

Almost Two Decades Without Diagnosis: Sheehan Syndrome

Taqwa Miftah Mousa 2348

Supervisor: Rema H Faraj Saad

Libyan International Medical University Faculity Applied Medical Science (AMS)



Introduction

Every year, around 14 million women suffer from postpartum hemorrhage, according to the World Health Organization (WHO). A total blood loss of 1,000 mL or more with concomitant hypovolemia within the first 24 hours after giving delivery is characterized as postpartum hemorrhage. A serious hemorrhage is defined as a blood loss of more than 3,000 milliliters. Although postpartum hemorrhage can be lethal, it has been on the decline for the past two decades. The decrease in mortality has resulted in a surge in the recognition and diagnosis of Sheehan's Syndrome (SS), an uncommon consequence of postpartum hemorrhage [1].

Sheehan's Syndrome

Harold Leeming, a British Pathologist, was the first to Recognize it in 1937. is a long-term health problem that affects women who have had a postpartum hemorrhage and is characterized by varying degrees of Pituitary hormone imbalance produced by the pathological process of pituitary gland necrosis following the hemorrhage [2].

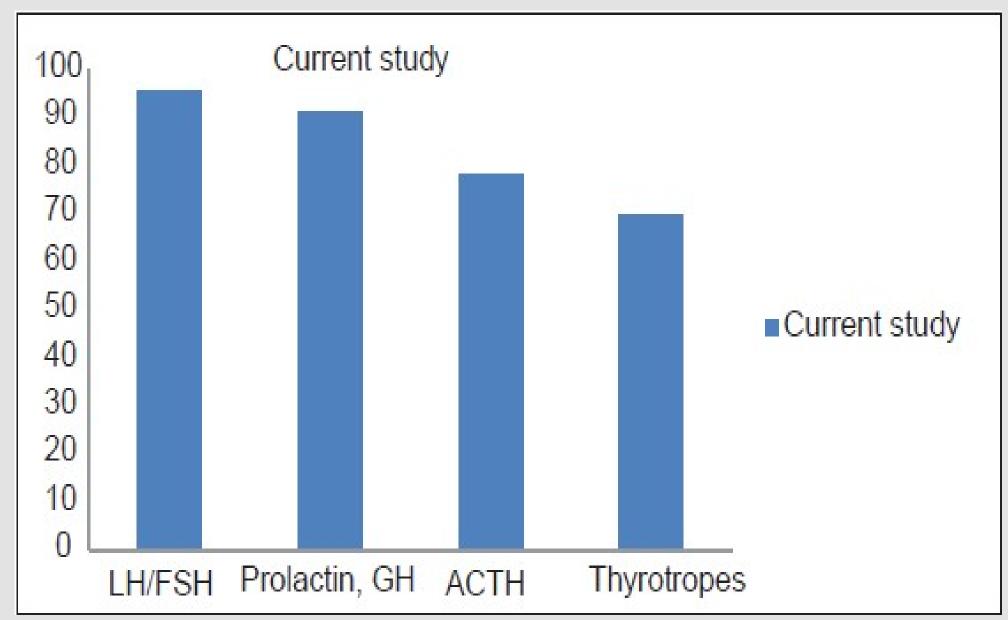


Figure 1: Anterior pituitary assessment [3].

Case Presentation

We describe the case of a 45-year-old Ethiopian woman, presented to the emergency department of our hospital on February 18, 2020, with a complaint of generalized fatigue for 18 years and worsening abdominal pain, vomiting, and diarrhea of 1 month duration. The history dated back 18 years to the birth of her seventh child in home delivery, following which she experienced excessive vaginal bleeding which progressed to anorexia, nausea, vomiting, diarrhea, and abdominal pain of 6 years' duration, for which she was treated symptomatically throughout these years, she had associated weight loss, anorexia, dizziness, cold intolerance, myalgia, arthralgia, and progressive loss of axillary and pubic hair. Complete clinical evaluation, endocrine studies, and pituitary magnetic resonance scan revealed hypopituitarism secondary to Sheehan's syndrome. She had significant improvement noted following the commencement of hormone replacement therapy

Diagnosis

To diagnose Sheehan's, doctor likely will:

- Compile a comprehensive medical history. It's Critical to report any Issues you've had during childbirth.
- Run blood Test. Your pituitary hormone levels will be checked.
- Pituitary Hormone Stimulation Test. You may need pituitary Hormone Stimulation Testing, which is injecting hormones and repeating blood Tests to evaluate how your pituitary reacts.
- Imaging Testing. Imaging studies, such as MRI or CT scan, may be required [6].

Plate 1-17 Postpartum hemorrhage Postpartum hemorrhage Postpartum hemorrhage Postpartum hemorrhage Postpartum hemorrhage Rapid drop in blood pressure Rim of relatively normal pituitary of pregnancy Information Failure of lactation (often first sign postpartum) Adrenal cortical insufficiency (acute initial shock, loss of public and body hair, asthenia, hypoglycemia) Hypothyroidism Hypothyroidism

Figure 2: Pathophysiology of sheehan syndrome

Mangment

Sheehan's Syndrome is treated with long-term hormone replacement therapy. One or more of the drugs listed below:

- Corticosteroids.
- Levothyroxine.
- Estrogen.
- Growth hormone.
- Fertility hormones [6].

Conclusion

We discuss SS, which is postpartum hypopituitarism caused by pituitary infarction and necrosis. SS is caused primarily by a substantial postpartum hemorrhage after labor. manifest with, abdominal pain, vomiting, and diarrhea, anorexia, nausea, vomiting, weight loss, myalgia, arthralgia, and loss of axillary and pubic hair, menstrual irregularities. To replace the hormones that are missing in Sheehan's syndrome, lifelong hormone replacement treatment is used. (Corticosteroids,Levothyroxine,Estrogen,Growth hormone).

References

- 1- E. Gaspar B. Diagnosis and treatment challenges of Sheehan's Syndrome. *Front Womens Health*. 2018;3(3).
- 2-Schury M, Adigun R. Sheehan Syndrome. *StatPearls*. Published online 2021. Accessed January 21, 2022.
- 3-Gutch M, Kumar S, Razi S, et al. Sheehan's syndrome: A clinical, biochemical, hormonal, radiological, bone mineral density and quality of life; assessment. *CHRISMED J Health Res.* 2014;1(2).
- 4-Genetu A, Anemen Y, Abay S, Bante SA, Mihrete KM. A 45-year-old female patient with Sheehan's syndrome presenting with imminent adrenal crisis: a case report. *J Med Case Rep.* 2021;15(1).
- 5-Sheehan's syndrome. Mayoclinic.org. Accessed January 21, 2022.
- 6-Hypopituitarism. Mayo Clinic. Accessed March 19, 2022. https://www.mayoclinic.org/diseases-conditions/hypopituitarism/symptoms-causes.