Noncystic white matter injury – a case report

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Magnetic resonance imaging (MRI) of the brain in preterm infants at term-equivalent age demonstrated that apart from cystic periventricular leukomalacia (PVL), noncystic white matter injury may take place, detected as diffuse excessive high signal intensity (DEHSI) in the white matter on T2-weighted imaging.

Magnetic resonance imaging of the brain is conducted in few neonatal intensive care units. Consequently, the literature on the subject lacks descriptions of sequelae of noncystic white matter injury in premature newborns with very low birth weight (VLBW).

We present the results of a three-year long observation of a child born at the 27th week of pregnancy diagnosed with DEHSI. The boy exhibited cerebral palsy, hyperexcitability and hypoacusis.

In the authors’ opinion, noncystic white matter injury may not just be one of the reasons for cognitive/behavioral deficits - it may also be responsible for some cases of cerebral palsy in premature infants.

Key words: magnetic resonance imaging, diffuse excessive high signal intensity (DEHSI), brain, preterm, very low birth weight, developmental disabilities (disturbances).
cellular target in this pathology\textsuperscript{25-29}. According to Volpe\textsuperscript{2}, the diffuse white matter injury may correlate with the cognitive/behavioral deficits in VLBW newborns.

**Case Report**

The authors present an offspring of middle-age parents: 36-year-old mother, 35-year-old father, with no history of chronic disease in either. The mother’s obstetric history was complicated with one stillborn intrauterine pregnancy and one ectopic pregnancy. There were two older offspring developing properly. The boy was born vaginally at the 27\textsuperscript{th} week of pregnancy, with birth weight of 1200 g. The pregnancy was complicated with an infection of the mother’s urinary system, intrauterine infection and placental abruption. The Apgar score was 4, 4, 5. The maturity of the baby according to Ballard’s scale was equivalent to 27\textsuperscript{th} week of pregnancy. Because of severe respiratory distress syndrome observed from the first day of life, he received surfactant and assisted ventilation was applied. Symptoms of generalized infection, progressing with meningitis, thrombocytopenia, significant leukocytosis, and cholestasis (the culture of the blood, cerebrospinal fluid and discharge taken from endotracheal tube was positive for *Staphylococcus epidermidis*) were also recognized. Additionally, the newborn was diagnosed with a congenital heart defect (ventricular septal defect, VSD), necrotizing enterocolitis, retinopathy of prematurity and subclinical hypothyroidism. Ultrasound of the brain performed in the first week of life detected grade 2 intraventricular hemmorhage, which was not present in follow-up examinations in the first month of life.

A complex assessment of his neurological condition was carried out at approximately the 40\textsuperscript{th} week of the baby’s conceptual age, including the assessment of the baby’s position, his spontaneous movement, muscle tonus, and primary reflexes, revealing that his neurological condition was equivalent to corrected age. Despite normal ultrasound of the brain, MRI demonstrated DEHSI on T2-weighted images corresponding to noncystic white matter injury (Fig. 1). Additionally, the baby was diagnosed with severe hearing loss (right ear-100 dB, left ear-80 dB), for which he was treated with hearing aids. Within the first six months of the baby’s life some neurological disturbances occurred. It was observed that the symmetrical tonic neck response and placing reflex continued longer than the physiological norm. These disturbances preceded the symptoms of increased tone in lower limbs and in one upper limb, which were detected after the sixth month of life. The cranial ultrasound carried out before each neurological examination showed mild lateral ventricular enlargement, increasing after the six month of life.

The boy was rehabilitated according to NDT-Bobath’s method since the first months of life and has attended therapeutic sessions with a logopaedics specialist and a psychologist. His hearing loss was treated.

At the end of his 1\textsuperscript{st} corrected year, the boy was diagnosed with cerebral palsy in the form of mild-degree spastic diplegia. After having finished his second year, the boy regained his independent motion. He articulated separate, consciously pronounced words. At the same time, some psycho-motor hyperexcitability was observed.

Another complex assessment of the boy’s neurological condition (including neurological, psychological, auditory brainstem response [ABR] examinations and MRI) was carried out at the end of his third year. The neurological
examinations demonstrated hearing impairment, a slight weakening of the muscle strength in his right upper limb, excessive patellar reflex as well as positive Babinski’s sign in both lower limbs. The gait was independent, but ungraceful. The psychological examination after the third year revealed speech impairment, perceptual organization disorder, neurosensory impairment and irritability and hyperactivity.

Small hypersensitive areas were demonstrated in the white matter of both parietal lobes in MRI on T2-weighted images, which could have been equivalent to earlier anoxic changes (Fig. 2). Additionally, in close proximity to the lateral wall of the lateral right cerebral ventricle, a thin–walled fluid lesion 1.3 cm x 1.5 cm in size was detected. The lesion could be related to the ventricular loculation (Fig. 3).

Other disorders which may contribute to the development of cerebral palsy include PVL, periventricular hemorrhagic infarction, and major intraventricular hemorrhage. These abnormalities are well demonstrated in head ultrasound examinations. However, cerebral palsy is also diagnosed in premature newborns without the sonographic findings described above.

In presented case, ultrasound examination of the boy in question showed only a mild hemorrhage of second degree, which resolved later. No feature related to increased periventricular echogenicity or cystic form of PVL was found. On the other hand, MRI of the brain confirmed noncystic white matter injury. A serial ultrasound examination carried out several months later showed mild widening of the frontal horns of the lateral ventricles, which was a symptom of cerebral atrophy related to the white matter injury.

The ineffectiveness of transfontanel ultrasound examination in the diagnosis of noncystic white matter injury was demonstrated by other authors. According to Volpe, diagnosis of DEHSI can be responsible for developmental disturbances in VLBW newborns. Apart from developmental disturbances, our subject was diagnosed with a mild form of cerebral palsy. In spite of his very complex history (prematurity, general infection with meningitis, intrauterine hypoxia, postnatal hypoxia), the authors

Discussion

Premature newborns are a special group of newborns particularly susceptible to developing spastic diplegia later in life.
conclude that the noncystic white matter injury demonstrated in brain MRI is a sequela of his past history and is responsible not only for the boy’s developmental disturbances, but also for a mild form of cerebral palsy.

To the best of our knowledge, no report on clinical sequelae of noncystic white matter injury has been available until now. Domizio et al.\textsuperscript{32} were the first who reported results of the two-year follow-up in 16 term children diagnosed with DEHSI: half of them had severe and mild mental impairment, and 10 children had severe and mild motor deficits; one child presented with seizures. Their results suggested that DEHSI in preterm newborns is not related to white matter immaturity, but can be considered as a form of the white matter injury.

Taking into consideration the frequency of mild developmental disturbances and DEHSI in VLBW newborns, it is possible that noncystic white matter injury may be responsible for developmental disturbances in this group of children\textsuperscript{3}. Our subject with DEHSI presented developmental disorders and cerebral palsy in his further development. The above presented case also points to a fact that DEHSI diagnosed with MRI may explain the presence of cerebral palsy in newborns in whom cranial ultrasound failed to detect cystic forms of PVL, periventricular hemorrhagic infarction, major intraventricular hemorrhage or hydrocephalus.

We believe that more reports describing the development of newborns with noncystic white matter injury are necessary.

REFERENCES


