



Definitions

- **Short stature:** height that is below the 3rd percentile
- **Familial short stature:** parents with short stature; normal growth velocity, proportionally normal weight, normal timing of pubertal onset, and normal bone age; predicted final height based on current growth pattern within expected range of mid-parental height
- **Constitutional delay of growth and puberty:** parents without short stature; normal pre-pubertal growth velocity, proportionally normal weight, late pubertal onset, delayed bone age, similar family history in one or both parents; should achieve normal final height without intervention

Measuring Stature

- 0 - 24 months = measure **length** (recumbent)
 - Calibrated length board, no shoes, eyes up, legs extended, toes upwards
- 2+ years = measure **height** (standing)
 - Wall-mounted stadiometer, no shoes, legs straight, looking straight ahead
- Normal growth in stature:
 - 0 - 24 months = rapid growth
 - pre-pubertal = slower consistent growth ~5 cm/year
 - puberty = rapid growth

History

- **Growth history** → onset of short stature, if child has always been short or recent growth velocity decrease, if weight is not proportional to height
- **Birth history** → gestation, birth weight and length, IUGR, neonatal hypoglycemia, jaundice, maternal illness, smoking
- **Developmental history** → developmental milestones, tooth eruption and loss, puberty onset
- **Nutrition** → calorie intake, feeding history
- **Social history** → indications of deprivation
- **Medications affecting growth** → stimulants, corticosteroids
- **Mid-parental height** using child's sex assigned at birth and biological parents' heights

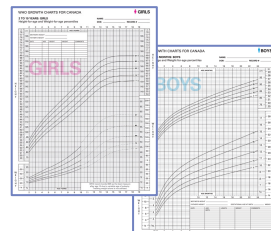
$$\text{male} = \frac{\text{mother} + \text{father} + 13 \text{ cm}}{2}$$

$$\text{female} = \frac{\text{mother} + \text{father} - 13 \text{ cm}}{2}$$
- **Review of symptoms** → GI, cardiac, renal, thyroid diseases; neurological symptoms suspicious of tumours including headaches, vision changes



Decreased growth velocity is **more concerning** than a short absolute height.

Growth should be plotted on the **WHO Growth Charts for Canada**. For children with genetic syndromes, growth should be plotted on **syndrome-specific growth charts**.



Physical

- Full physical exam including cardiac, respiratory, GI, extraintestinal manifestations of IBD including eyes
- Plot height and growth velocity **every 6 months**
- Upper-to-lower segment ratio and arm span to evaluate for skeletal dysplasia
- Sexual maturity rating (Tanner stages)
- Features associated with genetic syndromes including Down syndrome, Turner syndrome, Noonan syndrome

Differential Diagnosis (With Examples)

Normal variants

- Familial short stature
- Constitutional delay of growth and puberty

Endocrine

- Growth hormone deficiency
- Hypothyroidism
- Cushing's syndrome

Malabsorption

- Celiac disease
- Inflammatory bowel disease

Chromosomal abnormalities

- Skeletal dysplasia
- Syndromes associated with short stature

Potential Investigations

- Bone age X-ray
- No investigations is appropriate, given clinical context
- IGF-1, IGFBP-3, GH
- T4, TSH
- Cortisol, ACTH
- LH, FSH, testosterone, estradiol
- Glucose
- tTG-IgA
- CRP, ESR, fecal calprotectin
- Endoscopy, MR enterography
- Genetic tests (molecular, chromosomal)

October 2023

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