INTRODUCTION

Perinatal asphyxia, more appropriately known as hypoxic ischemic encephalopathy (HIE) is characterized by clinical and laboratory evidence of acute or subacute brain injury due to asphyxia. The primary causes of this condition are systemic hypoxia and/or reduced cerebral blood flow (CBF). Birth asphyxia causes an estimated 840,000 deaths or 23% of all neonatal deaths worldwide annually.(1,2,3) Despite major advances in monitoring technology and knowledge of fetal and neonatal pathologies, perinatal asphyxia or more appropriately hypoxic ischemic encephalopathy remains a significant condition that cause mortality and long term morbidity.

MRI is the imaging modality of choice for the diagnosis and follow-up of infants with moderate to severe hypoxic-ischemic encephalopathy. This study was undertaken to evaluate the various MRI patterns of term hypoxic ischemic encephalopathy and to correlate the MRI appearances with clinical outcome. It was also undertaken to compare wherever possible, the neurosonographic appearances with the MRI findings. Differences in the patterns of hypoxic ischemic encephalopathy are likely to result from severity of birth asphyxia. MRI is non-invasive and has no radiation hazards associated with it. The excellent gray–white matter resolution and multiplanar imaging capabilities provide an advantage to MRI over other modalities. The addition of DWI improves the diagnostic capacities of MRI. There is a strong relation between the MRI appearances of birth asphyxia and the clinical outcome. Therefore MRI has a strong role in prognosticating lesions.

MATERIALS AND METHODS

This is a prospective study. A total of 45 hemodynamically stable term neonates with a clinical staging of HIE, referred to the department of Radio-diagnosis for imaging studies, from pediatric department at tertiary care hospital. The study requires that the patients be adequately sedated to tolerate the prolonged examination time inside the MRI machine.

All patients who fulfill the criteria for the study, referred to the department of Radio diagnosis with clinically suspected Hypoxic Ischemic Encephalopathy in a period of 1 year from July 2015 to June 2016 will be subjected for...
the study. Study sample was calculated by power based sample size calculation formula.

Following MRI patients were clinically followed up for a period of five months to assess for neurological outcome. The study exclude preterm neonates with birth asphyxia, haemodynamically unstable patients who may not tolerate prolonged examination times in the isolated setting of MRI.

**MRI Protocol**

The MRI imaging protocol is part of our routine protocol and comprise T2W (axial and coronal), T1W (axial), FLAIR (axial), DWI (axial), GRE (axial), SWI (axial) and MRA (TOF-circle of willis and neck). The total imaging time with 1.5T whole body MR imager was approximately 20 minutes 53 seconds and head coil was used.

**RESULTS**

A total of 45 children underwent for MRI brain. Most of the neonates included in our study belonged to the age group of 6 to 10 days. This study recognizes the limitations imposed because of this selection bias as diffusion findings have been classically known to pseudonormalize after a period of 7 days. 75 % of the patients included in our study were male. The rest were female. 90% of the patients included in our study were normal weight. Only 10% patients were classed as low birth weight.

Out of 45 babies who were part of the study, 35 babies (77.8%) were clinically staged as Stage II, 6 patients were graded Stage I (13.3%) and 4 out of 45 were categorized as Stage III, corresponding to 8.9% of the total study population. The number of subjects with Grade III injuries that who could be included in the study was less due to the unstable clinical picture associated with the same and the difficulties in obtaining MRI in patients on respiratory support.

**Clinical outcome**

Out of the 45 patients included in the study, 21 (46.67%) had abnormal outcomes at 5 months follow up, 18 (40%) had normal outcomes at 5 months and 6 (13.33%) were lost to follow up. This study recognizes the limitations of a five month follow up as many of the neurological sequelae of birth asphyxia are known to manifest in a delayed fashion.

**MRI findings**

Out of 45 patients, 36 patients demonstrated findings positive for HIE. The remaining 9 MRI scans were either normal or demonstrated pathology different from HIE.

Out of these 9 patients, 4 showed presence of subdural haemorrhage, 3 showed evidence of venous sinus thrombosis, 2 patients showed intra-axial hemorrhages.

MRI findings that correlated with HIE in our patients fit into the four categories, i.e. central, periventricular, watershed territory infarcts and mixed findings.

The largest group (17 patients – 37.78%) demonstrated changes in the central structures only. These structures included the thalami, basal ganglia, internal capsules and the corpus callosum.

The next significant group by numbers was the patient population with mixed findings. 14 patients out of 45 (31.1%) had findings that categorized them in the mixed category.

The next group in our study was three study belonged to the watershed infarction corresponding to 6.67% of the study population and two with periventricular changes alone. Two patient in our study belonged to this category, corresponding to 4.44% of the study population.

In a review article titled Prognostic tests in term neonates with Hypoxic Ischemic encephalopathy: A systematic Review(8), the following results were published. In imaging, diffusion weighted MRI performed best on specificity (0.89 [0.62-0.98]) and T1/T2- weighted MRI performed best on sensitivity (0.98 [0.80-1.00]). Magnetic resonance spectroscopy demonstrated a sensitivity of 0.75 (0.26-0.96) with poor specificity (0.58 [0.23-0.87]). The results obtained in our study are in close correlation with the statistical results for MRI published in the review article.

It was also found that neurosonography had a poorer sensitivity compared to MRI as it detected abnormalities in only 44% cases where it was performed as compared to MRI which was abnormal in a much larger percentage of patients. However, the positive predictable value and the specificity are comparable to that of MRI.

**Table - 1 Distribution of Neurosonographic Findings**

| Neurosonogram       | Number | Percent |
|---------------------|--------|---------|
| Abnormal            | 11     | 24.44   |
| Normal              | 14     | 31.11   |
| Not performed       | 20     | 44.45   |
| Total               | 45     | 100     |

**Table - 2 MRI Pattern of Hie**

| MRI pattern        | Number | Percentage |
|--------------------|--------|------------|
| Central            | 17     | 37.78      |
| Mixed              | 14     | 31.11      |
| Watershed infarction | 3   | 6.67       |
| Periventricular    | 2      | 4.44       |
| Other              | 9      | 20         |

**Table - 3 Distribution of Clinical Outcome**

| Outcome             | Number | Percent |
|---------------------|--------|---------|
| Abnormal            | 21     | 46.67   |
| Normal              | 18     | 40      |
| Lost of Followup    | 6      | 13.33   |
| Total               | 45     | 100     |

**Table - 4 Correlation of Restricted Diffusion In Posterior Limb of Internal Capsule With Clinical Outcome**

| Diffusion PLIC | Abnormal outcome | Normal outcome | Lost to follow up | Total |
|----------------|------------------|----------------|-------------------|-------|
| Positive       | 11               | 7              | 2                 | 20    |
DISCUSSION

James Barkovich et al (9) undertook a study to detect changes of neonatal Hypoxic Ischemic Encephalopathy. They evaluated the results for MRI patterns and divided into four groups based on the tissue structures affected.

1) Primarily deep grey matter involvement.
2) Primarily cortical injury.
3) Periventricular pattern of injury.
4) Mixed pattern of injury.

In our study a larger number of patients (20 patients) had restricted diffusion in the posterior limb of the internal capsule as compared to the number of patients (11 patients) who had loss of the normal signal intensity on T1 in the posterior limb of the internal capsule. 55% of patients with loss of the normal signal in the posterior limb of the internal capsule had abnormal outcomes at 5 months compared to 55.55% of PLIC diffusion restriction cases. This is in partial concordance with the observations provided by Rutherford et al (10) in their studies who found that all patients with abnormal signal intensity in the Posterior limb of the Internal capsule had abnormal neuro developmental outcome.

In our study three out of the 45 patients were found to have restricted diffusion in the region of the thalamus and out of these 3 patients, 1 patient also had T1 hyperintensity in the thalamus. All these patients had abnormal neuro developmental outcomes at 5 months follow up.

Seven patients had restricted diffusion within the basal ganglia, of which five (71.4%) had abnormal outcomes and 2 patient (28.5%) had a normal outcome.

10 patients had T1 hyperintensity in the basal ganglia of which 7 (70%) had an abnormal outcome, 2 had a normal outcome (20%) and 1 patient (10%) was lost to follow up. Rutherford et al (11) conducted studies on infants with history of birth asphyxia. They found that outcome was severe in patients with bilateral basal ganglia abnormalities and our study matches with the data obtained in this study.

Out of 45 neonates, 25 (55.56%) neonates showed presence of restricted diffusion within the splenium of the corpus callosum, 10 neonates (22.22%) showed restriction in the genu and 8 neonates (17.78%) showed restriction in the body. All these three groups had comparable outcomes with outcomes from diffusion restriction in the genu and that in the splenium matching more closely.

It was found that neurosonographic findings correlated poorly with clinical outcome as patients in both groups (those with abnormal and normal neurosonograms) had similar outcomes.

Case 1 Mri Findings In Case of Central Hypoxicischemic Encephalopathy

![Restricted diffusion in the genu of the corpus callosum and both internal capsules](image1)

![Diffusion restriction in the corpus callosal splenium](image2)

![ADC image showing reduced ADC values in the genu of the corpus callosum & both internal capsules](image3)
ADC image showing reduced ADC values in the corpus callosal splenium

Loss of normal signal intensity in the posterior limb of internal capsule bilaterally on T1W images

Case 2 Mixed Pattern of Hypoxic Ischemic Encephalopathy in a Neonate

T1 hyperintensity noted involving bilateral basal ganglia

Restriction of diffusion in the splenium of the Corpus Callosum and the posterior limbs of bilateral internal capsules

Restriction of water diffusion in thalami bilateral basal ganglia

Corresponding ADC images showing reduced ADC values

Restricted diffusion in bilateral perirolandic cortices, indicative of cortical involvement
Limitation

The limitation of this study is the lack of systematic follow-up neuro-imaging. Also, patients could be followed up for a maximum of 5 months time at follow-up clinics and it has been recognized that subtle neuro-developmental abnormalities can occur years after birth asphyxia.

CONCLUSION

This study attempted to evaluate the various magnetic resonance imaging changes in hypoxic ischemic encephalopathy in term infants. It set out to categorize and suitably classify the various MRI patterns of disease in these patients. Additionally, an attempt to correlate the MRI findings with neurosonographic appearances was undertaken wherever feasible.

It was found that MRI is superior to other imaging modalities in the evaluation of neonatal hypoxic ischemic encephalopathy. There is a strong and consistent correlation between the various MRI findings and the final clinical outcome.

MRI has a high sensitivity and specificity in the evaluation of the above condition. It is non-invasive and has no radiation hazards. It offers excellent gray-white matter resolution which Computed Tomography is incapable of especially in the neonate with a large proportion of unmyelinated brain parenchyma.

Over study found that diffusion weighted imaging adds sensitivity and provides information not seen on the other conventional sequences. Many of the signs of hypoxic ischemic encephalopathy are based on diffusion weighted imaging.

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