Postpartum hyponatremic coma

Coma hiponatréxico posparto

A 33-year-old primigravida was admitted to our center for hyponatremic coma 7 days after delivery of a healthy infant. She had no remarkable personal or family history, and was admitted to a private hospital for delivery after an uneventful full term pregnancy. High blood pressure was found upon admission, and a cesarean section was therefore decided, which was complicated by heavy bleeding due to a right uterine ligament tear. Subsequent laboratory tests showed AST levels of 100 U/L (5–40), anemia (hemoglobin 6.5 g/dL 135–145), proteinuria in the nephrotic syndrome range. Four RBC units and seven platelet units were transfused, and urapidil, labetalol 200 mg/6 h, and methyldopa 500 mg/6 h were infused due to sustained HBP. A single dose of cabergoline 1 mg was administered to inhibit lactation on the patient decision.

Seven days after birth, patient reported severe asthenia and headache. Physical examination showed no changes and BP values of 150–155/95–100 mmHg. A complete blood count revealed persistent anemia. A few hours later the patient experienced a tonic–clonic seizure followed by coma and was intubated. She remained afebrile and hemodynamically stable, with a BP of 135/80 mmHg and a HR of 67 bpm. Laboratory tests revealed worsening of anemia, blood glucose of 73 mg/dL, CPK of 1710 U/L, and anemia (hemoglobin 6.5 g/dL). Upon arrival to the ICU, sodium levels of 104 mmol/L (5–40), anemia (hemoglobin 6.5 g/dL), thrombocytopenia (21 × 10^3/μL, 150–450), and proteinuria in the nephrotic syndrome range. Four RBC units and seven platelet units were transfused, and urapidil, labetalol 200 mg/6 h, and methyldopa 500 mg/6 h were infused due to sustained HBP. A single dose of cabergoline 1 mg was administered to inhibit lactation on the patient decision.

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Upon arrival to the ICU, sodium levels of 104 mmol/L (135–145) were found, as well as glucose 78 mg/dL (60–100), urea 19 mg/dL (15–50), creatinine 0.57 mg/dL (0.5–1.2), and serum osmolality of 224 mOsm/kg. Urinary analysis showed Na 41 mmol/L and osmolality of 384 mOsm/kg. There was persistent normocytic and normochromic anemia with hemoglobin 8.2 g/dL, and slightly elevated transaminase levels. Chest X-rays and a CT scan of the head showed no changes, and diffuse brain involvement was found in an electroencephalogram. Isotonic saline, 2000 mL over 5 h, was started at the ICU with no improvement. Urine output during the first admission day ranged from 15 to 50 mL/h. Corticosteroids were therefore decided based on recovery of sodium levels and patient remained conscious and oriented, with no focal neurological signs or new seizures. MRI showed a pituitary gland of normal shape and size, with peripheral enhancement and no central uptake with gadolinium contrast, consistent with non-hemorrhagic subacute adeno-hypophyseal ischemia. Sodium levels remained stable, and a gradual improvement occurred in thyroid hormone levels (free T4 0.62 ng/dL on the sixth day).

Patient was transferred to the endocrinology ward. Improvement in BP levels and disappearance of proteinuria allowed for tapering of antihypertensive drugs. Based on improved hormone control, decreased ischemia in the control MRI, and negative immune tests, corticosteroid tapering was decided. On hospitalization day 27, hormone tests were again performed before discharge, showing basal cortisol of 9 ng/dL, increasing to 14.8 ng/dL (30′) and 5.5 ng/dL (60′) after short stimulation with intravenous ACTH 250 mcg, and improvement in all other parameters. It was therefore decided to prescribe corticosteroid therapy under stress conditions only (Fig. 1).

After discharge, patient did not still require background treatment. Menses returned at 5 months of discharge, and
became regular again at 8 months. Partial cortisol response to short ACTH stimulation test persists, while all other hormones have recovered. An ischemic image persisted at 6 months in MRI.

Analysis of data recorded during her stay at the ICU showed that the patient had TSH levels in the low range with inadequate thyroxine levels after pregnancy. Although this was initially attributed to hemodilution, the trend of free T4 to continue decreasing despite increased sodium levels and administration of high glucocorticoid doses led to suspect central hypothyroidism. As regards the corticotropic axis, patient had inappropriately low basal cortisol levels for the stress situation she experienced, with a virtually normal response to the short ACTH stimulation test. This suggested a recent onset central adrenal insufficiency with an almost preserved capacity to respond to exogenous ACTH of the adrenal glands, which had not atrophied yet. In addition, prolactin appeared to be too decreased for a single 1 mg dose of cabergoline.

It would have been helpful to perform specific pituitary function stimulation tests, but these were not feasible because of the patient’s condition. Thus, we think that the decision to start stress doses of hydrocortisone was adequate, and that thyroxine replacement would also have been appropriate.

Evidence of hypopituitarism combined with a history of obstetric complications with bleeding requiring transfusion and the ischemic image of adenohypophysis in MRI suggested a Sheehan syndrome (SS) as first diagnostic possibility.

While hyponatremia is the most common electrolyte change in SS, it is usually chronic in nature, and although rare in developed countries, hyponatremic coma is part of the acute presentation of SS. The mechanisms by which it occurs range from “adequate” vasopressin hypersecretion to cortisol deficiency and, to a lesser extent, volume depletion (due to labor complications) and hypothyroidism (due to impaired free water clearance), mimicking a syndrome of inappropriate antidiuretic hormone secretion, but we should not forget that this condition is diagnosed by exclusion. Because of its pathophysiology, hyponatremia in SS is characterized by the lack of response to isotonic and hypertonic
saline infusion until this is not associated to replacement glucocorticoid and thyroxine therapy to decrease compensatory vasopressin hypersecretion.3,6

Although thyroxine was not administered to our patient, an analysis of the flowchart tables (Fig. 1) showed that natremia and urine output did not start to improve until hydrocortisone was started. It is of note that 3% hypertonic saline was not administered at the ICU to a patient with severe (Na level of 115 mmol/L or less) symptomatic hyponatremia with seizures and subsequent coma suggesting acute hyponatremic encephalopathy, because women of childbearing age are known to be part of the group at a greatest risk of ischemia, brain herniation with brain stem compression, and death.3

Although the fear to use hypertonic saline because of the risk of occurrence of an osmotic demyelination syndrome is understandable, the risk does not lie so much in its use but in its delayed discontinuation. Close monitoring is therefore required to prevent increases in sodium levels greater than 2 mmol/L/h, 10 mmol/L/24 h or 18 mmol/L/48 h. This would explain desmopressin administration (although this was a debatable treatment) at the ICU when rapid increases were seen in sodium levels associated to urinary frequency.

In addition to the exceptional acute presentation of SS in this patient, a decrease in ischemic surface was seen in MRI during its course, as well as a gradual recovery of thyroid hormones (Fig. 2), TSH, cortisol, and prolactin which allowed for treatment discontinuation. The condition therefore appears to be a transient SS, already reported in the literature, although most recoveries from it were only seen in the gonadotropic and/or lactotropic axes,8,9 and recovery of the corticotropic and somatotropic occurred in a single case.2

Finally, this case emphasizes the significance of laboratory monitoring and especially electrolyte levels in patients with complicated labor, and the need for early recognition of a condition that may be fatal for women.7

References

Visceral leishmaniasis in a type 1 diabetic patient with isolated pancreas transplant

Leishmaniasis visceral en un paciente con diabetes tipo 1 y trasplante aislado de páncreas

Isolated pancreas transplant (IPT) is a treatment allowing for long-term normalization of carbohydrate metabolism in patients with type 1 diabetes mellitus (T1DM). It is currently indiced for patients with labile T1DM with frequent and severe acute metabolic complications and preserved kidney function.1,2

We report below the case of a 31-year-old male patient with T1DM starting 16 years before who showed poor metabolic control, highly variable blood glucose levels, acute diabetic complications (repeated severe undetected hypoglycemic episodes and frequent episodes of ketosis and ketoacidosis), and chronic complications (proliferative retinopathy, established nephropathy with preserved glomerular filtration rate, peripheral and autonomic sensorimotor neuropathy: gastroparesis requiring jejunostomy for feeding and neurogenic bladder with daily self-catheterization and recurrent urinary tract infection). Patient attended the endocrinology department of another hospital and was referred to our center to be assessed for IPT. He was treated with rapid-acting insulin analogues and insulin glargine using a basal-bolus regimen, and had glycosylated hemoglobin (HbA1c) levels ranging from 10% to 12%.

Despite treatment optimization with an integrated system of continuous glucose monitoring and continuous subcutaneous insulin infusion, recurrent undetected hypoglycemia persisted, and IPT was therefore performed on November 2008. Patient had an uneventful postoperative course, and although insulin therapy could not be discontinued (insulin detemir at approximately 30 IU/day continued to be given), controls showed stable blood glucose levels, hypoglycemia did not recur, and tests showed HbA1c values around 5.5% and C peptide levels of 1.5 ng/mL.

Four months after transplant, patient reported malaise, fever up to 40°C with chills, irritative cough, watery diarrhea, and weight loss for the past 20 days. He was taking immunosuppressants (prednisone, mycophenolate mofetil, and tacrolimus), trimethoprim–sulfamethoxazole at prophylactic doses, insulin detemir, and antihypertensives (losartan, atenolol, manidipine, furosemide, and doxazosin). On admission, patient was conscious and oriented, with a moderately impaired general condition, tachycardia, fever, and mild skin and mucosal pallor. Small purpuric lesions were seen in right nostril mucosa. Cardiorespiratory auscultation was normal. No changes were found in abdomen and limbs. Supplemental tests on admission showed the following results: hemoglobin 8.5 g/dL, hematocrit 27%, platelet count 62,000/L, WBC count 1800/L (absolute neutrophils 1300/L, absolute lymphocytes 290/L), and creatinine 2.9 mg/dL; results of all other biochemical tests, including liver profile, were normal.

Blood, urine, sputum, nasal exudate, and catheter tip cultures were all negative. Cytomegalovirus, Epstein–Barr virus, parvovirus B19, and herpesvirus 6 viral loads were all negative. An echocardiogram and a Doppler abdominal ultrasound of the graft were normal, and computed tomography (CT) of the chest only showed a mild pericardial effusion. An abdominal CT revealed an enlarged spleen with infarction of 50% of its volume, no pancreatic graft changes, and no abdominal collections. Patient was initially treated with broad spectrum antibiotic therapy, with no response of fever. Bone marrow aspiration was performed because of the presence of pancytopenia and splenomegaly, showing