Abstracts

Methods unclear. The significance of an enlarged stomach detected on a second trimester scan is unclear: “double bubble” sign and polyhydramnios in the third trimester. The significance of an enlarged stomach detected on a second trimester ultrasound is unclear.

Case 2: The second cases involved a 32 year old multip with confirmed Parvovirus infection who was referred with severe fetal hydrops. Severe thrombocytopenia was again noted however a successful fetal transfusion was performed. Unfortunately the mother subsequently developed Ballantyne (Mirror) syndrome which resolved with expectant management.

Case 3: The final case involved a 28 year old multip with a dichorionic twin pair both of which were severely anaemic with similar haematocrit levels at cordocentesis. Both twins received the same treatment course however different outcomes were encountered.

This case series demonstrates the various complications that add further challenging features to the management of pregnancies affected by Parvovirus infection.

Introduction Duodenal atresia classically presents with a “double bubble” sign and polyhydramnios in the third trimester. The significance of an enlarged stomach detected on a second trimester scan is unclear.

Methods A retrospective review of cases identified from the Wessex Fetal Medicine and Antenataly Detected Anomaly (WANDA) regional databases from 1995 to 2012. Scan reports were reviewed and correlated with outcome.

Results 33 cases of an enlarged stomach in the second trimester were identified. In nine there were additional major anomalies: four with gastrochisis, three with cardiac anomalies (including two trisomies), one severe growth restriction with dilated bowel loops and one with renal cystic dysplasia. In the 24 fetuses without additional anomalies, one with renal cystic dysplasia. In the 24 fetuses without additional anomalies, one with renal cystic dysplasia. In these 19 babies there was one neonatal death following preterm labour at 26 weeks post amenorrhea. In the 18 cases with postnatal follow up, there were no gastro-intestinal anomalies or feeding problems detected.

Conclusions The finding of an isolated enlarged stomach in the second trimester appears to have a good outcome with no associated feeding problems. However if an early “double bubble” sign is seen there is a significant risk of an underlying duodenal atresia.

Introduction Echogenic lung lesions (ELL) are a heterogeneous group of lung abnormalities that display a variety of features and are inherently difficult to diagnose and characterise antenatally. Included in this group are congenital cystic adenomatoid malformations (CCAM), pulmonary sequestration, broncho-pulmonary atresia, and congenital emphysema.

Objectives To investigate the changing incidence of echogenic lung lesions detected both antenatally and postnatally.

Methods A retrospective review of cases identified from the Fetal Medicine database and the Wessex Antenatally Detected Anomalies (WANDA) congenital anomalies register from 1994 to 2011.

Results We identified a total of 111 cases of ELL in 492,559 births during the stated period. All but six cases were identified on antenatal ultrasound.

In 1994, the total incidence of ELL was 0.37 per 10,000 births. By 2011 this had risen to 5.39 per 10,000 births, with a progressive incline during the intervening years.

Conclusion We found a nearly 15-fold increase in the incidence of ELL found antenatally in the Wessex region between 1994 and 2011. It is unclear whether this is due to a true rise in the incidence of this abnormality, or due to increased antenatal recognition as a result of improved ultrasound resolution and operator experience.

REFERENCES

PF.68 TORCH SCREENING, WHERE ARE WE NOW? doi:10.1136/archdischild-2013-303966.078
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TORCH screening is used in pregnancy in a wide number of indications. In 1990 the Public Health Laboratory service advised that individual serology tests, rather than a TORCH screen, should be performed depending upon clinical circumstances1. Our group confirmed these findings for fetal medicine indications. The aim of this retrospective study was to determine our progress with the use of TORCH in pregnancy.

Methods A retrospective review of all TORCH tests requested in St Michael’s Hospital in obstetrics and gynaecology between 01/10/2006 and 31/01/2012 was undertaken via the pathology database.

Results 742 tests were undertaken over the study period. 40 indications were identified. There were 4 positive tests for CMV (1%), with no cases of confirmed toxoplasmosis or rubella. CMV was found in late miscarriage, recurrent miscarriage and multiple fetal abnormalities.

Conclusions The incidence of toxoplasma in the UK is 1-2 infections per 1000 pregnancies and is normally associated with a maternal illness. Rubella is screened for as part of the routine antenatal screen. Our findings have further confirmed the targeted approach to serology screening. We therefore now only perform CMV serology unless there is an overwhelming clinical indication for the addition of toxoplasma testing.

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Introduction Echogenic lung lesions (ELL) are a heterogeneous group of lung abnormalities that display a variety of features and are inherently difficult to diagnose and characterise antenatally. Included in this group are congenital cystic adenomatoid malformations (CCAM), pulmonary sequestration, broncho-pulmonary atresia, and congenital emphysema.

Objectives To investigate the changing incidence of echogenic lung lesions detected both antenatally and postnatally.
Background Anomalies of the fetal central nervous system (CNS) require considerable expertise. In 2010 we launched a Perinatal Neurorsurgical Clinic, wherein fetal CNS anomalies are assessed and counselled by a multi-disciplinary team, comprising a fetal medicine specialist, a paediatric neurosurgeon, a paediatric radiologist and a specialist nurse.

Materials and Methods A prospective review of all patients referred to our clinic from Jan 2010 to July 2012. All cases were triaged initially by a fetal medicine specialist to confirm diagnosis. If warranted, a fetal MRI was obtained prior to referral. Individual patient records were examined to determine pregnancy outcome.

Results Over the study period, 122 fetal CNS anomalies (excluding choroid plexus cysts) were seen. Of these, 41 women (34%) were referred to the Neurosurgical Clinic, including 1 case each of caudal regression syndrome, neuronal migration disorder, sacroccygial teratoma, AV malformation, agenesis corpus callosum, cerebellar hypoplasia and thoracolumbar gibbus (Table). Median gestation at referral was 30.5 weeks. Three women opted for termination. Of the remaining cases, 53% underwent fetal MRI. In 55%, MRI was considered to alter the diagnosis and counselling. 60% of women underwent caesarean delivery, at a median 38.8 weeks. Excluding 3 cases of holoprosencephaly, the perinatal mortality rate in this high risk cohort was 0%.

Conclusions A multi-disciplinary Perinatal Neurosurgical Clinic offers excellent potential, both in expert management of common CNS anomalies, such as neural tube defects, and in the assessment of much rarer fetal CNS anomalies.

Abstract PF.69 Table

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>Total</th>
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<tbody>
<tr>
<td>Ventriculomegaly</td>
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<td>7</td>
</tr>
<tr>
<td>Neural Tube Defect</td>
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<td>18</td>
</tr>
<tr>
<td>Holoprosencephaly</td>
<td>10</td>
<td>3</td>
</tr>
<tr>
<td>Dandy-Walker syndrome</td>
<td>19</td>
<td>4</td>
</tr>
<tr>
<td>Other</td>
<td>9</td>
<td>9</td>
</tr>
<tr>
<td>TOTAL</td>
<td>122</td>
<td>41</td>
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</table>

Abstract PF.71 Table

<table>
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<tr>
<th>No. of cases</th>
<th>Size (mm)</th>
<th>Bilateral</th>
<th>Unilateral</th>
<th>Subsequent scans showed</th>
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<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Progressive in size</td>
</tr>
<tr>
<td>17</td>
<td>10–12</td>
<td>9</td>
<td>8</td>
<td>4</td>
</tr>
<tr>
<td>7</td>
<td>13–15</td>
<td>5</td>
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<td>2</td>
</tr>
<tr>
<td>2</td>
<td>≥16</td>
<td>2</td>
<td>0</td>
<td>2</td>
</tr>
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</table>

(9 babies are undelivered)

Conclusion A multi-disciplinary Perinatal Neurosurgical Clinic offers excellent potential, both in expert management of common CNS anomalies, such as neural tube defects, and in the assessment of much rarer fetal CNS anomalies.

Abstract PF.70

Prenatal Diagnosis, Maintaining High Standards: Reflections on 200 Consecutive Procedures

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Aims To assess the local compliance for CVS and Amniocentesis, with RCOG Green-top standards1. Additionally, examine the indications for prenatal diagnosis, results and outcomes.

Background 1:20 pregnant women in the UK are offered prenatal diagnosis, amounting to 30,000 procedures annually1. CVS is offered 11–13 weeks, Amniocentesis from 15 weeks1. Type and timing of diagnostic test screening provided. Changes are predicted with adoption of combined first trimester screening for Down’s Syndrome1.

Methods Retrospective case-note audit of 200 consecutive prenatal diagnosis cases, performed by operators in Ormskirk District General Hospital, with 3200 deliveries per annum.

Results 200 procedures (25 CVS, 175 Amniocentesis) performed with continuous ultrasound, by 2 experienced operators, from August 2007 to November 2011. Rate of pregnancy loss following any procedure was 1.7% (3 cases of amniocentesis), and 0.6% loss <14 days of procedure (1 Amniocentesis). All miscarriages were <20 weeks gestation. 5 CVS and 1 Amniocentesis required 2 attempts. Bloodstaining complicated one procedure clinically, and 11 cases were recognised by the lab. 20% CVS and 2.9% of amniocentesis had inadequate sample volumes. Overall culture failure was 1%. 2 women reattended with vaginal discharge, 1 with abdominal pain. There was one liquor leak, and one case of sepsis.

Conclusions To assess the aetiology and neurological outcomes of ventriculomegaly at a University Hospital from 2009–2012.

Methods 35 women with ventriculomegaly were identified on anomaly/fetal medicine scans between 20 to 30 weeks gestation. They were investigated for aneuploidy, infections and further abnormalities. Long term neurological outcomes including the differences between mild (10–12 mm), moderate (13–15 mm) and severe (≥16 mm) ventriculomegaly in unilateral/bilaterial/progressive cases were examined.

Results 26 women were offered invasive testing for aneuploidy, it was contra-indicated or not appropriate in 9 patients. 5 women accepted invasive testing, 12 declined and 9 are still considering. Karyotyping was normal in all 5 patients. 31 women had a negative TORCH screen. 15 women had a fetal MRI scan.

Apgar scores were normal in all babies. 7 out of 18 babies with stable or progressive ventriculomegaly had abnormal cranial USS or MRI postnatally. All babies with mild ventriculomegaly at diagnosis have no developmental abnormalities currently. 2/7 babies with moderate ventriculomegaly and 1/2 babies with severe ventriculomegaly have delayed development.

REFERENCE
1. RCOG. Amniocentesis and Chorionic Villus Sampling (Green-top 8). RCOG. 2010.