Hydranencephaly in a newborn due to occupational toluene exposure during pregnancy: a case report

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The etiopathogenesis of hydranencephaly remains unclear; however, exposure to toxic substances during pregnancy likely increases hydranencephaly risk. Head computed tomography (CT) was performed in a neonate 9 hours post-delivery because the anterior fontanelle was large and there were clinical signs of encephalopathy. Head CT revealed a lack of both cerebral hemispheres and significant cystic enlargement, while the cerebellar hemispheres and pons were found to have developed normally. History-taking revealed that the mother worked in the automotive industry, specifically in the car paint cleaning business and was exposed to toluene during the pregnancy. The patient was diagnosed with hydranencephaly, central diabetes insipidus and central hypothyroidism. Due to the increased head circumference and tense anterior fontanelle, a ventriculoperitoneal shunt was placed. Toluene exposure during pregnancy should be considered among the causes of hydranencephaly. Furthermore, central diabetes insipidus and central hypothyroidism may develop in such cases.

Key Words: hydranencephaly, hypothyroidism; newborn, toluene

Hydranencephaly (HE) is a rare congenital neurological disorder that usually occurs in the second trimester of gestation as a result of the destruction of the cerebral hemispheres. The cerebral hemispheres are commonly replaced by a membranous sac filled with cerebrospinal fluid (CSF) [1]. The etiopathogenesis of HE remains unclear, but most researchers think that HE is associated with brain damage due to the involvement of bilateral internal carotid arteries in the early antenatal period [2]. Other associated conditions are intrauterine infections, rare genetic syndromes, and exposure to toxic substances during pregnancy [3]. This brain malformation often results in fetal loss and therefore is rarely seen postnatally. The incidence of HE also remains unclear. Toluene is an industrial chemical produced during petroleum distillation that is widely used as a solvent. Here, we present the case of a newborn who was born with HE from a mother with occupational exposure to toluene. Toxic exposure started before pregnancy and continued throughout the pregnancy. The neonate also developed central diabetes insipidus (CDI) and central hypothyroidism as a result of the HE.

Case Report

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CASE REPORT

Written informed consent for the study and publication of the photographs was obtained from the patient’s family. A female neonate was born as the fourth pregnancy of a 22-year-old mother who had no prenatal care. The delivery was spontaneous at home at term with a weight of 3,300 g and the neonate cried spontaneously at birth. She was brought to our institution 9 hours after birth due to respiratory distress. On physical examination, weight was 3,300 g (25–50 percentile), height 50 cm (50–75 percentile), and head circumference was 34 cm (50–75 percentile). Vital signs showed arterial blood pressure, 54/32(24) mm Hg; heart rate, 186 beats/minute; and oxygen saturation, 80%. The patient was in poor condition and was intubated. The anterior fontanelle was open and large. Newborn reflexes were weak. Moro and rooting reflexes were absent. There was biting when the suck reflex was tested.

Given the large anterior fontanelle and signs of encephalopathy, head computed tomography (CT) and cranial magnetic resonance imaging were performed on the day of admission. Both cerebral hemispheres were absent on neuroimaging. There was cystic enlargement bilaterally with intact cerebellar hemispheres and pons (Figure 1). Echocardiography and abdominal ultrasonography were both normal. The patient’s 7th-day TORCH serology was negative. There was no growth in CSF culture. CSF polymerase chain reaction for *Toxoplasma gondii* was negative. Chromosome analysis revealed that the patient was 46XX.

The mother stated that she did not know she was pregnant until approximately 24 weeks of gestation, and that she had not received antenatal care. She had been working in the auto industry for 8 hours a day on average before and throughout the pregnancy until delivery, including cleaning cars with paint thinner. On postnatal day 21, the neonate’s urine output was polyuric (10 ml/kg/hr) with blood osmolarity of 353 mmol/kg, serum sodium 172 mEq/L, urine osmolarity 400 mOsm/kg. Tandem metabolic screening with urine and blood amino acids was normal. Thyroid stimulating hormone was 100 µU/mL and free T4 was 0.04 ng/dl. In addition to HE, CDI was diagnosed due to hypernatremia, polyuria, high serum osmolarity and low urine osmolarity, and oral desmopressin (5 µg/day) and L-thyroxine 12 µg/kg/day were started for central hypothyroidism.

On postnatal day 32, the neonate was taken off the mechanical ventilator and given oxygen using a hood. Full oral feeds were also started as the suck reflex was active. Oxygen therapy was discontinued on postnatal day 42. Neurosurgery was consulted because of an increase in head circumference of 2 cm over 3 days on postnatal day 65. The patient also developed tachypnea and an increased oxygen need on day 51 of life. The patient’s weekly head circumference is shown in Figure 2. Head CT continued to show a lack of cerebral tissue bilaterally with cystic enlargement. The cerebellar hemispheres and pons remained normal. Phenobarbital treatment was started due

![Figure 1. Cranial computed tomography (A) and cranial magnetic resonance imaging (B) taken due to head circumference on the 1 postnatal day.](https://www.accjournal.org)
to pedaling and convulsive movements of the arms, leading to suspicion of neonatal seizures. Due to the increase in the patient's head circumference and anterior fontanelle tension, a ventriculoperitoneal shunt was placed by the neurosurgeon (Figure 3). The patient required a tracheostomy and percutaneous endoscopic gastrostomy because she could not be extubated on postnatal day 222. The patient was transferred to a center with a pediatric intensive care unit.

**DISCUSSION**

HE is a very rare brain anomaly that typically occurs in isola-
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growth retardation, convulsive seizures, central hypothyroidism between 4 days and 4 years of age. They found macrocephaly, midline congenital malformations such as holoprosencephaly and septo-optic dysplasia, but its relationship with HE is less well understood. Omar et al. [10] reported seven cases of HE between 4 days and 4 years of age. They found macrocephaly, growth retardation, convulsive seizures, central hypothyroidism, hypocortisolemia, and panhypopituitarism in these cases. In our case, CDI and central hypothyroidism were observed. In conclusion, HE may develop as a result of occupational exposure to toluene in women of reproductive age. CDI may also occur in neonates with HE.

CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.

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Conceptualization: BA, EB. Data curation: BA, EB. Formal analysis: EB, SA. Methodology: BA, EB. Project administration: BA, BG. Visualization: BA, EB. Writing—original draft: BA, BG. Writing—review & editing: EB, BA.

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