5). In our study, both patients presented good response to lower dose of meprednisone.

In conclusion, we report two cases of IgG4-related hypophysitis in young nulliparous women, the youngest to date, without typical radiological pituitary images, in whom the diagnosis was histologically confirmed. Notably, the disease was restricted to the pituitary and response to relatively low-dose glucocorticoids has been favourable.

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
Written informed consent was obtained from the two patients.

References