JCIMCR Journal of

OPEN ACCESS Clinical Images and Medical Case Reports

ISSN 2766-7820

Short Report

Open Access, Volume 4

Sheehan syndrome: Clues to underlying diagnosis

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Received: Jun 05, 2023 Accepted: Jul 14, 2023 Published: Jul 21, 2023 Archived: www.jcimcr.org Copyright: © Agrawal SS (2023). DOI: www.doi.org/10.52768/2766-7820/2510

Abstract

As Hypoglycemic disorders are rare in patients without Diabetes Mellitus, a thorough evaluation should be performed only in individuals presenting with Whipple's Triad (Symptoms or signs consistent with hypoglycaemia, a low reliably measured plasma glucose concentration and resolution of these symptoms after plasma glucose is raised). In a seemingly Ill-looking individual, hormonal deficiencies including Cortisol and thyroid should be ruled out after excluding more common causes such as drugs and critical illnesses.

Keywords: Hypoglycemia; Hyponatremia; Cortisol.

Case details

A 46-year-old lady presented to the emergency department with sweating, palpitations, and tremors for the last fifteen minutes. Her plasma glucose was documented to be 46 mg/dl (reference range – 70-180 mg/dl) in emergency. Her Symptoms resolved after the administration of 25% dextrose solution. She had a recent history of low trauma fracture of the neck of the femur. There was no history of Anti-diabetic agents or insulin use in the past. There was a history of traumatic delivery twelve years back, which was followed by secondary amenorrhea and loss of secondary sexual characteristics. Physical examination revealed a pale-alabaster-looking skin, wrinkled face, loss of lateral eyebrows (Figure 1), and hoarse voice. Blood pressure was 92/60 mm Hg while pulse rate was 56/minute at presentation.

Laboratory studies showed Glycated hemoglobin of 5.3%, serum sodium - 128 mEq/L, low free T4 - 0.2 ng/dl with normal TSH of 2.2 mlu/ml and low serum cortisol - 1.5 mcg/dl. Other parameters of anterior pituitary function including, ACTH, Prolactin, IGF-1, were also low. LH and FSH were inappropriately normal for menopausal age.

MRI of the pituitary revealed an Empty sella with thinnedout anterior pituitary (Figure 2). A diagnosis of Sheehan syndrome was made and she was started on intravenous dextrose containing fluids with hydrocortisone supplementation followed by thyroxine replacement. After 6 weeks of follow-up, her serum sodium had improved and there was no new episode of hypoglycaemia.

Discussion

Sheehan syndrome occurs due to ischemic pituitary necrosis induced by vasospasm and thrombosis of hypophyseal arteries [1]. In Sheehan syndrome, Hypoglycaemia can occur due to secondary adrenal insufficiency associated with loss of counterregulatory hormones like Growth Hormone while hypothyroidism and cortisol deficiency decrease free water clearance and cause subsequent hyponatremia [2]. Low bone mineral density occurs due to long-standing estrogen deficiency.

Pale-alabaster-looking-skin occurs due to loss of ACTH and pro-opiomelanocortin related peptides and subsequent decreased stimulation of the skin MC1 receptor and is a clue to **Citation:** Agrawal SS, Bhardwaj S. Sheehan syndrome: Clues to underlying diagnosis. J Clin Images Med Case Rep. 2023; 4(7): 2510.



Figure 1: Showing pale-alabaster-looking skin, wrinkled face and loss of lateral eyebrows.

the underlying secondary adrenal insufficiency [3]. The loss of lateral one third of eyebrows known as "Hertoghe Sign" occurs in hypothyroidism due to increased apoptosis of hair follicles [4].

Central hypoadrenalism and hypothyroidism should always be suspected in patients presenting with unexplained hypoglycaemia and hyponatremia in emergency.

Acknowledgements: None.



Figure 2: Showing Empty Sella with thinned out anterior pituitary.

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