Development of an Empty Sella in Neurosarcoidosis

Arshpreet Kaur1, Pankaj Sharma2, Zeng Wang2, Charles E Moore3, Mary Koutourousiou4, Aly Abayazeed5, Robert Bert2 and Stephen J Winters1

1Division of Endocrinology, Metabolism and Diabetes, University of Louisville, USA
2Department of Neurology, University of Louisville, USA
3Division of Rheumatology, University of Louisville, USA
4Department of Neurosurgery, University of Louisville, USA
5Department of Radiology, University of Louisville, USA

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*Corresponding author: Arshpreet Kaur, MBBS, 2435 West Belvedere Avenue, Suite #56, Sinai Hospital of Baltimore, Baltimore, MD -21215, USA, Tel: 443-642-7607; Email: arshpreet.31@gmail.com

Abstract

We report the case of a 56 year old African American woman who presented with headache, difficulty walking and loss of hearing and vision who had undiagnosed diabetes insipidus and pan-hypopituitarism. MRI revealed inflammation of the meninges with thickening of the pituitary-infundibulum and hypothalamus with spread to the optic chiasm and optic nerves. Dura biopsy confirmed the clinical diagnosis of isolated neurosarcoidosis. During 8 months of treatment with prednisone and methotrexate, there was partial resolution of the dura thickening but the neurological manifestations and the endocrine disorder were unaffected. Swelling of the pituitary became an empty sella.

Learning Points:

A. The diagnosis and treatment of isolated neurosarcoidosis can be challenging.

B. Neurosarcoidosis can present with hypopituitarism, diabetes insipidus and hyperprolactinemia.

C. This case report demonstrates the evolution of a pituitary mass in a patient with neurosarcoidosis into an empty sella.

Keywords: Neurosarcoidosis; Empty sella; Panhypopituitarism; Pituitary

Introduction

An empty sella develops when arachnoid herniates into the sella turcica through a congenital or acquired defect in the dura, and cerebrospinal fluid (CSF) fills the sella and compresses the pituitary gland. Radiologically, the expanded sella is defined as partially empty if less than 50% is filled with CSF and the pituitary gland thickness is ≥ 3mm, or total, when more than 50% of the sella is filled with CSF and the gland thickness is ≤ 2 mm. A partial empty sella is a common radiographic finding, and is usually asymptomatic, and found incidentally. Endocrine deficiencies may occur, but are much more common in total empty sella [1].

Empty sella is also divided into two groups clinically, primary and secondary. Patients with primary empty sella are more often female and overweight [2], and some have idiopathic intracranial hypertension [3]. An empty sella may also develop following pituitary injury. Disorders associated with secondary empty sella include wrents in the diaphragmata sella, pituitary macroadenomas treated surgically or by radiotherapy or following the spontaneous necrosis of pituitary apoplexy, Sheehan’s syndrome, vasculitis, infection, head trauma, and autoimmune hypophysitis. In several cases, a pituitary mass, believed to represent autoimmune hypophysitis, was documented to convert to an empty sella [4]. For neurosarcoidosis, there are case reports of empty sella by CT [5] and pneumoencephalogram [6] but the condition is rarely mentioned among the causes of empty sella. This case report demonstrates the evolution of a pituitary mass in a patient with neurosarcoidosis into an empty sella.
Case Report

A 56 year-old African American woman with hypertension presented with progressive bilateral vision loss, difficulty walking, vertigo, headache, hearing loss, extreme polyuria and polydipsia, and a 20lb weight loss over 4 months. She had deficits in cranial nerves II, VII and VIII, and a wide based gait.

Results

Endocrine tests indicated pan-hypopituitarism: LH 0.21U/L (reference range in menopause 13.1-86.5), FSH 1.01U/L (reference range in menopause 21.5-131), free T4 0.35ng/dL (reference range 0.78-2.19), TSH 0.44mlU/L (reference range 0.46-4.68), GH 0.2ng/ml (reference range <10), IGF-1 118ng/ml (reference range 50-317; -1.4 SD for age), IGF-BP3 2.5mg/L (reference range 0.78-2.19), TSH 0.44mU/L (reference range in menopause 21.5-131), free T4 0.35ng/dL (reference range 4.5-22.7), and ACTH 6pg/mL (reference range 3.4-6.9), A.M. cortisol 0.83μg/dL (reference range 50-317; -1.4 SD for age), IGF-1 118ng/ml (reference range 0.78-2.19), TSH 0.44mU/L (reference range in menopause 21.5-131), free T4 0.35ng/dL (reference range 4.5-22.7), and ACTH 6pg/mL (reference range 6-50). The PRL level was increased at 56ng/mL (reference range 4.5-22.7). The plasma osmolality was 290mOsm/kg while the urine osmolality was 236mOsm/kg which rose to 827mOsm/kg following dDAVP, consistent with central diabetes insipidus. Magnetic Resonance Imaging (MRI) revealed diffuse dural thickening, pituitary enlargement and mild leftward deviation of the pituitary stalk (Figure 1A). The level of angiotensin converting enzyme in serum 40U/L (reference range 9-67) and CSF 4U/L (reference range <15) was normal. Dural biopsy showed both non-caseating and caseating granulomas with focal scantly necrosis. Chest CT was normal. Isolated neurosarcoidosis was diagnosed after extensive testing to exclude infection and malignancy. She was treated with prednisone and methotrexate and biopsy is needed for definitive diagnosis. Neurosarcoidosis may present with partial or complete anterior hypopituitarism, diabetes insipidus (DI), and hyperprolactinemia [8]. Excess thirst and polyuria may also occur without DI [9]. The diagnosis of isolated neurosarcoidosis can be challenging, as it can present with a wide variety of symptoms and imaging findings [9,10]. Involvement of the leptomeninges can be diffuse, focal or multifocal. The brain parenchyma and spinal cord may be affected, and in some cases the MRI appears normal. Basilar leptomeningeal disease can involve the cranial nerves and the hypothalamus-pituitary unit where the differential diagnosis includes lymphocytic hypophysitis, histiocytosis, tuberculosis, leukemia and metastasis. Measurement of the angiotensin converting enzyme level in serum and CSF is rarely helpful [11], and biopsy is needed for definitive diagnosis. Neurosarcoidosis patients are generally treated with high dose corticosteroids and immunosuppressive agents, and while the neurosarcoid lesions may regress, endocrine dysfunction is usually irreversible [8]. This case demonstrates that empty sella can be the direct outcome of sarcoidosis affecting the pituitary.

Outcome and follow-up

With treatment, her weight increased 37kg, her neurological symptoms were unchanged, and pan-hypopituitarism and diabetes insipidus persisted. She continued to complain of polydipsia and polyuria even when the urine specific gravity was maintained at 1.018. After 12 months of treatment, methotrexate was stopped, and prednisone was slowly reduced to a maintenance dose, with addition of azathioprine for steroid sparing.

Discussion

Sarcoidosis is a multisystem inflammatory disorder that is defined by the histological finding of noncaseating granulomas. While the disorder can affect any organ, the lung, liver, eye and skin are most commonly involved. The nervous system is affected in 5-14% of cases, which may occur in isolation [7]. Patients with neurosarcoidosis may present with partial or complete anterior hypopituitarism, diabetes insipidus (DI), and hyperprolactinemia [8]. Excess thirst and polyuria may also occur without DI [9]. The diagnosis of isolated neurosarcoidosis can be challenging, as it can present with a wide variety of symptoms and imaging findings [9,10]. Involvement of the leptomeninges can be diffuse, focal or multifocal. The brain parenchyma and spinal cord may be affected, and in some cases the MRI appears normal. Basilar leptomeningeal disease can involve the cranial nerves and the hypothalamus-pituitary unit where the differential diagnosis includes lymphocytic hypophysitis, histiocytosis, tuberculosis, leukemia and metastasis. Measurement of the angiotensin converting enzyme level in serum and CSF is rarely helpful [11], and biopsy is needed for definitive diagnosis. Neurosarcoidosis patients are generally treated with high dose corticosteroids and immunosuppressive agents, and while the neurosarcoid lesions may regress, endocrine dysfunction is usually irreversible [8]. This case demonstrates that empty sella can be the direct outcome of sarcoidosis affecting the pituitary.

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