Pictorial essay: MRI of the fetal brain

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Abstract

MRI is a useful supplement to USG for the assessment of fetal brain malformations. Superior soft tissue contrast and the ability to depict sulcation and myelination are the strengths of MRI. Subtle or inconclusive USG abnormalities can be confirmed or ruled out by MRI. In some cases, additional findings detected with MRI often help in arriving at a definitive diagnosis, which is necessary for parental counseling and for guiding management. Fast T2W sequences form the basis of fetal MRI. There have been no reports of deleterious effects of MRI on the fetus. A few case examples are presented to illustrate the advantages of MRI.

Key words: Brain, fetal, MRI

Introduction

USG is the primary modality used to assess the fetus. USG examination by a skilled operator, in most cases, provides adequate information regarding fetal morphology, its environment, and its well-being. The quality of USG, however, is adversely affected by factors such as maternal obesity, unfavorable fetal position, multiple gestation, decreased amniotic fluid, or the near-field reverberation artifact (relevant for fetal cranial imaging) [Table 1]. The abnormalities detected on USG may at times be very subtle or inconclusive. In such cases, several studies have shown that MRI is a helpful modality.[1-3] In the context of the fetal brain, some aspects such as maturation and myelination can only be assessed by MRI.[4,5] This essay describes the ideal timing of the MRI examination, safety issues, technique and various indications illustrated and explained by typical examples and cases.

Timing of the MRI study

The need for early prenatal diagnosis, parental anxiety (after a suspicious finding on USG), and local legislation[6] regulating termination of pregnancy are the factors which necessitate early MRI examination of the fetus.

It is relevant to note that the corpus callosum and the vermis are completely formed by the end of 18 weeks of gestation. Fetal brain sulcation begins at around the same gestational age. When deciding on the timing of MRI, one needs to consider the natural history of the fetal disorder in question. For example, in the case of fetal cytomegalovirus infection and tuberous sclerosis, the cranial findings may become apparent only in the third trimester and MRI done too early may do more harm than good. Since most USG abnormalities are generally detected in an anomaly scan done at around 20 weeks of gestation, most cases are referred for MRI around the same time.

Safety issues

Many studies have shown that fetal MRI examination is not associated with any major deleterious effects.[7] No health risks have been reported at field strength of 1.5-Tesla (1.5-T). No adverse outcomes have been observed in pregnant MRI workers.[8] The Safety Committee of the Society for MRI

Table 1: Comparison USG vs. MRI

| Feature                   | US | MRI |
|---------------------------|----|-----|
| Multiplanar               | ✓  | ✓   |
| Field of View             | x  | ✓   |
| Soft Tissue Contrast      | x  | ✓   |
| Obesity                   | x  | ✓   |
| Oligohyramnios            | x  | ✓   |
| Foetal position           | ✓  | ✓   |
| Near half resolution      | x  | ✓   |
| Foetal movements          | ✓  | x   |
| Claustrophobia            | ✓  | x   |
| Cost, availability        | ✓  | x   |
| Spatial Resolution        | ✓  | x   |

✓-has an advantage, x-is at a disadvantage
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has concluded that prenatal MRI is indicated when other nonionizing diagnostic imaging methods are inadequate or when MRI examination can provide important information that would otherwise require the use of ionizing radiation.[9] There is currently no data regarding the level of acoustic noise experienced by the fetus during the MRI procedure.[10,11] The use of gadolinium is not recommended because of the need to avoid potential deleterious effects on the fetus.[12-14] The National Radiation Protection Board and the Food and Drug Administration have approved MRI only after the first trimester.[15,16] The safety of the newer techniques of diffusion-weighted imaging (DWI), diffusion tensor imaging (DTI), MRI spectroscopy (MRS), and functional MRI (F-MRI) has not yet been proven.[17]

**MRI technique**

Informed consent from the mother has to be obtained before the procedure. Fetal MRI is performed on a high-field strength MRI scanner (1.5-T); a phased-array surface coil is placed over the abdomen and pelvis, with the patient in the supine or lateral decubitus position. Polyhydramnios and

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**Figure 1 (A,B):** T2W sagittal MRI image (B) at 27 weeks shows the parieto-occipital sulcus (arrowhead), calcarine sulcus (open arrow), and primary cerebellar fissure (closed arrow). A T2W axial MRI image (B) shows myelination (hypointense region) in the posterior brain stem (arrowhead).

**Figure 2 (A,B):** Single live gestation at 33 weeks. T2W (A) and gradient-echo T2W (T2*) MRI images show intraventricular (arrows) and periventricular (arrowhead) hemorrhage. Fetal blood sampling was negative for TORCH IgMs. A platelet count of 50,000 / mm3 was noted. Pregnancy ended with a stillbirth at 35 weeks.

**Figure 3:** Single live gestation at 30 weeks. T1W axial MRI image at the level of the lateral ventricles shows tubers as subependymal hyperintense nodules (arrows).

**Figure 4 (A-D):** Single live gestation at 37 weeks. USG scans of the heart (A) and brain (B) show a cardiac rhabdomyoma (arrow in A) and mild ventriculomegaly (arrow in B). Axial (C) and coronal (D) T2W MRI images show subependymal tubers (arrowheads), confirming the diagnosis of tuberous sclerosis. The large left lateral ventricular subependymal nodule at the foramen of Monroe is probably responsible for the unilateral ventriculomegaly. The infant developed ash-leaf macules at 1 month and myoclonic jerks at 2 months of age. It died at 3 months (cradle death).
multiple gestations may increase the distance between the region of interest and the surface coil, resulting in a reduction in the signal strength.[18,19] Maternal premedication has been used to achieve fetal sedation, although the fast sequences available on the newer scanners obviate the need for sedation. Maternal breathholding during sequences is desirable.[5,20] Fetal motion significantly degrades image quality. For this reason, fast T2W single-shot imaging techniques such as single-shot fast spin-echo (SSFSE), half-Fourier acquisition turbo spin-echo (HASTE), and rapid acquisition with relaxation enhancement (RARE) have been the most widely used pulse sequences for fetal MRI.[21-24] Such fast acquisition sequences provide good contrast and spatial resolution and are suitable for surface delineation, sulcational analysis, and biometry [Figure 1]. Studies mention a lag of one week of the brain development compared to the neuroanatomic findings.[25] Gradient-echo T2W images (T2*) are accurate in the detection of chronic hemorrhagic lesions and calcifications[8] [Figure 2]. T1W images are well suited for demonstrating small hyperintense lesions like tubers, calcification, lipomas, and laminar necrosis[21,26-29] [Figure 3]. There are reports which mention a good correlation between USG and MRI measurements of biparietal diameter, head circumference and cerebellar width.[30-32] An initial fast localizing sequence is performed; sequences are then acquired in orthogonal planes relative to the immediately preceding plane. Imaging planes are chosen to represent sections relative to the fetus.[5,20]

We normally use the following sequences [Table 2] on our 1.5-T MRI (Avanto, Siemens, Erlangen, Germany) unit:

**Table 2: Sequences**

| Sequence | TR  | TE   | Flip angle | FOV | BW  | Slice thickness | Distance factor | Dimension |
|----------|-----|------|------------|-----|-----|-----------------|-----------------|----------|
| T2 HASTE | 1000| 120  | 144        | 300 | 400 | 2-4 mm          | 10%             | 2D echo  |
| T1 Fl 2D | 120 | 5    | 70         | 370 | 120 | 3-6 mm          | 10%             | 2D echo  |
| T2 Trufi | 4.07| 2.04 | 58         | 350 | 500 | 2-4 mm          | 10%             | 2D echo  |

**Indications and case studies**

MRI is indicated in situations when a brain anomaly has been detected on USG, but the diagnosis is not obvious or definite and needs confirmation.

Marginal ventriculomegaly is one such instance. MRI may detect the presence of heterotopia or sulcational disorders, which have profound prognostic and counseling implications. In the case illustrated in Figure 4, the presence of fetal cardiac rhabdomyoma and unilateral marginal ventriculomegaly prompted the MRI examination. Multiple subependymal tubers were demonstrated on MRI, thus clinching the diagnosis of tuberous sclerosis. Bilateral...
ventriculomegaly, especially of the occipital horns, was the indication for MRI in the case illustrated in Figure 5. Here, MRI clearly demonstrates bilateral, symmetric, neuroparenchymal loss in the parieto-occipital watershed regions as against the USG possibility of clefing; these findings suggest an ischemic etiopathogenesis rather than a neuronal migrational disorder such as schizencephaly. MRI demonstrated the extent, location, symmetry, and morphology of the defects better than USG. In the case illustrated in Figure 6, USG detection of ventriculomegaly, periventricular calcification, cortical thinning, and cerebellar hypoplasia led to a referral for MRI. MRI confirmed the severity of cerebral and cerebellar atrophy but failed to demonstrate calcification. Congenital cytomegalovirus infection was then considered.

Uncertain findings on USG can be supplemented by an MRI examination. In the case described in Figure 7, USG detected inferior vermian agenesis and postaxial polydactyly. Fetal
MRI was done specifically to look for the ‘molar tooth sign’ which, however, was not seen. Postnatally, at 3 months of age, this infant presented with the clinical features of Joubert’s syndrome. A repeat MRI demonstrated the presence of the ‘molar tooth sign,’ thus confirming the diagnosis of Joubert’s syndrome. This is an instance of a disease that evolved with the passage of time.

In Figure 8, cerebellar hypoplasia & suspected cortical atrophy at 31 weeks gestational age were the clues on USG. MRI demonstrated poor sulcation, poor opercularization, and shallow Sylvian fissures, confirming lissencephaly. In some instances, overt intracranial findings on USG may necessitate MRI examination as a second confirmatory tool. The example of a case of USG-detected frontal horn fusion with agenesis of septum pellucidum is illustrated in Figure 9. Here, MRI confirmed the USG findings and also revealed an indistinct optic chiasma and possible hypophyseal hypoplasia. This clinched the diagnosis of septo-optic dysplasia.

MRI can confirm callosal agenesis or dysgenesis in cases suspected by USG. Additionally, MRI may also demonstrate heterotopia. USG-detected fetal intracranial tumors can be characterized by MRI. When USG is unable to differentiate between fetal intracranial bleed and tumor, MRI can help resolve the issue.

Fetal MRI is increasingly being used as a means of surveillance in situations where fetal brain lesions are anticipated. An example of such a situation is the monitoring of the surviving twin after the co-twin’s demise in monochorionic twinning. MRI is also indicated when there is a history of a previous child with a genetic disorder, e.g., Miller-Dieker syndrome.

MRI examination of the fetal brain requires extensive training. As with USG examination, an MRI examination should also be performed in a systematic manner. The USG findings and the history should be known to the radiologist. It is best if the radiologist performing the MRI is provided with the questions that have to be specifically answered. The systematic approach typically includes biometry, maturational / sulcational analysis, and myelination analysis. The ability to accurately measure cerebral and bone biparietal diameters enables quantification of any fetal cerebral neuroparenchymal loss. Callosal length and vermian height and area are measurements that MRI can provide with ease, though with the advent of three-dimensional studies these measurements may also be assessed by USG. A detailed discussion of sulcation and myelination analysis is beyond the scope of this article.

Newer techniques such as DWI, DTI, MRS, and F-MRI are being tried out. DWI may be useful to study hypoxia and ischemia affecting the fetal brain and DTI for studying the development of neural tracts like the corpus callosum, the optic tracts, and the anterior commissure. MRS has the potential to be useful in the evaluation of myelination, ischemia, hemorrhage, metabolic variations in brain damage and inborn errors of metabolism.

Fetal MRI is a powerful supplement to USG and enables us to demonstrate findings that cannot be recognized on USG. Appropriate integration of fetal MRI into the prenatal evaluation algorithm can improve decision-making and patient care. Clinching of a diagnosis enables cross-specialty consultation amongst radiologists, obstetricians, neuropediatricians, and geneticists; this allows more targeted and meaningful counseling and management strategies.

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