hypotension preceded by a 6 month history of intense fatigue. She did not take any steroids in the 6 months prior. Initial tests were consistent with secondary adrenal insufficiency; low plasma cortisol (<1ug/dl n 5-23 ug/dl) and ACTH-concentrations below detection (< 4 ng/l), central hypothyroidism: abnormal low TSH (0. 04uU/ml n 0.4-4.5 uU/ml) and low free T4 (0.56 ng/dl n 0.8-2 ng/dl), and growth hormone deficiency (IGF1 below age and sex normal values). Craneal CT and pituitary MRI didn't show any anomaly. Based on symptoms, clinical context and test results the pressumtive diagnosis was AH related to her lupus. The patient was treated with hydrocortisone and levothyroxine with excellent response and complete resolution of her symptoms. Clinical Lesson: This case demonstrates that hypophysitis is a clinical entity that can be suspected and treated based on clinical and past medical history even in the context of normal imaging were an invasive procedure such as a biopsy would likely not change the final outcome.

Presentation: No date and time listed

## Abstract citation ID: bvac150.990

## Neuroendocrinology and Pituitary LBODP074

## Autoimmune Hypophysitis: Diagnosis Beyond Imaging

Juanita Gonzalez, MD<sup>1</sup>, Claudia Monsalve, MD<sup>1</sup>, Carolina Aguilar, MD<sup>2</sup>, Natalia Aristizabal, MD<sup>1</sup>, Jose Luis Torres, MD<sup>2</sup>, Nestor Alfonso Lopez Pompey, MD<sup>3</sup>, Laura Valentina Estupiñan Vargas, MD<sup>1</sup>, Andres Felipe Garcia Ramos, MD<sup>1</sup>, Daisy Buenaventura, MD<sup>1</sup>, Alex Ramirez, MD<sup>1</sup>, Hernando Vargas, MD<sup>1</sup>, and Sebastian Herrera, MD<sup>2</sup> <sup>1</sup>UNIVERSIDAD PONTIFICIA BOLIVARIANA MEDELLN, Medellin, Colombia<sup>2</sup>CLINICA LAS AMERICAS AUNA, Medellin,

Colombia; <sup>3</sup>Hospital San Vicente Fundacin Rionegro, Rionegro Antioquia, Colombia

Background: Lymphocytic (LH) or autoimmune hypophysitis (AH) is a rare inflammatory disorder of the hypophyseal gland, often miss-diagnosed. LH is predominant in females, occasionally related with other rheumatic disorders (1-20%). The disease can lead to pituitary dysfunction and symptoms of hypopituitarism, the most affected axes are the ACTH and TSH. Definitive diagnosis of AH can be based only on pathological examination of a pituitary biopsy sample but such invasive procedure is seldom needed. Different imaging modalities, especially MRI, can be useful but up to 30% can be unremarkable, thus clinical manifestations and biochemical evaluation is of great importance in guiding the diagnosis. Hormonal replacement therapy is the main treatment strategy. Immunosuppressive drugs are indicated by the severity of the symptoms and the underlying autoimmune disease. Clinical Case: A 36 year old patient with active lupus and Sjogren disease presented to the emergency department with syncope and