An Uncommon Cause of Hypernatremia in Very Low Birth Weight Premature Infants: Idiopathic Central Diabetes Insipidus

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Central diabetes insipidus (CDI) is a water homeostasis disorder characterized by an inability to concentrate urine because of insufficient production of antidiuretic hormone. Dehydration with hypernatremia can occur during the neonatal period in preterm neonates in association with insensible water loss, high urine output, and reduced sodium excretion. A high index of suspicion is required to diagnose CDI in preterm neonates. We report two cases, who presented persistent hypernatremia with polyuria despite increased fluid supply and low sodium intake. CDI diagnosis was confirmed by the therapeutic test with oral vasopressin analog. Investigations were all normal; CDI was considered idiopathic. Persistent hypernatremia despite increased fluid intake with polyuria, hypothenuria, low urine output, and high plasma osmolality is the key point for the diagnosis.

Keywords: Central diabetes insipidus, premature infants

INTRODUCTION

Diabetes insipidus (DI) is a water homeostasis disorder characterized by an inability to concentrate urine because of insufficient production of antidiuretic hormone (ADH) (central diabetes insipidus, CDI) or due to impaired kidney response to ADH (nephrogenic diabetes insipidus).1,2 CDI in a neonate is usually associated with congenital abnormalities of the central nervous system, hypoxic-ischemic encephalopathy, meningitis, encephalitis, or severe intraventricular hemorrhage (IVH) in preterm neonates.3 During neonatal period, dehydration with hypernatremia can occur due to insensible water loss, high urine output, and reduced sodium excretion in preterm neonates. Therefore, a high index of suspicion is required to diagnose CDI in very low birth weight infants.2–4

To alert for CDI diagnosis and early treatment in the neonatal period, two cases of very low birth infants with idiopathic CDI, who were successfully controlled through lyophilized sublingual desmopressin, were reported.

CASE PRESENTATION

Case 1

A male preterm neonate with a gestational age of 31 weeks and a birth weight of 1215 g was born by cesarean section (C/S) with Apgar scores of 4 and 7 at 1st and 5th minutes, respectively. He was on nasal intermittent positive-pressure ventilation and did not require surfactant therapy. Antibiotic therapy was begun because of high CRP level. On the 5th day of life due to increase in CRP, oxygen requirement (FiO2 > 40%), and respiratory acidosis, he was intubated, surfactant was given, and antibiotic treatment was changed. He was extubated the next day followed by noninvasive ventilation for 2 weeks. On the 13th day of life, although the daily Na intake was 1.2 meq kg–1 d–1 and the daily fluid supply was 180 mL kg–1 d–1, the serum Na level and urine output were 152 mmol L–1 and 8 mL kg–1 h–1, respectively. The density of urine was around 1005. The ADH level was 2.30 pmol L–1. Thus, DI was diagnosed and desmopressin treatment was begun (2 μg per oral). The urine output decreased to 4-5 mL kg–1 h–1 and the plasma Na level decreased to 140 mmol L–1. Transfontanelle sonography did not demonstrate IVH. The evaluation of the pituitary axes revealed normal thyroid, adrenal, and gonadal functions. Serological tests for syphilis, toxoplasmosis, cytomegalovirus, herpes simplex, and rubella (TORCHS) ruled out
these congenital infections. No abnormalities were observed in serum urea nitrogen, creatinine, potassium, calcium, and bicarbonate levels. Magnetic resonance imaging (MRI) of the brain was planned to evaluate pituitary gland and the other structures of the brain, but parents did not approve it because of sedation. The etiology remains unknown. He was discharged on postnatal on 52nd day of life, at 38/4 corrected age, weighing 2285 g, with a prescription of 2 × 1.5 mcg of oral desmopressin per day.

Case 2
A male preterm neonate with a gestational age of 31 weeks and a birth weight of 1015 g was born by C/S with Apgar scores of 4 and 7 at 1st and 5th minutes, respectively. He was intubated in the delivery room, surfactant was given for the diagnosis of respiratory distress syndrome, and then extubated followed by noninvasive ventilation for 10 days. On the 12th day of life, respiratory distress reappeared and oxygen requirement increased (FiO₂ 40%). He was thought to develop pneumonia and re-intubated. He was extubated on the next day followed by noninvasive ventilation for 2 weeks. On the 24th day of life, we noticed that the urine output (7.78 mL kg⁻¹ h⁻¹) and plasma Na level were high, while he was receiving Na supply at a maintenance dose of 3 meq kg⁻¹ d⁻¹. The urine osmolality was low (153 mOsm kg⁻¹), serum osmolality (292 mOsm kg⁻¹) and plasma Na level (146 mmol L⁻¹) were high. DI was thought and lyophilized sublingual desmopressin (2 × 3 mcg) was begun. After treatment, the serum Na level was between 139 and 142 mmol L⁻¹, and the urine output decreased to 4-5 mL kg⁻¹ d⁻¹. Due to desmopressin response, CDI diagnosis was made. Transfontanelle sonography did not demonstrate IVH. The evaluation of pituitary axes revealed normal thyroid, adrenal, and gonadal functions. Serological tests for TORCHS were normal. No abnormalities were observed in serum urea nitrogen, creatinine, potassium, calcium, and bicarbonate levels. Desmopressin was used for 23 days. Before discharge, we stopped desmopressin treatment. After 2 days without desmopressin treatment, the urine density was 1015, and plasma and urine osmolalities were 281 and 123 mOsm kg⁻¹, respectively; the plasma Na and plasma ADH levels were 142 mmol L⁻¹ and 3.41 pmol L⁻¹. Therefore, the patient was discharged without treatment. Ten days after discharge, the serum Na level and urine density were 141 mmol L⁻¹ and 1030, respectively. CDI disappeared, but the etiology remained unknown.

DISCUSSION
In very-low-birth weight infants, fluid-electrolyte homeostasis in the first week of life is generally characterized by low urine output in the first few days and polyuria thereafter. This causes physiological weight loss and an increase in the serum Na concentration, which is aggravated by transepidermal water loss. Generally, this diuretic phase gets over by the end of the first week of life, and hyponatremia is rare thereafter. Persistent hyponatremia despite increased fluid intake with polyuria, hyposthenuria, low urinary osmolality, and high plasma osmolality should alert the clinician for the diagnosis of DI. In present cases, despite the increase in the total fluid intake, patients’ serum Na and urine outputs remained elevated. The detection of low urine osmolality with high serum osmolality and good response to desmopressin confirmed the CDI diagnosis. Until now, underlying causes of CDI reported are intraventricular hemorrhage, meningitis, septo-optic dysplasia, Listeria monocytogenes sepsis, congenital cytomegalovirus infection, midline intracranial defects, and following surgical resection of a suprasellar mass. Idiopathic CDI accounts for 12-24% of the cases, but idiopathic CDI prevalence in premature infants is higher. Diagnostic work-up, including the level of hypothalamic-pituitary axis hormones, infectious causes such as serological tests for TORCHS of present cases, were all normal. Transfontanelle sonography revealed no intraventricular hemorrhage.

ADH analog—desmopressin, which is available in three different forms, namely, oral, parenteral, and intranasal preparations, is used to reduce urine output and to decrease serum Na levels to normal range. Many studies recommended oral desmopressin lyophilisate because of its efficiency, ease of use, and better tolerance. Regardless of the route, treatment requires careful adjustment of dose as neonates rely on liquid diet and desmopressin may cause fluid overload with wide fluctuations in serum Na levels. Thus, beginning with low dosage twice a day and then increasing according to serum Na levels and urine output should be better. Both the cases were treated with sublingual lyophilized desmopressin twice a day. Similarly, Ozaydin et al. and Atasay et al. used sublingual desmopressin lyophilisate at a dosage of 2.5 μg kg⁻¹ d⁻¹ to manage CDI in very low birth weight premature infants.

Karthiskeyan et al. reported the first case series of CDI. The most common cause was septo-optic dysplasia but of note idiopathic isolated CDI was diagnosed in three of every premature infants. None of the preterm infants had significant intracranial hemorrhage. The median serum Na and serum and urine osmolalities at diagnosis were 156 mmol L⁻¹ (range: 145-175), 320 (range: 300-345) and 112 mOsm kg⁻¹ (range: 66-322), respectively, as reported cases.

In conclusion, polyuria despite increased fluid intake, hyposthenuria, low urinary, and high plasma osmolality and persistent hyponatremia are the key points for the diagnosis.

Main Point
- Central diapedes insipidus is rare and diagnosis is difficult in preterm infants.
- Persistent hyponatremia, dehydration and polyuria beyond first week of life should alert neonatologists.
- Oral desmopressin should be started as soon as the diagnosis is suspected.
- This treatment should be begun with low dosage and be closely monitored.

Ethics Committee Approval: N/A
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