



GLOBAL JOURNAL OF MEDICAL RESEARCH: F  
DISEASES

Volume 23 Issue 10 Version 1.0 Year 2023

Type: Double Blind Peer Reviewed International Research Journal

Publisher: Global Journals

Online ISSN: 2249-4618 & Print ISSN: 0975-5888

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By Aida Correia de Azevedo, Ana Sofia Rodrigues,  
Beatriz Parreira de Andrade, Raquel Cardoso & Susana Lopes

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A female newborn, 12 days old (birth weight: 2800g), was brought to the Emergency Department due to a 30% decrease in birth weight and reduced urine output. The newborn was solely breastfed every 2 hours, with a good notion of adequate reflexes and tolerance. Physical examination revealed a skeletal appearance, jaundice, sunken eyes and skin turgor. Blood pressure measurement was unsuccessful, while capillary blood glucose was 96g/dL. A saline bolus (10 mL/Kg) was administered. Venous blood gas analysis showed pH 7.39, lactate 4.3 mmol/L, HCO<sub>3</sub><sup>-</sup> 21.8 mmol/L, and Na<sup>+</sup> 180 mEq/L.

**Keywords:** *breastmilk; exclusive breastfeeding; hypernatremia; hypernatremic dehydration; newborn.*

**GJMR-F Classification:** *NLMC Code: RJ45*



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# Severe Hypernatremic Dehydration in a Neonate

Aida Correia de Azevedo <sup>α</sup>, Ana Sofia Rodrigues <sup>σ</sup>, Beatriz Parreira de Andrade <sup>ρ</sup>,  
Raquel Cardoso <sup>ω</sup> & Susana Lopes <sup>¥</sup>

**Abstract-** Hypernatremia, a rare condition in newborns, is marked by elevated plasma sodium levels exceeding 150 mEq/L. It is more common in newborns who are exclusively breastfed or with excessive weight loss. Hypernatremic dehydration (HD) presents severe risks, including cerebral edema and other neurological complications.

A female newborn, 12 days old (birth weight: 2800g), was brought to the Emergency Department due to a 30% decrease in birth weight and reduced urine output. The newborn was solely breastfed every 2 hours, with a good notion of adequate reflexes and tolerance. Physical examination revealed a skeletal appearance, jaundice, sunken eyes and skin turgor. Blood pressure measurement was unsuccessful, while capillary blood glucose was 96g/dL. A saline bolus (10 mL/Kg) was administered. Venous blood gas analysis showed pH 7.39, lactate 4.3 mmol/L, HCO<sub>3</sub><sup>-</sup> 21.8 mmol/L, and Na<sup>+</sup> 180 mEq/L. Treatment for dehydration began intravenously following the protocol. Laboratory tests showed normal blood count, Na<sup>+</sup> 178 mEq/L, K<sup>+</sup> 4.5 mEq/L, Cl<sup>-</sup> 143 mEq/L, urea 191 mg/dL, creatinine 1.39 mg/dL, and C-reactive protein 1.45 mg/dL. Formula milk was introduced, and the NB was admitted to the Neonatology unit for observation. The newborn remained stable, with gradual normalization of analytical parameters, along with sterile blood and urine cultures and a normal cephalic ultrasound. Post-discharge, the newborn underwent follow-up appointments, showing good weight evolution and normal neurodevelopment.

HD is potentially life-threatening, emphasizing the critical need for early diagnosis and appropriate treatment. It stands as a significant preventable cause of infant morbidity and mortality. Providing information and guidance on newborn feeding, as well as maintaining vigilant monitoring of risk factors for breastfeeding failure in an outpatient setting, is crucial for success, particularly in cases where the mother lacks necessary information upon discharge.

**Keywords:** *breastmilk; exclusive breastfeeding; hypernatremia; hypernatremic dehydration; newborn.*

## I. INTRODUCTION

Hypernatremia consists of a serum sodium concentration exceeding 145 mEq/L, constituting a relatively infrequent condition in newborns [1-7]. The exact incidence of hypernatremia in newborns remains uncertain but is estimated to range between 1% to 5.6% [1,6,8]. In neonates, hypernatremic dehydration (HD) is more commonly observed in those exclusively breastfed or with excessive weight loss, with early

discharge from the hospital being a significant risk factor. In these cases, HD is associated with ineffective lactation due to insufficient milk production or inadequate support [1-7,9-11].

Initial symptoms of HD are nonspecific, which can lead to delayed diagnosis and subsequent treatment. Consequently, many newborns with HD experience prolonged and severe hypernatremia, increasing the risk of serious complications such as seizures, thrombosis, intracranial hemorrhage, metabolic acidosis, acute kidney injury (AKI) and disseminated intravascular coagulation [1-11]. These complications are directly associated with hypernatremia and its treatment, and the severity of the symptoms does not correlate with blood osmolarity [1,2].

The management of HD involves identifying and addressing the underlying cause, as well as administering intravenous fluids to correct sodium levels.

This correction should occur at a controlled rate of 0.5-0.6 mEq/L per hour, aiming to prevent complications from a rapid decrease in sodium levels, particularly cerebral edema [1-3].

Despite the potential for severe consequences, most newborns recover fully without any lasting effects [1-3,9].

The authors have presented a case report of a newborn with severe HD. This report was previously showcased as an abstract at the 2021 European Academy of Pediatrics Meeting on April 25, 2021.

## II. CASE PRESENTATION

A 12-day-old female newborn was referred by her family doctor to the Emergency Room due to significant weight loss and decreased urination. The infant was delivered at 39 weeks via cesarean section because of a pelvic presentation and the Apgar score was 9/10. Maternal serological tests showed negative results, and all prenatal ultrasounds reported normal findings.

At birth, the newborn weighed 2800 grams (15th percentile;  $-2 < z\text{-score} < 0$ ) and measured 46 centimeters in length (3rd-15th percentile;  $-2 < z\text{-score} < 0$ ). Since delivery, she had been exclusively breastfed every two hours, exhibiting satisfactory suction reflexes and tolerance (the newborn was discharged home at the age of two days).

*Corresponding Author α: Department of Pediatrics and Neonatology, Médio Ave Hospital Center, Vila Nova de Famalicão, Portugal.  
e-mail: aidac.azevedo@gmail.com*

*Author σ ρ ω ¥: Department of Pediatrics and Neonatology, Médio Ave Hospital Center, Vila Nova de Famalicão, Portugal.*

During the ER assessment, the newborn's weight was measured at 1940 grams, indicating a 30% weight loss compared to birth weight. She was emaciated, jaundiced and had sunken eyes (refer to Figure 1) and fontanelle. Capillary refill time was 3 seconds, heart rate was 123 bpm, blood pressure was immeasurable, and capillary blood glucose levels measured 96 mg/dL. A saline bolus (10 mL/kg) was administered with a good response in blood pressure. The newborn also appeared less responsive, with weak crying and mild global hypotonia; the remainder of the neurological examination was ordinary.

Venous blood gas analysis displayed the following values: pH 7.39, lactate 4.3 mmol/L, HCO<sub>3</sub><sup>-</sup> 21.8 mmol/L, and Na<sup>+</sup> 180 mEq/L. Following HD's approach guidelines, intravenous treatment for dehydration was initiated. Laboratory results indicated a normal blood count, with Na<sup>+</sup> levels 178 mEq/L, K<sup>+</sup> 4.5 mEq/L, Cl<sup>-</sup> 143 mEq/L, urea 191 mg/dL, creatinine 1.39 mg/dL, and C-reactive protein 1.45 mg/dL. The coagulation study yielded average results. Both blood and urine cultures returned negative results.

The protocol implemented to address the newborn's hyponatremia ensured a gradual reduction in sodium levels, aiming for a decrease of 0.5-0.6 mEq/L per hour. This process necessitated the adjustment of fluid infusion rates according to sodium levels, which were monitored every one to two hours during the initial phase of treatment. The infant remained stable, exhibiting a gradual normalization of analytical parameters, achieving complete sodium level normalization 60 hours after admission. Throughout the treatment period, formula milk was introduced and adjusted per newborn's tolerance.

An encephalic ultrasound was conducted to rule out any potential brain damage resulting from hyponatremia and its treatment, which revealed no abnormalities.

After a seven-day hospital stay, the newborn was discharged and continued follow-up care, showing positive weight gain and normal neurodevelopment.



*Figure 1:* Newborn with hypernatremic dehydration. Physical examination shows an emaciated appearance, jaundice, sunken eyes and skin turgor.

### III. DISCUSSION

HD is a severe condition defined by elevated sodium levels above 145-150 mEq/L, often stemming from excessive fluid loss or inadequate fluid intake, with the latter being a prevalent cause in neonates. While it occurs more frequently in premature infants due to feeding immaturity, it can also affect full-term healthy newborns, mainly due to breastfeeding issues in the initial weeks of life [1-3,5-7,9-11]. In our case, the newborn exhibited unmistakable signs of dehydration linked to insufficient milk intake, notably a 30% decrease in birth weight and a skeletal appearance. The likely cause for breastfeeding failure was inadequate support provided to the mother, compounded by limited guidance during the prevailing pandemic, where the mother did not receive adequate support from both family and healthcare providers. This situation allowed inadequate breastfeeding to persist until the first appointment for the newborn, which, within the Portuguese national healthcare system, can take place up to the fifteenth day of life.

Several risk factors for HD exist, including cesarean delivery, primiparity, breast anomalies, excessive pre-pregnancy maternal weight, delayed initiation of the first breastfeeding, and lack of prior breastfeeding experience [3,4,6]. In the case described, the mother was a primipara and had no previous experience or knowledge about breastfeeding. Additionally, the newborn was delivered via cesarean section.

Neonates with HD often present well, as classic signs of dehydration, such as sunken eyes and depressed fontanelles, may not be initially evident. Parents may misinterpret certain signs, like the lack of crying or prolonged sleep, as indicators of satiety, leading to a delayed diagnosis [2,4,8-9]. Initial signs of HD are subtle and nonspecific, like lethargy, agitation, and irritability, progressing to severe symptoms including seizures, coma, and potentially death [1-5,8-9]. In our case, the parents perceived prolonged sleep and rare crying as positive signs, assuming the child's well-being.

The diagnosis of HD is typically incidental during newborn weight checks, revealing a weight loss higher than 10% compared to birth weight, prompting comprehensive evaluation, including medical history, physical examination, and blood tests [1,2,5,7,11]. In our case, the family doctor immediately referred the infant to the ER upon finding a 30% weight loss for further investigation and treatment.

Treating HD involves identifying and addressing the underlying cause while administering intravenous fluids to restore sodium levels to normal values. The challenge lies in preventing rapid sodium level decreases, which can lead to severe problems such as intracranial hemorrhage, thrombosis, and cerebral

edema, particularly within the first 24 hours of dehydration [1,2,5-7,10-11]. Sodium correction might take up to 48 hours or more, maintaining a decrease rate of 15 mEq/L in 24 hours or 0.5-0.6 mEq/L per hour [1-4,6-7]. In mild cases (sodium values below 160mEq/L), the infant may be given oral feeds at 1.5 times the maintenance with breast milk or formula milk [3,11]. For more severe cases, the fluid choice depends on the sodium level, aiming to prevent rapid changes. If the sodium levels are higher than 170 mEq/L, a customized solution aligning with the patient's osmolarity is recommended to avoid complications [1,2,5-7,9]. Regular monitoring and adjusted infusion rates prevented complications, resulting in normalized sodium levels 60 hours after admission with no evidence of cerebral edema.

While higher mortality rates are associated with sodium levels exceeding 160 mmol/L, newborns with HD typically make a full recovery without sequelae [2,3,8]. Indeed, in a cross-sectional study involving 46 infants diagnosed with HD, none exhibited central nervous system complications. Approximately half of the participants presented with AKI upon admission, which subsequently resolved entirely [3]. In a separate study encompassing 65 infants diagnosed with HD, over 12% of the neonates developed intracranial hemorrhage, while 12.8% experienced cerebral edema [8]. In our particular case, the newborn presented with AKI, which was resolved through appropriate correction of dehydration. Subsequently, the newborn showed complete recovery without cerebral complications, evidenced in follow-up appointments with normal encephalic ultrasounds and adequate psychomotor development.

It is crucial to maintain close follow-up in newborns who are exclusively breastfed to ensure adequate milk intake and to avoid serious conditions such as the one presented.

### IV. CONCLUSIONS

HD carries the potential for significant complications, such as intracranial hemorrhage and disseminated intravascular coagulation. The primary recognized risk factor for HD is inadequate milk intake, primarily stemming from ineffective lactation and insufficient knowledge about breastfeeding practices.

Therefore, this case report draws attention to the severe consequences resulting from breastfeeding failure, underscoring the critical need to educate and support mothers in breastfeeding. Providing ample guidance and information on breastfeeding techniques, healthy practices, and identifying signs of breastfeeding failure is imperative for successful breastfeeding. Additionally, emphasizing the necessity for close monitoring of newborns during the initial weeks of life through regular weight assessments and

comprehensive physical examinations is essential to detect and manage conditions like HD promptly.

The benefits of breastfeeding are extensive and irreplaceable. Hence, breastfeeding promotion is crucial, requiring adequate support and guidance for mothers throughout this process.

#### Disclosures

Informed consent was obtained for the use of the newborn's photo.

The authors don't have any conflict of interest to declare.

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