## **Adolescent With Suspected Hirsutism**

| Suggestive history and physical findings   | Initial laboratory and/or radiologic work-up can include:   | When to refer  | Items useful for consultation  | Additional information  |
|--|---|--|--|---|
| <ul> <li>Symptoms/signs:         <ul> <li>Excessive terminal hair growth in locations typically seen in adult males (face, sternum, lower abdomen, back, and thighs)</li> <li>Can be associated with other signs associated with androgen excess, such as acne and irregular periods</li> <li>History of premature adrenarche may be present</li> </ul> </li> <li>Family history:         <ul> <li>Family history of hirsutism and/or polycystic ovarian syndrome (PCOS) may be present</li> </ul> </li> <li>Differential Diagnosis</li> </ul> | <ul> <li>Blood tests:         <ul> <li>Total and free testosterone (assay for women and children)</li> <li>DHEAS</li> <li>Androstenedione</li> <li>17 OH progesterone</li> </ul> </li> <li>TSH</li> <li>Radiologic studies:         <ul> <li>Pelvic ultrasound for very elevated testosterone levels</li> </ul> </li> <li>Other tests to consider after consultation with Pediatric Endocrinologist:         <ul> <li>Prolactin</li> <li>ACTH stimulation test for androgens</li> </ul> </li> </ul> | Urgent: Concern for tumor:  Total testosterone >200 ng/dl DHEAS >700 mcg/dl  Concern for non classic CAH: Elevated 17 OH progesterone  Routine: Laboratory findings at or just above the normal ranges  Find a Pediatric Endocrinologist | Previous growth data/growth charts  Pertinent medical records  Recent laboratory studies | Polycystic Ovarian Syndrome: A Guide for Families  References |

## **Differential diagnosis of Hirsutism:**

- Physiologic hyperandrogenism of puberty
- Idiopathic hyperandrogenism
- PCOS
  - Less commonly:
- Congenital Adrenal Hyperplasia (CAH): late onset CAH, mild CAH, non classic/virilizing CAH
- Androgen secreting tumors of the adrenal glands or ovaries
- Hypothyroidism
- Cushing's disease
- Severe hyperprolactinemia
- Hypertrichosis
- Exposure to androgenic drugs

## **Additional Information:**

- Hirsutism affects 5–10% of reproductive-aged females
- Diagnosis of hyperandrogenism can be based on clinical symptoms or measurement of serum androgens. In females, androgens originate from three primary sources: (1) the ovarian theca, (2) the adrenal cortex, and (3) within end organs by peripheral conversion.

The American College of Obstetricians and Gynecologists makes the following recommendations and conclusions:

- Pelvic ultrasonography is not routinely indicated unless serum androgen levels or the degree of virilization is concerning for an ovarian tumor.
- Multimodal therapy is the most effective approach to the treatment of hirsutism; this includes lifestyle changes, physical hair removal, and androgen suppression or blockade with medication that slows or prevents new hair growth.
- If hormonal therapy is initiated, patients should be counseled that it may take >6 months before they see the benefits of treatment.
- Patients should be assessed at routine intervals (every 3–6 months) for adverse effects and response to treatment until their condition is stable; they then should be monitored annually.
- Monitoring serum androgens is not recommended.

## **Suggested References and Additional Reading:**

 $\underline{https://www.acog.org/Clinical-Guidance-and-Publications/Committee-Opinions/Committee-on-Adolescent-Health-Care/Screening-and-Management-of-the-Hyperandrogenic-Adolescent? \underline{IsMobileSet=false}$ 

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