Abstract 406 Table 1  Structure of discussion with patient’s family

<table>
<thead>
<tr>
<th>Sex of Rearing</th>
<th>Surgical Procedures Needed</th>
<th>Future Hormone Therapy</th>
<th>Fertility</th>
<th>Sexual Function</th>
<th>Urinary Function</th>
<th>Risk of Cancer in Tests</th>
</tr>
</thead>
<tbody>
<tr>
<td>Female</td>
<td>vClitoral reduction</td>
<td>Estrogen during puberty</td>
<td>vInfertile</td>
<td>Intercourse possible</td>
<td>Intact</td>
<td>Nil</td>
</tr>
<tr>
<td>Male</td>
<td>vConstruction of penis, scrotal sac and urethra</td>
<td>May not need testosterone</td>
<td>vSub-fertile</td>
<td>Can conceive with egg donation but with risk as uterus is abnormal</td>
<td>May need to sit for urination</td>
<td>Approximately 10%</td>
</tr>
<tr>
<td></td>
<td>vOrchidopexy Removal of uterus and fallopian tubes</td>
<td>vSub-fertile as male</td>
<td>Sperm banking may be possible</td>
<td>Penile function affected (erection during intercourse)</td>
<td>Sperm banking may be possible</td>
<td>Approximately 10%</td>
</tr>
<tr>
<td>No assignment currently; decision later</td>
<td>vTo decide at 21 years of age vConstruction may be more difficult at later age. Genitalia may resemble male appearance as testosterone is active up to 1 year.</td>
<td>vSub-fertile as male</td>
<td>Infertile as female</td>
<td>Depends on chosen sex</td>
<td>Depends on chosen sex</td>
<td>Approximately 10%</td>
</tr>
</tbody>
</table>

Background 45, X/46, XY mixed gonadal dysgenesis (MGD) is a disorder of sex development characterized by a broad phenotypic spectrum. Patients may have unilateral, bilateral or no testis, streak gonads and/or persistent Mullerian structures. It poses a great clinical challenge due to known effects on growth, hormonal balance and gonadal development.

Objectives We present a case of 45, X/46, XY mixed gonadal dysgenesis (MGD) and the challenges of gender assignment discussion with the patient’s family.

Methods Clinical Case Report

A term new-born infant, delivered at home, presented to the neonatal unit with ambiguous genitalia. Examination revealed a 1.5 cm midline phallic structure, with labial-scrotal folds with rugosity but no urethral opening (figure 1), two external openings at introitus (urethral and vaginal) (figures 2 and 3), and bilateral palpable inguinal gonads, right larger than left.

Results Ultrasound of the pelvis showed a uterus, cervix and vagina with possible right intra-abdominal testis. The neonate passed a synacthen test and had normal 17-Alpha-Hydroxyprogesterone. Gonadotropins (Follicle stimulating hormone and Luteinizing hormone), testosterone and Anti-mullerian hormone were normal but there was no detectable oestradiol. Cytogenetic investigation included FISH and karyotype which showed 45,X[22]/46,XY(idic(Y)(q11)(8).ish idic(Y)(p11.3) (SRYx2), of which 25% of cells were Y-containing, while 75% showed 45,X\[22]/46,X,idic(Y)(q11)\[8].ish idic(Y)(p11.3) (SRYx2), of which 75% of cells were Y-containing, while 25% were X-containing. Exploratory laparoscopy with biopsy of the right gonad showed right gonadal tissue (figure. 4) and left hemi-uterus and fallopian tube with no ovary (figure. 5).

Histology confirmed right testicular tissue and left sided structures resembling fallopian tube showed no ovarian stroma, primordial follicles, nor seminiferous tubules. Findings and options regarding sex of rearing, surgical and medical treatment, were discussed in the family conference. The structure of discussion is appended in table 1.

Conclusions The management of MGD is multi-disciplinary. Gender assignment is based on the consideration of several factors, including external and internal genital findings, the role of surgical procedures required, future prospects of hormone replacement, fertility, urinary & sexual function and risk of gonadal malignancy. In addition, social and psychological support is important as the family makes the decision on gender assignment.

Background Retinopathy of prematurity (ROP) is a potentially vision threatening disease affecting preterm babies. Progress in neonatal intensive care in recent years has led to an increased survival of preterm & sick babies and subsequently, to an increasing incidence of ROP.

Objectives To analyze the incidence, risk factors and outcome of ROP.

Methods

STUDY POPULATION 50 Babies ≤32 weeks gestational age and 50 preterm babies >32 weeks gestational age.

INCLUSION CRITERIA Babies with birth weight ≤1500 g. Babies born at ≤32 weeks of gestation. Selected preterm babies with a birth weight between 1500 grams and 2000 grams or gestational age of more than 32 weeks with additional risk factors (eg. oxygen therapy, sepsis, apnea, birth asphyxia, RDS, NEC, use of surfactant, exchange transfusion, IVH, PRBC transfusion).

Group 1: Babies with gestation ≤32 weeks and/or babies with birth weight ≤1500 g.

Group 2: Selected preterm babies with a birth weight between 1500 grams and 2000 grams or gestational age of more than 32 weeks with additional risk factors as mentioned above.

EXCLUSION CRITERIA Outborn babies treated in our NICU

STUDY PERIOD: 2017-2018 When to screen: First screening examination should be carried out at 31 weeks of gestation or 4 weeks of age, whichever is later.
Abstracts

409 CAN WE REDUCE NEONATAL ADMISSIONS DUE TO JAUNDICE?

Sonal Datta, Gopa Sarkar, UK

10.1136/bmjpo-2021-RCPCH.227

Background Neonatal hyperbilirubinemia is a common cause for neonatal admission in term and preterm infants. These infants are primarily managed on postnatal wards; however, they require admission to neonatal unit due to jaundice above the exchange level, rapidly increasing bilirubin levels, pathological jaundice or sepsis.

Objective The objectives of this project included evaluation of the management of infants admitted to neonatal unit with jaundice including the feeding practices; and identifying the areas of improvement to reduce admissions.

Methods This was a retrospective observational project including infants born at ≥ 35 weeks admitted to neonatal unit at District Hospital with a diagnosis of jaundice from January 1, 2017 to December 2018. Data was collected using proforma, medical records and blood results on computer system.

Results A total of 519 infants ≥ 35 weeks were admitted to the neonatal unit of which 12% (60) infants were admitted due to jaundice. 42% (25) were preterm and 58% (35) were term infants.

The risk factors for jaundice were identified as male infant (66%), first born (49%), gestation, prematurity, and breast fed babies.

The mean birth weight was 2892.5 grams and 20% (12) were low birth weight. 35% (21) infants were admitted from home (average 3.9 days) and 65% (39) from postnatal wards.

27% (16) infants had jaundice < 24 hours. 6 infants had > 10% weight loss on admission. DAT was positive in 23% (14) infants of which 8 infants presented < 24 hours.

Sepsis was suspected in 72% (43) infants but was proven in none.

Only 20% (12) received lactation support on the postnatal ward prior to admission whereas 43% (26) did not receive any support. 36 infants were exclusively breastfed, 13 were formula fed and 11 were mixed-fed. After admission, formula feeds were added to 32 infants while 4 infants exclusively breast fed. 10 infants were given intravenous fluids.

The causes of jaundice included prematurity (25), ABO incompatibility (11), Rh incompatibility (3), poor feeding or exaggerated jaundice (25).

33 infants had bilirubin above exchange line, and required a mean of 24 hours (range 6-144 hours) of phototherapy. None of them required immunoglobulins or exchange transfusion.

The mean length of stay was 3 days (range 1-14 days) and there was a remarkable decrease in breast feeding as only 7% (11.6%) infants were breast fed on discharge.

Conclusions There is a scope to decrease the admissions due to jaundice by optimising the postnatal support on the postnatal wards and community in the presence of risk factors.

Transitional care for late preterm infants is important to reduce admission to the neonatal unit thereby reducing the separation of mother and baby.

The opportunities to support breast feeding on postnatal ward and neonatal unit are often missed.

Effective measures should be taken to promote lactation support at all levels.

410 SYRINGE AIR FLUSH TECHNIQUE ELIMINATES SURFACTANT REFUX AS A LIMITING FACTOR IN USING HIGHER VOLUME SURFACTANT FOR MINIMALLY INVASIVE SURFACTANT THERAPY

Karthikeyan Gengaimuthu, United Arab Emirates

10.1136/bmjpo-2021-RCPCH.228

Background Minimally Invasive Surfactant Therapy or Less Invasive Surfactant Administration (LISA) is the preferred way of administration of surfactant in neonates. Surfactant reflux resulting in ineffective surfactant delivery is considered a limiting factor in administering a higher volume surfactant like bovine surfactant by MIST.

Objectives To analyze if syringe air flush technique as incorporated in our MIST procedure protocol eliminates surfactant reflux with higher dose volume bovine surfactant in our cohort of babies that received surfactant by MIST.

Methods Syringe air flush after the surfactant administration is incorporated in our MIST procedure protocol, the default standard of administering surfactant in our neonatal units.

Both porcine surfactant (1.5 ml/kg) and bovine surfactant