Progress in Neonatal Neurology with a Focus on Neuroimaging in the Preterm Infant

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Abstract

There have been tremendous changes in the methods used to evaluate brain injury in the preterm infant in the past 30 years. In particular, major improvements have been made in how we use neuroimaging techniques and now magnetic resonance imaging (MRI) is used more often and considered complimentary to routine and sequential cranial ultrasound. The focus has shifted from severe lesions such as large intraventricular and parenchymal hemorrhages and cystic periventricular leukomalacia to assessing and understanding the etiology of more subtle noncystic white matter injury, punctate hemorrhage, and cerebellar lesions. The more severe lesions that dominated the early period of preterm neonatal brain imaging occur less frequently but are still associated with major disabilities, such as, cerebral palsy, while subtle white matter injury and cerebellar lesions are more often associated with cognitive and behavioral problems, which have become the most prevalent issues among the survivors of extremely preterm birth.

Keywords

► neuro-imaging
► cranial ultrasound
► MRI
► cerebral palsy

Introduction

The first images of the neonatal brain were obtained using computed tomography in the late 70s¹ and there was concern about the high percentage of hemorrhages, usually in the absence of apparent clinical symptoms. Cranial ultrasound (cUS) was introduced into the neonatal intensive care unit only a few years later, initially through the temporal bone using a linear probe, but soon after the anterior fontanelle was used as the main acoustic window. The linear probe was replaced by a mechanical sector probe, which had better resolution and a wider field of insonation. This bedside technique, without ionizing radiation, allowed repeated imaging of the preterm infant and within a few years, we learned that 80% of germinal matrix–intraventricular hemorrhages (GMH–IVH) occurred within the first 72 hours after birth and that the hemorrhage could become more severe over the next day or so and it was sometimes associated with adjacent parenchymal involvement and the development of ventriculomegaly. Recently, we have also become aware that GMH–IVH may already be present at birth or develop beyond 96 hours after birth and this atypical presentation was associated with factor V Leiden mutation in 41% of these infants.² By performing daily examinations, risk factors for hemorrhage were identified, mostly related to complications of mechanical ventilation (pneumothorax, hypercarbia, “fighting the ventilator”) or fluctuations in blood pressure or blood flow. With this knowledge, these risk factors could be minimized and a gradual decrease in the incidence of GMH–IVH was noted. However, antenatal administration of corticosteroids to enhance lung maturation has been the single most important factor for this reduction in GMH–IVH.³

The associated parenchymal involvement now known as a periventricular hemorrhagic infarction (PVHI) adjacent to the ventricle was initially considered to be due to the rupture of the ependymal lining of the ventricle from pressure due to the IVH itself. Most of us now consider a PVHI to be because of the hemorrhage due to impaired venous drainage of the...
mediullary veins in the periventricular white matter. The PVHI tended to be large and globular in shape, the ventricular margin was not maintained allowing communication with the lateral ventricle and usually evolving into a porencephalic cyst. Now the lesions we see are more often smaller and triangular in shape and often are not or only partly communicating with the lateral ventricle and they evolve into one larger or several smaller cysts adjacent to the ventricle (Fig. 1). The size and site of these PVHI lesions as seen on early cUS are predictive of outcome. A PVHI in the frontal white matter carries the lowest risk of developing a unilateral spastic cerebral palsy (USCP) while infants with a PVHI involving the trigone are most at risk. Infants with a PVHI in the temporal lobe are more at risk of cognitive and visual problems. A magnetic resonance imaging (MRI) at term-equivalent age (TEA) allows visual assessment of myelination of the posterior limb of the internal capsule (PLIC) and asymmetry in myelination with poor or absent myelin ipsilateral to the PVHI is highly predictive of the subsequent development of a contralateral USCP. The use of diffusion tensor imaging (DTI) and a direction-encoded color map within 1 month of the PVHI onset, and well before visual assessment of PLIC myelination is possible, has shown asymmetry with delayed PLIC maturation in all who subsequently developed USCP.

**Posthemorrhagic Ventricular Dilatation**

Posthemorrhagic ventricular dilatation (PHVD) develops in approximately 25 to 50% of preterm infants within 7 to 14 days after the onset of a severe GMH–IVH. cUS is a useful bedside technique for following this process (Fig. 2). Measurements can be taken of the ventricular index (VI), anterior horn width and occipital horn width, and these measurements can be used to optimize timing of intervention. When performing cUS before and after a lumbar puncture (LP) that was successful in draining a reasonable amount of cerebrospinal fluid (CSF) (10 mg/kg), one may see a decrease especially in the anterior horn width. Doppler ultrasound can be used to assess changes in cerebral hemodynamics in infants with PHVD, showing an increase in peak systolic flow velocity, followed by a decrease or absence of the
end diastolic flow velocity with increasing intracranial pressure. When LPs are not successful in alleviating the PHVD within a week, the pediatric neurosurgeon may be asked to insert a ventricular reservoir. cUS appearances and measurements will guide the neonatologist in how much CSF to tap from the reservoir to decrease the ventricular size below the 97th centile. There is an ongoing and often heated discussion both in the literature and between neonatologists and neurosurgeons about the optimal time to intervene in PHVD. It is well known that these very preterm infants have a large extracerebral space and it takes several weeks before classical clinical symptoms of increased intracranial pressure (apneas, bradycardia, vomiting, a full fontanelle or rapid increase in head circumference) will occur. Should one wait for clinical symptoms more applicable to the older child or should one treat earlier based on ventricular measurements made with cUS? Those who advocate early intervention are concerned about the adverse effect of intraventricular blood and progressive ventricular dilatation on the adjacent vulnerable periventricular white matter of the preterm infant and have reported better outcome data than others who were less proactive, but these data were retrospective and the need for a randomized controlled trial was acknowledged. In a randomized controlled trial, early (i.e., initiated once the VI line has crossed the 97th centile line) versus late intervention (i.e., initiated after the VI line has exceeded 4 mm above the 97th centile line) is compared (ELVIS trial, number ISRCTN43171322). Because of the decrease in the incidence of severe GMH–IVH, enrollment has been slow and the trial is still ongoing.

**Cerebellar Hemorrhage**

With the increased survival of extremely preterm infants (gestational age [GA], 24–28 weeks) and more routine use of the mastoid window when performing cUS, cerebellar hemorrhages (CBH) are now recognized as a common problem in a very immature infants. The cerebellum has an extremely rapid and complex development during the preterm period. From 24 to 40 weeks gestation, the cerebellar volume, as assessed with in vivo three-dimensional volumetric ultrasound, increases fivefold, and the surface area of the cerebellar cortex increases more than 30-fold during this period.

The reported incidence of CBH when using cUS ranges from 2 to 9% depending on the GA of the population studied. When MRI is performed as well, the incidence is much higher and ranges from 15 to 20%. cUS will only allow recognition of the hemorrhages which are more than 4 to 5 mm in size. Larger CBHs tend to be associated with supratentorial lesions, most often severe GMH–IVH. Smaller (punctate) hemorrhages in the cerebellum are far more common but can only be diagnosed with MRI. Susceptibility-weighted imaging (SWI) further improves the recognition of small punctate cerebellar lesions (Fig. 3). Focal unilateral lesions occurring in the cerebellar hemisphere may originate in the external granular layer, covering the surface of the cerebellum, whereas the less common vermian hemorrhages may originate in residual GMH of the ventricular zone in the roof of the fourth ventricle. CBHs vary from a single to multiple punctate lesions present throughout both cerebellar hemispheres, a single larger hemorrhage in one cerebellar hemisphere, or large bilateral CBH. When the MRI is repeated at TEA, atrophy of the affected cerebellar hemisphere can be seen following larger CBHs (Fig. 3c). A unilateral PVHI can also be associated with a marked loss of contralateral cerebellar volume, so-called crossed cerebellar atrophy. In contrast, Limperopoulos et al showed that unilateral cerebellar lesions had an adverse effect on the contralateral cerebellar volume.

The large CBHs occur within days of birth and as with GMH–IVH, cardiovascular factors appear to be important in the pathogenesis. In the study by Limperopoulos et al, a persistent ductus arteriosus, the minimum pH on day 5 and being born by emergency cesarean delivery were identified as independent risk factors. Ventilation using high-frequency oscillation and the presence of a supratentorial hemorrhage were identified as independent risk factors for punctate cerebellar lesions. While one would expect motor problems such as hypotonia, gait abnormalities, and ataxia, other deficits are more commonly reported in infants with CBH in the absence of severe supratentorial lesions. Among a group of 35 infants with an isolated CBH, impaired expressive language (37%), receptive language (42%), and cognitive deficits (40%), behavioral deficits (34%), and abnormal results on autism screeners measures (37%) were common, and involvement of the vermis almost exclusively accounted for those with socialization difficulties and abnormal autism screening. As MRI is more often used routinely, we will get more insight in how the site and size of the CBHs will affect outcome. Steggerda et al diagnosed small CBHs in 16 of 108 preterm infants did not find an association with neurodevelopmental outcome at 2 years of corrected age. In another study enrolling 131 preterm infants with a cUS diagnosis of a CBH in 3 and an MRI diagnosis in a further 10 infants, there was a fivefold increase in the odds for abnormal neurological examination for those infants with hemorrhages only detected by MRI compared with preterm infants without CBHs and adjusted for GA, presence of associated IVH, and white matter injury (WMI). They did, however, not find an association with the Wechsler Preschool and Primary Scale of Intelligence assessment at the age of 3 to 6 years.

**White Matter Injury**

With higher resolution ultrasound probes and a wider view of insonation, assessment of the white matter has improved considerably although it remains difficult as subtle white matter seen as increased echogenicity with cUS is a very subjective finding. Cystic WMI referred to as cystic periventricular leukomalacia (c-PVL), a term coined in 1962 by Banker and Larroche, “softening” (malacia) of the “white” (leukos) matter is nowadays no longer a common finding and sequential cUS is needed to recognize the cysts which take 2 to 4 weeks to develop. These cysts in the white matter were
first diagnosed with cUS in the early 80s. When severe echogenicity is seen with cUS, an MRI, and especially diffusion-weighted imaging (DWI) performed within 7 to 10 days after the presumed insult may assist in the prediction of cystic evolution. Even though not all with increased signal intensity (SI) on DWI will show cystic evolution, most do in the presence of confluent areas of restricted diffusion. The fluid in these cysts subsequently resorbs with adhesion of the walls of the cysts over the next weeks or months, resulting in the white matter loss and irregular dilatation of the adjacent ventricle. As with PVHI, the extent and especially the site of the cysts are important. Infants with extensive cysts in the parieto-occipital white matter are especially at risk for developing c-PVL. They subsequently reported that these infants with c-PVL and comparable to those with overt WMI, such as infants with IVH, risk factors were more difficult to elucidate for WMI. Hypotension of the boundary zones was considered to be a likely risk factor, but this has not been confirmed and over the last decade or so, it has become clear that WMI is a more multifactorial problem, with (antenatal) inflammation and excitotoxic injury being the key players. The time of onset may be antenatal, but it extends into the neonatal period and some infants may have so-called late-onset c-PVL following an acute deterioration, for example, after developing sepsis or necrotizing enterocolitis. Enterovirus, parechovirus, or rotavirus infection may also result in c-PVL, even though CSF polymerase chain reaction will only be positive for enterovirus and parechovirus but not for rotavirus infection. It is therefore recommended to repeat cUS following any clinical deterioration occurring at any time until discharge home as well as in infants readmitted with a rash, fever, and diarrhea.

A significant decline in the incidence of c-PVL has been noted in several centers over the last couple of decades and this coincided in one, but not in the other study with a decrease in the number of infants who developed CP as well as the severity of their disability, assessed using the gross motor function classification system. In the study by van Haastert et al., the decline was associated with an increased use of antenatal corticosteroids, the antenatal use of antibiotics, birth by cesarean delivery and insertion of an arterial catheter after birth, the latter allowing better control of blood pressure and carbon dioxide levels. It is well known that hypocalcemia is associated with c-PVL, most likely due to vasoconstriction. As a result of this reduction in c-PVL, several studies performing routine MRI at TEA only found a very small number of infants with c-PVL and it seemed therefore more appropriate to talk about WMI, also including subtle WMI, which is much more common in the preterm population. Several studies performed in the last 10 to 15 years showed that the role of cUS was limited when it comes to recognizing milder WMI and MRI is considered to be the “gold standard.” Counsell et al coined the term DEHSI (diffuse excessive high SI) and reported that high SI in the white matter was a common finding in very preterm infants. They were also able to measure apparent diffusion coefficient (ADC) values using DWI and showed that ADC values were significantly higher than in preterm infants without DEHSI and comparable to those with overt WMI, such as infants with c-PVL. They subsequently reported that these infants with increased ADC values in the white matter at TEA had volume reduction in the periventricular white matter, the corona radiate, and within the central region of the centrum semiovale dorsomedial nucleus and also the thalamus and the globus pallidus and these imaging findings were associated with a significantly lower developmental quotient using the Griffiths mental development scale at 24 months of corrected age. Several groups have since then however reported that
they were unable to find an association between DEHSI and neurodevelopmental outcome at 18 to 24 months. More often white matter abnormalities have been graded as mild, or moderate to severe. Woodward et al reported that moderate-to-severe WMI were associated with problems at school age. Others have focused on punctate white matter lesions (PWMLs), seen in approximately 25% of the preterm infants. The presence of inhomogeneous periventricular echogenicity on cUS may suggest the presence of PWMLs, but the sensitivity is quite low. PWMLs are seen as areas of low SI on a T2-weighted sequence and as high SI on a T1-weighted sequence. Because of this, the lesions were initially considered to be hemorrhagic in origin, but with the use of additional MRI sequences it appears that some PWMLs are hemorrhagic and some more ischemic. When the MRI is performed within a week after the presumed insult, ischemic lesions are seen as high SI on DWI. The lesions are often seen in clusters and are even more florid on the T1-weighted sequence performed at TEA, possibly due to early gliosis. Findings from SWI further support the ischemic nature of these lesions, due to the lack of low SI. When low SI is seen on the SWI, this is suggestive of hemorrhage in the lesion. Hemorrhagic PWMLs are more often seen in the presence of GMH–IVH and tend to have a more linear appearance.

**Perinatal Arterial Ischemic Stroke**

While most data about perinatal arterial ischemic stroke (PAIS) only includes full-term infants, PAIS is not uncommon in preterm infants (0.7% in preterm infants with a GA of ≤ 35 weeks). Similar to full-term infants, PAIS was more common on the left (61%), and 7% had bilateral lesions. The majority of strokes involved the middle cerebral artery (MCA) distribution. Twin-to-twin transfusion syndrome, fetal heart rate abnormality, and hypoglycemia were identified as independent risk factors for preterm PAIS. It was of interest that involvement of one or more MCA lenticulostriate branches was common in the preterm infant, referred to as “perforator stroke” by Ecuyer-Goossen et al. In this study, 25 of the 55 infants with perforator stroke were born preterm. Similar to our study, perforator stroke was first diagnosed beyond the first week in 40% of the infants, illustrating the importance of sequential cUS. In a recent study, comparing cUS with an early MRI perforator stroke was more reliably diagnosed with cUS than with MRI.

Bilateral injury to the thalami is unlikely to be because of the bilateral stroke, but it can occur in preterm infants with severe HIE. These abnormalities have only been reported a few times so far. Logitharajah et al only reported the MRI findings. The basal ganglia and brain stem were also often involved and outcome was poor with only one-third of the infants having a normal outcome at the age of 2 years. A third died, and nearly a quarter developed quadriplegic CP. Symmetrical echogenicity in the thalami can also be seen in late-preterm or full-term infants who are considered to have had an acute antenatal insult, even though the history is not always clear. The abnormalities are usually recognized with cUS and become more marked with time (Fig. 4). They tend to present with low Apgar scores, no spontaneous movements, are hypertonic often associated with contractions, sometimes following initial hypotonia, are unable to suck and swallow, and may have facial diplegia. The prognosis is very poor, with lack of psychomotor development and they usually die within weeks or months.

**Future Directions**

In this review, the focus was on the most commonly used neuroimaging techniques, cUS, and conventional MRI. While we tend to perform both cUS and MRI to look for

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**Fig. 4** Male infant, gestational age 34 weeks. (a) Coronal and (b) parasagittal ultrasound scans performed on day 1, showing bilateral thalamic echogenicity. (a) Axial magnetic resonance imaging T1-weighted images in the first week (c) and (d) at term-equivalent age (TEA) show increased signal intensity in the thalami. At TEA (d), the thalami look small and there is no myelination of the posterior limb of the internal capsule.
the presence of abnormalities, we should not forget that sequentially normal cUS and a normal MRI at TEA, especially when combined with a normal neurological examination are also very important in predicting a good outcome and very reassuring for the parents. Abnormalities found on an MRI may however also cause parental concern and some parents would prefer to have an informed choice about performing an MRI and may prefer not to hear the results.56

Other quantitative and more advanced MRI techniques are still mainly used for research but the additional value of DTI, allowing early detection of asymmetry of the corticospinal tracts was mentioned. These advanced techniques are especially promising for a better understanding of the impact of neonatal intensive care on subsequent brain development, particularly promising for a better understanding of the impact of severe intraventricular hemorrhage and therapy for post-hemorrhagic ventricular dilatation. J Pediatri 2008;152(5):648–654

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