**Normogram for the Frontomaxillary Facial Angle in a Lebanese Population**

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**Objective**
The frontomaxillary facial angle (FMFA) is a first trimester (FT) marker for trisomy. Recently, its role as a FT marker for spina bifida (SB) has been evaluated. Our objective was to determine the normogram for the FMFA in an unselected low risk Lebanese population and to compare our results to the established normogram of Borenstein et al to assess for any ethnic variations precluding its application in our population as an early FT marker of trisomy and SB.

**Methods**
Prospective study on 361 fetuses with confirmed dating undergoing a FT scan at 11 to 14 weeks. All scans were performed by a single sonologist certified by the Fetal Medicine Foundation. The FMFA was measured in all fetuses in a mid sagittal plane according to the guidelines of the Fetal Medicine Foundation. Regression analysis was used to establish the relationship between fetal crown rump length (CRL) and FMFA. ANOVA test of means was employed to compare the mean FMFA in relation to the CRL. P < 0.05 was considered statistically significant. All fetuses were healthy term live births.

**Results**
A total of 361 patients were included in the analysis. Mean CRL was 72.48 mm. Mean FMFA was 77.29°. Regression analysis was employed to establish the relationship of FMFA to CRL and it revealed no significant decrease in the FMFA with advancing gestation (P=0.609) as has previously been described by Borenstein et al. Statistical analysis using ANOVA test of means comparing the mean FMFA with CRL indicated that the mean FMFA is significantly unchanged as CRL increases or decreases. For a CRL of 50-59, 60-69, 70-79 and 80-89 mm, the mean FMFA was 79.3°, 77°, 77°, 78.1° and 77.3° respectively (p=0.421). The FMFA was > 85° in 7.8% of our patients.

<table>
<thead>
<tr>
<th>CRL mm</th>
<th>Total Cases</th>
<th>FMF Mean±SD</th>
<th>Min</th>
<th>Max</th>
<th>patients (%) with FMFA &gt;85</th>
</tr>
</thead>
<tbody>
<tr>
<td>[50-59]</td>
<td>20</td>
<td>79.3±1.1427</td>
<td>68.00</td>
<td>89.00</td>
<td>1 (5.0%)</td>
</tr>
<tr>
<td>[60-69]</td>
<td>93</td>
<td>77.0±6.989</td>
<td>57.00</td>
<td>94.00</td>
<td>5 (5.4%)</td>
</tr>
<tr>
<td>[70-79]</td>
<td>203</td>
<td>77.0±4.946</td>
<td>54.00</td>
<td>97.00</td>
<td>16 (7.9%)</td>
</tr>
<tr>
<td>[80-89]</td>
<td>45</td>
<td>78.1±1.0247</td>
<td>64.00</td>
<td>89.00</td>
<td>6 (13.3%)</td>
</tr>
<tr>
<td>Total</td>
<td>361</td>
<td>77.3±3.608</td>
<td>54.00</td>
<td>97.00</td>
<td>28 (7.8%)</td>
</tr>
</tbody>
</table>

**Conclusion**
Our study demonstrates that in an unselected low risk Lebanese population, there is no statistically significant decrease in the FMFA with advancing gestation as has been previously reported by Borenstein et al. In 7.8% of our population, the FMFA is > 85°. This ethnic variation calls for employing caution when using the FMFA in our FT screening for trisomy and when screening fetuses who may be at risk for SB.
Can the Fetal NT Help Predict the Evolution of Hypoplastic Left Heart Syndrome Suspected at 11-14 Weeks?

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Objective
To assess the evolution of hypoplastic left heart syndrome (HLHS) suspected at 11-14 weeks, and to evaluate the impact of the NT on predicting outcome.

Methods
Retrospective study comparing the findings and natural progression of 5 cases of HLHS suspected at 11-14 weeks. All scans were performed by a single sonologist certified by the Fetal Medicine Foundation. Fetuses suspected of having HLHS were referred to pediatric cardiology. The 5 cases were compared with respect to NT, presence of other structural/cardiac findings, karyotype (when available), final diagnosis and outcome. Chi square test was used in the analysis. P < 0.05 was considered statistically significant.

Results
There were 5 fetuses confirmed of having HLHS. Of those 2 (40%) elected termination at 15w0d and 21w6d, and 2 (40%) had spontaneous in utero demise at 14w2d and 18w0d. All 4 had an NT > 3 mm. The 5th case had a normal NT of 2.2 mm and evaluation at 2 week intervals demonstrated a decrease in the atrioventricular disproportion (AVD), a persistent left superior vena cava (LSVC) and a suspected dilated coronary sinus versus an atrioventricular (AV) canal. The fetus was delivered at term and the final diagnosis was an AV canal, persisting LSVC and a dysplastic mitral valve. The NT was highly predictive of the outcome in the 5 cases: the higher the NT the higher the chance for in utero demise. In the one case with a normal NT (20%), there was normalization of the AVD and a favorable neonatal outcome. Even though our sample size was very small, having an NT < 3 mm was associated with a favorable outcome (P=0.025).

Conclusion
In this small cohort of patients, our study emphasizes the varied outcome of HLHS suspected at 11-14 weeks with 40% ending with spontaneous in utero demise and 20% having a favorable outcome. As has been previously shown, the larger the NT, the worse the fetal prognosis and the higher the chances of spontaneous demise. This may lessen the burden on those families electing to terminate. Whenever HLHS is encountered with a normal NT, caution must be exercised without haste in offering termination.

<table>
<thead>
<tr>
<th>Age</th>
<th>G</th>
<th>P</th>
<th>GA</th>
<th>NT</th>
<th>NB</th>
<th>4CV</th>
<th>TR</th>
<th>Karyotype</th>
<th>Other Findings</th>
<th>Final Diagnosis</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>19</td>
<td>4</td>
<td>1021</td>
<td>14w2d</td>
<td>3.1</td>
<td>+</td>
<td>HLH</td>
<td>abn</td>
<td>n/a</td>
<td>Megacystis, limb amputation</td>
<td>Multiple anomalies with HLHS</td>
</tr>
<tr>
<td>2</td>
<td>34</td>
<td>5</td>
<td>3103</td>
<td>13w1d</td>
<td>4.2</td>
<td>+</td>
<td>HLH</td>
<td>nl</td>
<td>n/a</td>
<td>-</td>
<td>HLHS</td>
</tr>
<tr>
<td>3</td>
<td>37</td>
<td>4</td>
<td>3003</td>
<td>12w6d</td>
<td>7.9</td>
<td>+</td>
<td>HLH</td>
<td>abn</td>
<td>n/a</td>
<td>-</td>
<td>HLHS</td>
</tr>
<tr>
<td>4</td>
<td>22</td>
<td>2</td>
<td>1001</td>
<td>13w3d</td>
<td>8</td>
<td>+</td>
<td>HLH</td>
<td>nl</td>
<td>45XO</td>
<td>Cystic Hygroma</td>
<td>Turner Syndrome with HLHS</td>
</tr>
<tr>
<td>5</td>
<td>18</td>
<td>1</td>
<td>13w0d</td>
<td>2.2</td>
<td>+</td>
<td>HLH</td>
<td>nl</td>
<td>n/a</td>
<td>NI at 21w3d and 24 LSVC</td>
<td>AV Canal, Dysplastic MV, LSVC</td>
<td>Alive at 6 months</td>
</tr>
</tbody>
</table>
Objective

Our study aims at investigating the spatial relationships between 8 anatomic planes in the 11+6 - 13+6 week fetus.

Methods

This is a retrospective pilot study where three- and four-dimensional stored data sets were manipulated to retrieve 8 anatomic planes starting from the mid-sagittal plane of the fetus. Standardization of volumes was performed at the level of the transverse abdominal circumference plane. Parallel shift was utilized and the spatial relationships between 8 anatomic planes were established. The median and the range were calculated for each of the planes, and they were evaluated as a function of the fetal crown rump length. P < 0.05 was considered significant.

Results

A total of 63 volume data sets were analyzed. The 8 anatomic planes were found to adhere to normal distribution curves, and most of the planes were in a definable relationship to each other with statistically significant correlations.

Conclusion

To our knowledge, this is the first study to describe the possible spatial relationships between 8 two-dimensional anatomic planes in the 11+6 - 13+6 week fetus, utilizing a standardized approach. Defining these spatial relationships may serve as the first step for the potential future development of automation software for fetal anatomic assessment at 11+6 - 13+6 weeks.
Is the Size of the Fetal Renal Pelvis Gender-Dependent at 20-24 Weeks?

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Objective
Clinically we have noted that male fetuses tend to have a more prominent renal pelvis than female fetuses at 20-24 weeks. For this reason, this study was conducted with the aim of comparing the antero-posterior diameter of the renal pelvis in male fetuses versus female fetuses at 20-24 weeks of gestation given the role of pyelectasis as a second trimester marker for trisomy 21.

Methods
Prospective study on 203 euploid fetuses who had confirmed dating and who underwent first trimester scanning between 11 and 14 weeks. A second trimester scan was done at 20-24 weeks. When prominent, the antero-posterior diameter of both renal pelvises was measured in all fetuses in an axial plane. All scans were performed by a single sonologist certified by the Fetal Medicine Foundation. All fetuses had a known outcome and were healthy term live births. We excluded twin gestations, anomalous fetuses, fetuses born at less than 37 weeks, and those who were lost to follow up beyond the first trimester scan.

T-test and chi square were used for the statistical analysis. P < 0.05 was considered statistically significant.

Results
Using the paired test in comparing the right to left pelvises in all fetuses, no statistical difference was found. As a result, the average of each fetus’ renal pelvises was used in the analysis. There was a statistically significant difference between the mean female pelvis of 3.41 mm and the mean male pelvis of 3.84 mm (p=0.007). Looking at fetuses with an average renal pelvis of ≥ 6 mm, there was a statistically significant difference between the males and females. In 8% of males and 1.3% of females, the renal pelvis measured ≥ 6 mm (p=0.04).

Conclusion
In a representative sample of our population under study, the renal pelvis in the euploid male fetus is larger than that in the euploid female fetus. As a result, it may be worthwhile to establish gender specific normograms and gender specific cutoffs for pyelectasis when incorporating it into the second trimester risk assessment of trisomy 21.