Intraventricular haemorrhage and periventricular leucomalacia: ultrasound and autopsy correlation

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SUMMARY The brains of 30 infants who died after at least one real time ultrasound scan were examined after fixation. The ultrasound diagnosis of either periventricular haemorrhage or periventricular leucomalacia was compared with the macroscopic and histological appearances. Each hemisphere was considered separately for both periventricular haemorrhage and periventricular leucomalacia. The accuracy of ultrasound diagnosis for periventricular haemorrhage was 88%, with sensitivity of 91% and specificity of 85%. The accuracy for periventricular leucomalacia was 90%, with sensitivity of 85% and specificity of 93%. Ultrasound was shown to diagnose the entire range of periventricular leucomalacia lesions. Three hemispheres showed the appearance of prolonged flare, and this correlated with extensive spongiosis and microcalcification of the periventricular white matter, although no macroscopic lesion was seen.

There is now clear evidence that the long term neurodevelopmental prognosis for neonates with intraventricular haemorrhage is significantly better than for those with periventricular leucomalacia (PVL).1-3 If ultrasound imaging is to be effective in predicting prognosis then clearly an accurate differentiation of the pathological entities must be possible on scanning. Previous workers have reported accurate correlation between ultrasound demonstration of intraventricular haemorrhage and its finding on computed tomography or at autopsy,4-9 but in none of these studies was an attempt made to differentiate parenchymal lesions due to ‘extension of haemorrhage’ from PVL. As part of a prospective study of intraventricular haemorrhage and PVL in the preterm neonate we have had the opportunity to correlate ultrasound scan and autopsy findings and we now report our results.

Patients and methods

Over a 16 month period (1 January 1984 to 30 April 1985) 200 neonates of birth weight ≤1500 g admitted to two neonatal intensive care units (Leicester Royal Infirmary and Nottingham City Hospital) had regular cranial ultrasound scans. At least one scan had to have been performed during life for the patient to be included in the study. In both centres an ATL Mark III real time sector scanner fitted with a 3-5, 5, and 7-5 MHz multifrequency scanhead was used. Ultrasound diagnoses based on our previous definitions10 were made before autopsy in all cases. The brain was removed complete at the time of postmortem examination and suspended in 10% formalin for at least one month before sectioning. Coronal cuts were made at 1 cm intervals and the macroscopic appearances noted. Histological examination of the periventricular area was performed in most cases. For the purposes of analysis, each hemisphere was considered separately. The number of true positive (TP), true negative (TN), false positive (FP), and false negative (FN) ultrasound diagnoses for each cerebral hemisphere was tabulated. The accuracy of ultrasound was measured by calculating TP+TN/total number of hemispheres examined. Sensitivity was calculated as TP/(TP+FN) and specificity as TN/(FP+TN).

To show the range of PVL lesions we have included five additional babies who were not included in our previous study.10 Two of these had scans performed soon after death but had no in vivo scan, and a further two preterm infants had birth weights that had excluded them from the previous study. The final infant was born at 29 weeks’ gestation and suffered severe arrhythmias in the first 24 hours of life related to hyperkalaemia. An ultrasound scan at the age of 17 days showed extensive bilateral cystic PVL, involving temporal
and occipital regions, and this was still present when she was last scanned at the age of 7 weeks. She died suddenly aged 6 months and thus illustrates the more long term evolution of PVL.

Results

Forty two of the 200 study babies died, and all deaths occurred before discharge from hospital. Postmortem examination was performed in 31 of the 43 deaths, but in one infant the brain was too soft to fix. The results of the correlation between autopsy and ultrasound examinations are shown in Tables 1 and 2 for the 30 infants with complete postmortem data. Histological examination was performed on 25 brains.

Haemorrhage. Haemorrhage into the germinal matrix or ventricle or extension into the cerebral parenchyma was diagnosed both by ultrasound and subsequently at autopsy in 31 hemispheres. The results are summarised in Table 1. A false positive ultrasound diagnosis of haemorrhage was made in four hemispheres (three infants) in which postmortem examination subsequently showed a normal hemisphere. In three the ultrasound appearance was thought to consist of haemorrhage in the region of the germinal matrix, and the other seemed to show the appearances of more extensive clot in the lateral ventricles. A false negative diagnosis of haemorrhage was made in three hemispheres (two babies) (Table 2). All three hemispheres contained a small germinal matrix haemorrhage that had ruptured into the ventricles. The accuracy of ultrasound diagnosis of haemorrhage was 88%. The sensitivity of diagnosis was 91% and specificity 85%.

Ischaemic lesions. Echodensity in the periventricular white matter detected by ultrasound and thought not to be parenchymal extension of periventricular haemorrhage was seen in 20 hemispheres, and PVL lesions were diagnosed at autopsy in 20 hemispheres (Tables 1 and 2). The accuracy of the ultrasound diagnosis was 90%. A false positive ultrasound diagnosis of abnormal periventricular echodensity was made in three hemispheres (two infants). One child was only scanned once and the other had three scans, and mild bilateral flare was thought to be present on each occasion. A false negative ultrasound diagnosis of PVL was made in three hemispheres (two infants). In two hemispheres there was non-haemorrhagic PVL and in one hemisphere extensive haemorrhagic PVL was found and not previously detected by ultrasound. The sensitivity of ultrasound in the diagnosis of PVL was 85% and specificity 93%.

Table 1 Correlation between ultrasound and postmortem diagnoses for the study group of 60 hemispheres. As each hemisphere may have more than one form of disease the numbers do not add up to 60. Values are No (%)  

<table>
<thead>
<tr>
<th>Ultrasound observations</th>
<th>Postmortem observations</th>
<th>Normal</th>
<th>PVH</th>
<th>PVL</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td></td>
<td>22 (78)</td>
<td>3 (11)</td>
<td>3 (11)</td>
</tr>
<tr>
<td>PVH</td>
<td></td>
<td>4 (11)</td>
<td>31 (89)</td>
<td>0</td>
</tr>
<tr>
<td>PVL</td>
<td></td>
<td>3 (15)</td>
<td>0</td>
<td>17 (85)</td>
</tr>
</tbody>
</table>

PVH=Periventricular haemorrhage; PVL=periventricular leucomalacia.

Table 2 Correlation between postmortem and ultrasound diagnoses for the 60 hemispheres. Values are No (%)  

<table>
<thead>
<tr>
<th>Postmortem observations</th>
<th>Ultrasound observations</th>
<th>Normal</th>
<th>PVH</th>
<th>PVL</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td></td>
<td>22 (76)</td>
<td>5 (17)</td>
<td>2 (7)</td>
</tr>
<tr>
<td>PVH</td>
<td></td>
<td>3 (9)</td>
<td>31 (91)</td>
<td>0</td>
</tr>
<tr>
<td>PVL</td>
<td></td>
<td>3 (15)</td>
<td>0</td>
<td>17 (85)</td>
</tr>
</tbody>
</table>

PVH=Periventricular haemorrhage; PVL=periventricular leucomalacia.

Fig. 1 Spongiosis and microcalcification in an infant with prolonged flare. Haematoxylin and eosin stain, showing generalised spongiosis of periventricular white matter (arrow) and microcalcification (arrow head).
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Range of PVL lesions. To show the variety of PVL lesions an additional five infants have been considered together with the study infants. The ultrasound appearances we have described as prolonged flare occurred in two infants (three hemispheres) whose brains were examined at autopsy. The flares persisted for two weeks and three months, respectively, and were present on the last scan before death. No macroscopic abnormality was seen in either brain, but on histological examination extensive spongiosis and microcalcification was present in the periventricular white matter corresponding to the areas of echodensity (Fig. 1).

At autopsy one infant showed the characteristic 'white spots' described by Banker and Larroche. On ultrasound examination persistent bilateral flare was noted and it was thought that small echofree cavitations subsequently occurred shortly before death. The cysts were not confirmed at autopsy, but the echofree areas may have corresponded to the macroscopic white spots of lipid accumulation.

Seven infants (eight hemispheres) showed the pathological appearance of haemorrhagic infarction at autopsy. These infants died between the ages of 5 hours and 13 days. In all cases the floor of the ventricle appeared intact. In six of these infants in whom an ultrasound diagnosis of precystic PVL was made the necropsy appearance was consistent with haemorrhagic infarction in the region of the echodensity (Fig. 2).

Three infants were found to have periventricular cysts at autopsy. In all cases the initial ultrasound appearance was of a triangular shaped echodensity with its apex towards the angle of the lateral ventricle, from which echofree cavities developed in two of the three cases. A unilateral periventricular cyst was not detected in one baby on scanning,
although a flare had been seen in the corresponding area. This infant was scanned only once during life (Fig. 3).

One infant died aged 6 months of sudden infant death syndrome who at discharge from the neonatal unit showed extensive echofree cavitation in the periventricular white matter of both hemispheres. At autopsy there was irregular, ventricular dilatation, but the posteriorly placed cysts remained only as flattened cavities (Fig. 4), while the anterior cysts could no longer be seen at all.

Discussion

We have shown that real time ultrasound is an extremely good method for diagnosing both intraventricular haemorrhage (accuracy 88%) and PVL (accuracy 90%) in a series of 30 brains. Previous studies have produced similar autopsy correlation with the ultrasound diagnosis of periventricular haemorrhage in the region of 90% accuracy.1–9 In the two infants we report here with a false negative diagnosis of haemorrhage there was an interval of a few hours between the last scan and death, and the bleeding may have occurred during this time. A false positive diagnosis of haemorrhage was made in four hemispheres (three babies), all of whom died within 24 hours of birth. One baby had a massive contralateral parenchymal haemorrhage, and it is possible that on scanning we detected spread of thrombus from this site with no actual focus of bleeding within the ventricle in question. Szymonowicz and her colleagues reported congestion of the germinal matrix region that simulated germinal matrix haemorrhage,9 but this was not present in our infants.

Less information is available on the correlation between the ultrasound diagnosis of PVL and autopsy data. Nwaesey and colleagues reported 78% accuracy between ultrasound demonstration of increased echos or cysts in the periventricular region and the finding of PVL at autopsy in the corresponding site.12 In addition, they made the important point that both haemorrhagic and non-haemorrhagic infarction may produce identical changes on ultrasound. More recently, Fawer et al reported excellent autopsy correlation in five infants with the ultrasound appearances of PVL (three non-cystic and two cystic), and there were no false negative diagnoses.13 In our study ultrasound failed to detect PVL in two babies (three hemispheres). In one hemisphere the changes were early and in the other two there were histological abnormalities only (spongiosis and reactive astrocytosis but no microcalcification). A discrete cyst was missed in one infant whose only scan was performed 48 hours before his death aged 5 days. The scan showed some flaring in the area where the cyst was subsequently observed at postmortem examination. The cyst undoubtedly would have been present at the time of the scan, and we cannot explain why it was not detected, but distortion may possibly have occurred during fixation to give a cystic appearance larger than it may have been in vivo.

Armstrong and Norman reported bleeding into the ischaemic area in 25% of their infants with PVL,14 and we found this in six of 34 (18%) hemispheres. There is, however, considerable controversy concerning the differentiation between parenchymal extension of intraventricular haemorrhage and bleeds into a previously ischaemic lesion. Rushton and colleagues have recently reported eight infants with parenchymal abnormalities on ultrasound and at autopsy all showed PVL.15 These authors proposed that this is the underlying aetiology to all parenchymal haemorrhagic lesions. The evidence for this is still far from conclusive and it is beyond the scope of this study to resolve this question.

We have attempted to differentiate parenchymal extension of intraventricular haemorrhage from PVL by ultrasound scanning and report the relative incidence in a separate paper.10 The autopsy data reported here confirm that all cases of haemorrhagic infarction diagnosed by ultrasound were subse-
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...quenty found to be correct. No infant with parenchymal extension of intraventricular haemorrhage died in the acute stages and correlation in this group was not possible. We are encouraged, however, by the accuracy of ultrasound in diagnosing the range of PVL lesions reported here. This supports the validity of the ultrasound definitions that we have previously reported and used to evaluate prospectively a large group of very low birthweight infants.

We have had the opportunity of observing a variety of appearances in the immature brain that represent the range of perinatal ischaemic injury. The histologic appearances of PVL have been described originally by Virchow in 1867 and later by Banker and Larroche. These appearances may be present without macroscopic abnormality, and we have shown that ultrasound can show both gliosis and the 'white spots' of Banker and Larroche. Haemorrhagic infarction and subsequent cavitation is also readily diagnosed by ultrasound imaging. Two cases have been reported of infants showing extensive cavitation on sequential ultrasound scans in whom the echofree cystic lesions disappeared by 6 months. We report here pathological correlation with this sequence of scan findings. With collapse of the cystic cavities, compensatory ventricular dilatation occurred due to cerebral atrophy. As far as we know this is the only such pathological correlation reported to date.

In conclusion, real time ultrasound scanning seems to be an excellent method for identifying the entire range of PVL lesions. It is of particular interest that apparently normal macroscopic brain appearances are echodense in the presence of microscopic calcification and gliosis. In addition, both the specificity and sensitivity of ultrasound is high for the diagnosis and exclusion of these lesions. We have confirmed that the ultrasound distinction between types of PVL as described elsewhere is a realistic and reliable proposition.

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