An unusual cause of primary amenorrhoea suggested by the urine steroid profile

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The authors have nothing to disclose

Introduction
Primary amenorrhoea is a pathology with many possible aetiologies. The timing and appropriate first-line investigations for delayed menarche in an otherwise well teenage with secondary sexual characteristics are debatable.

Case report

Presentation
A 15-year-old female of African origin was referred to the regional paediatric endocrine clinic due to: Parental concern with incidental finding of “large clitoris”.

History
• The patient’s mother noticed the patient’s clitoris “looked big” incidentally when she was shown a non-specific rash on the legs without clothes on
• Pubertal development
  - Pubic hair since aged 10 years
  - Breast development since aged 13 years
  - Not achieved menarche
• Past medical history: None apart from acne
• Family history of delayed menarche (mother achieved menarche at aged 17 years)

Examination
• Clinically well and “slim-built”
• Facial acne
• Auxology: Height 172 cm (SDS 1.5), weight 60 kg (SDS 0.41) BMI 20.1 kg/m² (SDS -0.26)
• Puberty: Tanner stage 4 (B4,P4)
• Clitoris normal size and anatomy
• Rest of general examination unremarkable, in particular, no evidence of hepatosplenomegaly or masses on abdominal palpation

Initial impression
• Signs of hyperandrogenisation: acne and primary amenorrhoea
• To rule out polycystic ovarian syndrome and late onset congenital adrenal

Initial investigations
Biochemistry
- Na, K, Cl, HCO3, U&E, TFT, normal range
- LH 6.2 IU/L
- FSH 8.7 IU/L
- Oestradiol 234 pmol/L
- Testosterone 5.3 nmol/L [0.24-2.71]
- SHBG 87 nmol/L [20-110]
- T/SHBG ratio 7.9 [0-4]
- Prolactin 171 mU/L [102-496]
- Androstenedione 5.4 nmol/L [2-54]
- DHEA-S 1 umol/L [1.6-7.8]
- 17-OHP 1.8 nmol/L *

Pelvic Ultrasound Scan
Small adult uterus with endometrium, left ovary 4.6 ml, right ovary 2.5ml

Urine 24 hour steroid profile (figure 1 and table 1) showed unexpected findings suggestive of liver abnormality
Report: “High androstenediol and a-cortolone and low tetrahydrocortisol. Pattern found in association with portal hypertension”.

Further Investigations
Repeat urine steroid profiles
Due to the unexpected results from the initial urine steroid profile, the profiles were repeated pre and post dexamethasone suppression (Table 1).

Table 1: Results of initial and repeat urine steroid profile outcomes
<table>
<thead>
<tr>
<th>Steroid</th>
<th>Initial Urine</th>
<th>Post dexamethasone</th>
<th>Ratio</th>
<th>Normal</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Androstenediol metabolites</td>
<td>631</td>
<td>326</td>
<td>34</td>
<td>1054</td>
<td>703</td>
</tr>
<tr>
<td>Cortisol metabolites</td>
<td>3716</td>
<td>1953</td>
<td>&lt;30</td>
<td>4899</td>
<td>1076</td>
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<tr>
<td>Androstenediol</td>
<td>2548</td>
<td>1978</td>
<td>&gt;34</td>
<td>187</td>
<td>63</td>
</tr>
<tr>
<td>Androstenediol/cortisol metabolites</td>
<td>71</td>
<td>101</td>
<td>2</td>
<td></td>
<td></td>
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<tr>
<td>a-cortolone</td>
<td>1592</td>
<td>875</td>
<td>17</td>
<td>738</td>
<td>163</td>
</tr>
<tr>
<td>a-cortolone/cortisol metabolites</td>
<td>42</td>
<td>45</td>
<td>15</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Liver investigations
Liver profile: ALT 239 U/L (0-40), ALP 500 U/L (30-130), GGT 235 U/L (0-38), Alb 26 g/L (35-50), INR 1.5 (0.8-1.1), APTR 1.31 (0.85-1.35)
US liver: Parenchyma heterogeneous, right and left duct dilated, portal vein patent with normal flow. Enlarges spleen 13.8cm

Diagnosis
• Primary amenorrhoea due to previously undiagnosed liver disease

Progress
• Referred to regional liver centre
• Diagnosis of autoimmune liver disease confirmed by antibodies testing and liver biopsy
• Liver function improved since she was started on prednisolone and Azathioprine
• Patient achieved menarche with regular bleeds 6 month’s later

Discussion
The initial investigations were guided by the patient’s clinical features. The diagnosis was suggested by the urine steroid profile requested to exclude adrenal pathology in the absence of clinical stigmata of chronic liver disease.

The relative increase of a-cortolone in the urine steroid profile has previously been demonstrated in cirrhosis, but raised androstenediol has not been previously reported (Figure 2). There was also a low ratio of cortisone (11-ox) to cortisol (11-hydroxy) metabolites, which is not an invariant feature, but is likely to reflect impairment of 11-hydroxysteroid dehydrogenase-1 in hepatocytes.

Although primary amenorrhoea is often physiological in teenage girls, it may also be the first presenting sign of an underlying chronic disease. Careful evaluation is always needed to avoid missing an alternative diagnosis.

Figure 1: Urine steroid profile

Figure 2: Urinary steroid metabolites