

# Child With Suspected Sexual Precocity

Suggestive history and physical findings	Initial laboratory and/or radiologic work-up can include:	When to refer	Items useful for consultation	Additional information
<p><u>Females before 8 yrs of age:</u>  <b>Estrogenic:</b>            Breast development, vaginal discharge/bleeding  <b>Adrenergic:</b>            Pubic hair, axillary hair, body odor, acne</p> <p><u>Males before 9 yrs of age:</u>            Testicular enlargement, pubic hair, axillary hair, body odor, acne</p> <p>For both sexes:            Growth acceleration, exposure to exogenous hormones, history of brain injury or radiation, family history of early puberty, midline defects, severe hypothyroidism, café au lait macules, FH/history of neurofibromatosis</p> <p><u>Differential Diagnosis</u></p>	<p><b><u>Radiological tests:</u></b></p> <ul style="list-style-type: none"> <li>• Bone age</li> </ul> <p><b><u>Blood tests:*</u></b></p> <ul style="list-style-type: none"> <li>• LH</li> <li>• FSH</li> <li>• testosterone (males)</li> <li>• estradiol (females)</li> <li>• DHEAS</li> <li>• TSH</li> <li>• Free T4</li> </ul> <p>*Pubertal laboratory tests should be obtained in the <u>early AM</u> using <u>sensitive pediatric assays only</u></p> <p><u>Other tests to consider after consultation with Pediatric Endocrinologist:</u></p> <ul style="list-style-type: none"> <li>• 17-OH progesterone</li> <li>• androstenedione</li> <li>• Pelvic ultrasound</li> <li>• Brain MRI</li> </ul>	<p><b><u>Urgent:</u></b>            CNS abnormality such as headaches, seizures or visual changes, very elevated 17-OH progesterone; rapidly progressing puberty, very young age, initial presentation is vaginal bleeding</p> <p>Bone age greater than 2 SD above chronologic age; signs of rapid virilization</p> <p><b><u>Routine:</u></b>            Normal bone age, normal labs</p> <p><u><a href="#">Find a Pediatric Endocrinologist</a></u></p>	<p>Previous growth data/growth charts</p> <p>Pertinent medical records</p> <p>Recent laboratory and radiologic studies (including actual copy of bone age)</p>	<p><u><a href="#">Precocious Puberty: A Guide for Families</a></u></p> <p><u><a href="#">Premature Adrenarche: A Guide for Families</a></u></p> <p><u><a href="#">Premature Thelarche: A Guide for Families</a></u></p> <p><u><a href="#">References</a></u></p>

## Differential diagnosis of sexual precocity

### Gonadotropin-dependent sexual precocity

- Idiopathic (80% of cases in females, 10% of cases in males)
- CNS lesion
  - Hypothalamic Hamartoma
  - Other CNS tumors or lesions
- Genetic abnormality
  - Gain-of-function mutation of the G-protein-coupled kisspeptin-1 receptor (GPR54)
- Sex-steroid exposure causing bone age advancement

### Gonadotropin-independent sexual precocity

- Normal variants
  - Premature thelarche
  - Premature adrenarche
- Neuroendocrine
  - LH/hCG excess
  - Hypothyroidism
- Adrenal
  - Congenital adrenal hyperplasia
  - Adrenal tumor
- Gonadal
  - McCune-Albright Syndrome
  - Leydig cell tumor
  - Ovarian tumor
  - Familial male-limited precocious puberty (testotoxicosis)

**Suggested Reference and Additional Reading:**

Gad B. Kletter, Karen O. Klein, Yolanda Y. Wong . A pediatrician's guide to central precocious puberty. *Clinical Pediatrics* 2015 May; 54(5):414-24

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