

# Growth and Growth Disorders in Children

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# Outline

- Normal Growth and puberty
- Causes of short stature
- Approach to a short child

# Growth

- Growth in childhood: is a complex and dynamic biological processes, is tightly controlled and regulated.

Short stature: height below 3<sup>rd</sup> percentile (depending on what chart you use it could be < 5<sup>th</sup> percentile)  
Tall stature: > 95<sup>th</sup> percentile

Normal (full-term) birth weight: 2.4 kg - 4.2 kg (SGA: TORCH inf., eclampsia)

HC when born: 34 cm at 6 mon: 43 cm at 1 year: 49 cm

Baby should reach double their birth weight at 4 mon and triple by 1 yr

A newborn starts at 50 cm. They should reach double their height at 4 yrs old. Half of that growth (25 cm) should be completed in the 1st year of life.

Preemies catch up: weight: by 2 yrs  
height: by 2-4 yrs  
HC: by 1-2 yrs

# Growth Parameters

- Weight for age
- Height (length) for age
- Weight for length
- Body mass index for age
- Head circumference for age

# Stadiometer

Height  
↓  
Can  
stand



length - measured  
laying down

# Measuring Table

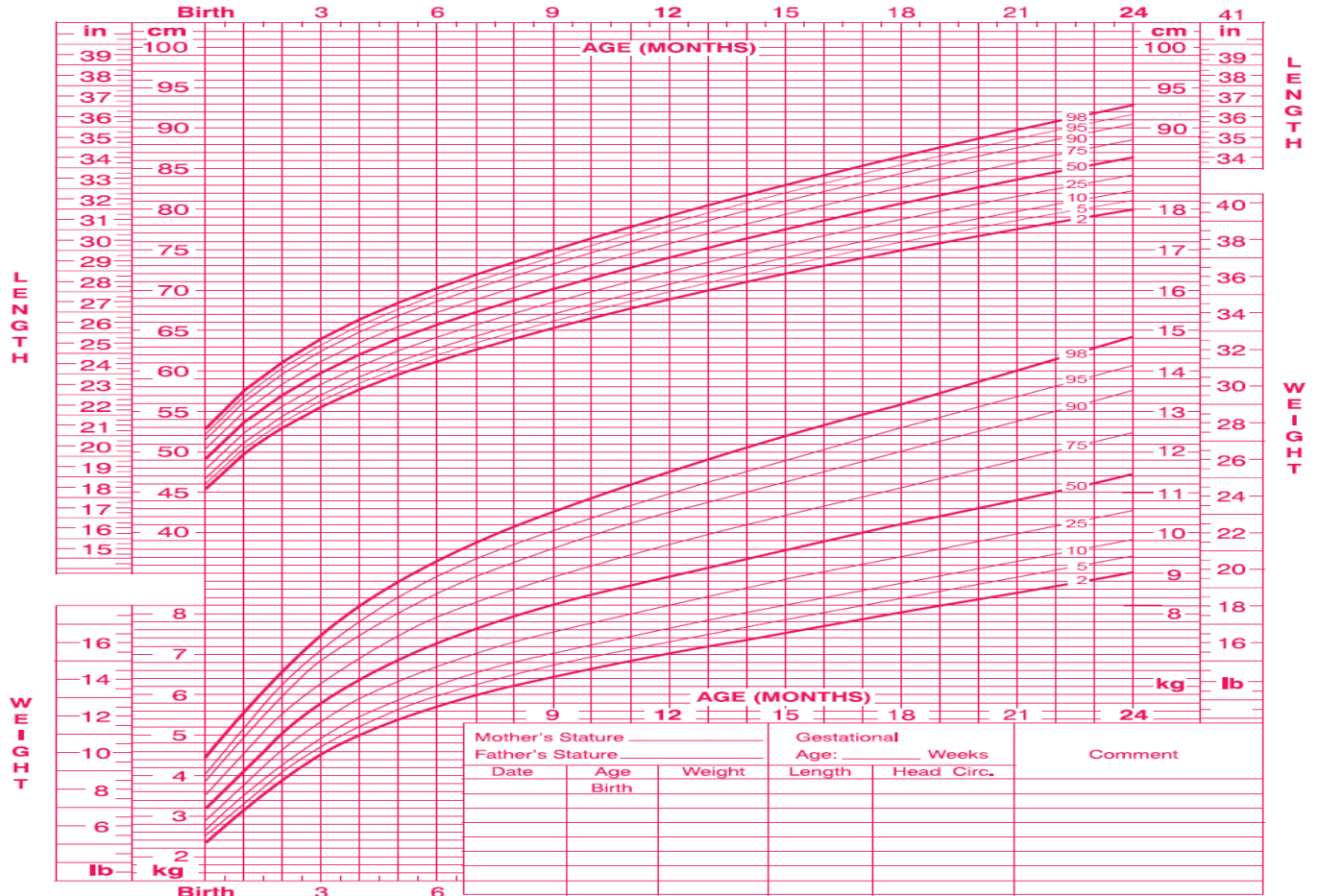




**Birth to 24 months: Girls**  
**Length-for-age and Weight-for-age percentiles**

NAME \_\_\_\_\_

RECORD # \_\_\_\_\_



Published by the Centers for Disease Control and Prevention, November 1, 2009  
 SOURCE: WHO Child Growth Standards (<http://www.who.int/childgrowth/en>)

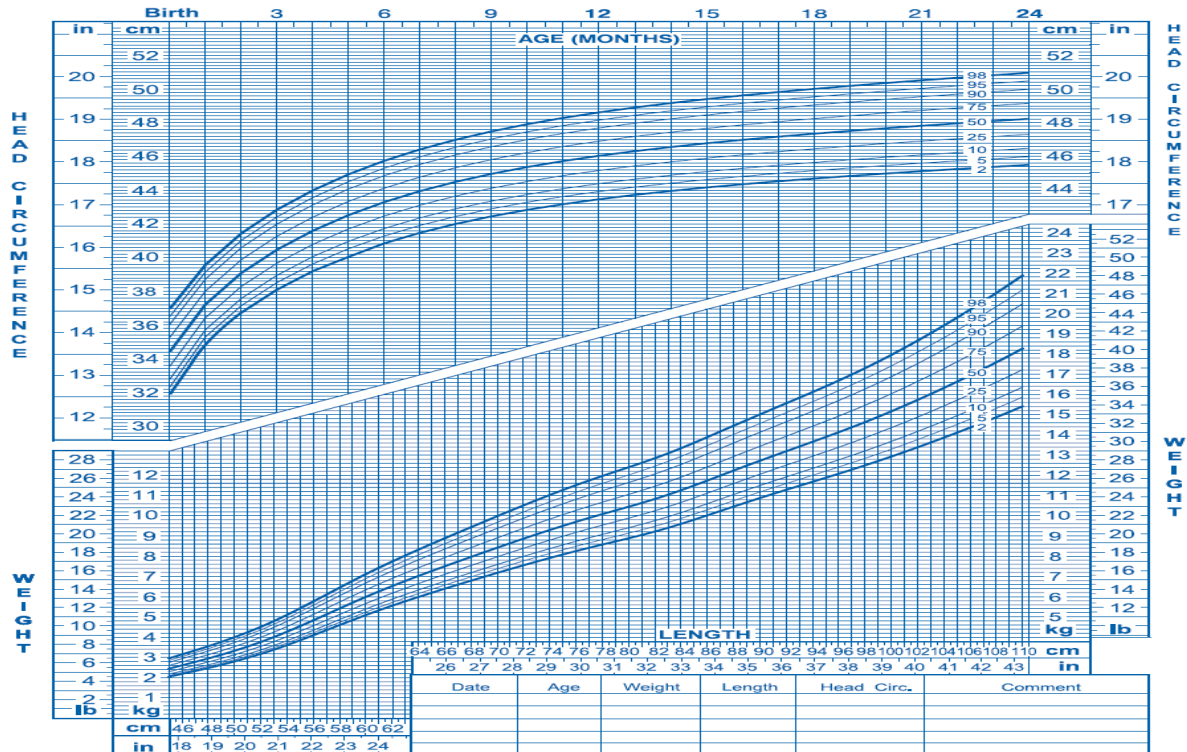




**Birth to 24 months: Boys**  
**Head circumference-for-age and**  
**Weight-for-length percentiles**

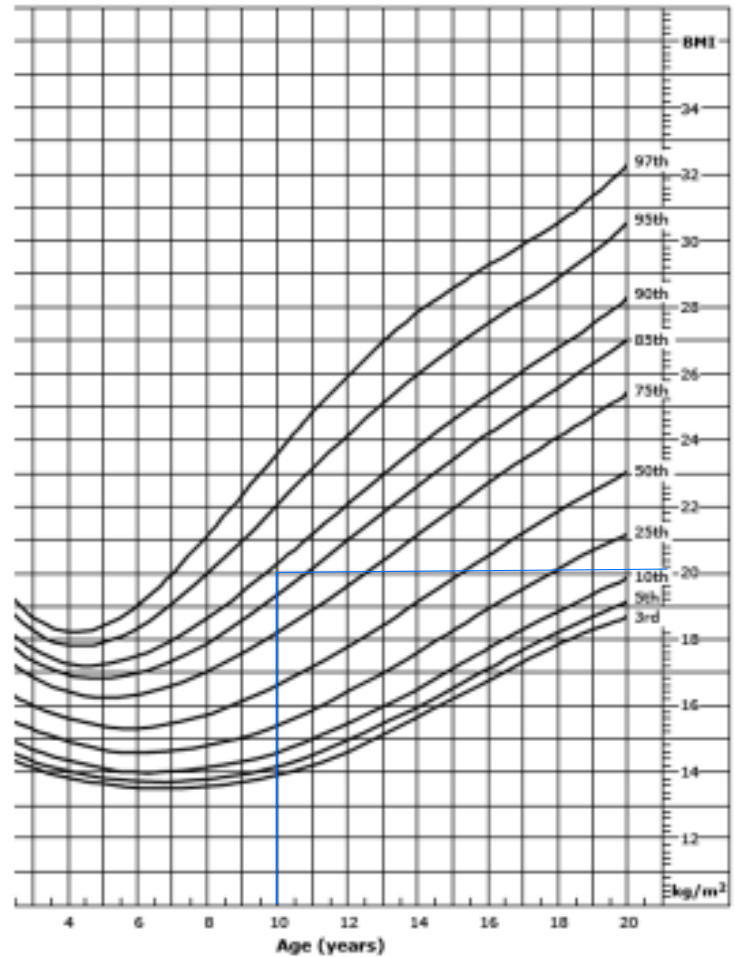
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overweight:  
 $\text{BMI} > 85^{\text{th}} < 95^{\text{th}}$   
percentile  
Obese:  
 $\text{BMI} > 95^{\text{th}}$   
percentile

*in children we don't base obesity/overweight classification on a specific number like adults (overweight  $\geq 25$ , obese  $\geq 30$ ) but instead base on percentile*



# Prader-Willi Syndrome

Obesity due to nutrition (mcc) tends to have children of normal height (or slightly tall). For very short + obese children suspect endocrine disorders (hypothyroidism, GH deficiency, Cushing's)

Prader-Willi:

1<sup>st</sup> 2 yrs of life → low weight

(hypotonia, poor suckling)

Afterwards → obese

(appetite ↑)

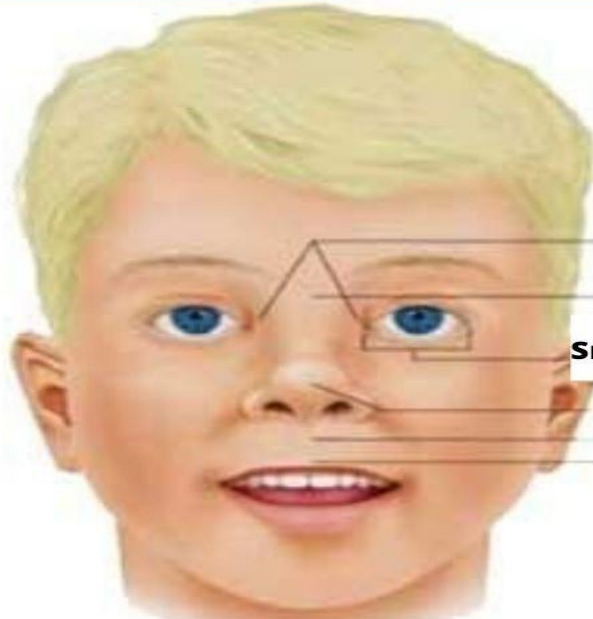


# PRADER WILLI SYNDROME

**A rare genetic disorder**

**Temper outbursts, stubbornness, compulsive behaviour**

**Weak muscle tone, feeding difficulties, poor growth, delayed development, chronic over eating and obesity**



*Almond eyes*

**Epicanthal folds**

**Flat nasal bridge**

**Small palpebral fissure**

**"Railroad track" ears**

**Upturned nose**

**Smooth philtrum**

**Thin upper lip**

\* **Height velocity** — A period of at least six months is necessary for reliable calculation of height velocity in children older than two years :

- 0 to 6 months — 2.5 cm per month
- 7 to 12 months — 1.25 cm per month
- 12 to 24 months — 12 cm per year
- 24 to 36 months — 8 cm per year
- 36 to 48 months — 7 cm per year
- 4 to 10 years — 4 to 6 cm per year

Growth velocity is imp. for  
determining if a pathology is present

A 95% of normal individuals will attain final heights within 2 standard deviations of mid-parental height ( $\pm 8.5$  cm)

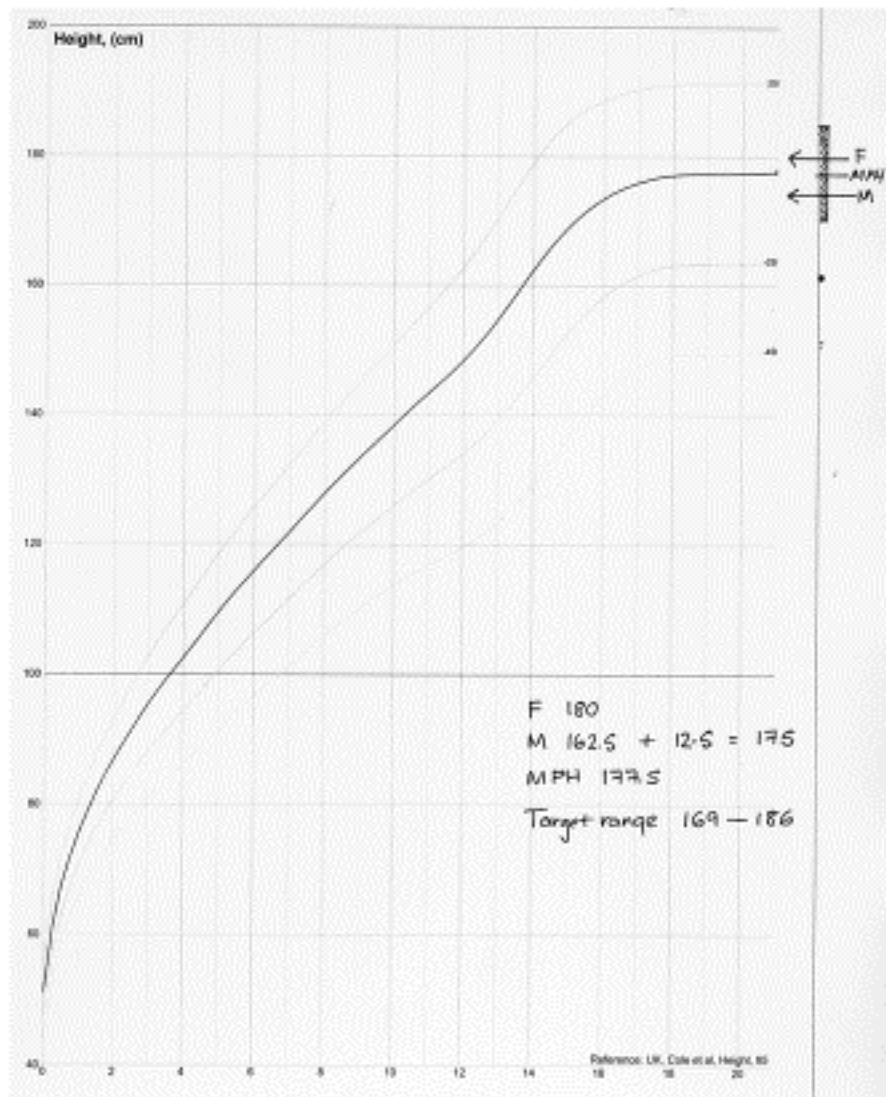


This is a simplified version of:

$$\text{Boy: } \frac{\text{height of dad} + (\text{height of mom} + 13)}{2}$$

(this is to account for diff b/w genders)

$$\text{Girl: } \frac{(\text{height of dad} - 13) + \text{height of mom}}{2}$$



# Evaluation of Growth





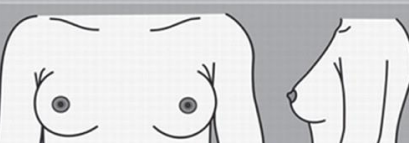
- How short is the child?
- Is the child's height velocity (HV) impaired?
- What is the child's likely adult height?
- Is the child growing on a centile appropriate for the genetic target height?








# Pubertal staging

- Thelarche - appearance of breast buds
- Pubarche - appearance of pubic hair
- Adrenarche - appearance of adrenal hormones (DHEA)
- Menarche - first menstrual cycle

# Tanner Staging- females

I		<b>Stage I:</b> prepubertal
II		<b>Stage II:</b> breast bud with elevation of breast and papilla; enlargement of areola <i>only elevation of areola</i>
III		<b>Stage III:</b> further enlargement of breast and areola; no separation of contour <i>Elevation of areola + skin</i>
IV		<b>Stage IV:</b> areola and papilla form secondary mound above level of breast <i>Projection of areola + skin</i>
V		<b>Stage V:</b> mature stage; projection of papilla only, related to recession of areola <i>Areola no longer protrudes</i>

# Tanner Staging- males

Childhood	I		3	25
Early Puberty	II		4	25-32
Mid Puberty	III		10	33
Late Puberty	IV		16	41-45
Adulthood	V		25	>45

Sexual hair,  
dark, coarse, curly

hair fills pubis

hair extend to thigh

pubertal size of testis  $\geq 4$  ml

Adult reaches 25 ml

# Causes of Short Stature

*but not every familial growth stature is benign  
(check if growth velocity is normal)*

- Normal (familial and constitutional delay)  
*↑*
- Small for gestational age
- Syndromes
- Skeletal dysplasias
- Systemic illnesses
- Endocrine disorders
- Psychosocial circumstances
- Idiopathic short stature



**2 to 20 years: Boys**  
**Stature-for-age and Weight-for-age percentiles**

NAME \_\_\_\_\_

RECORD # \_\_\_\_\_

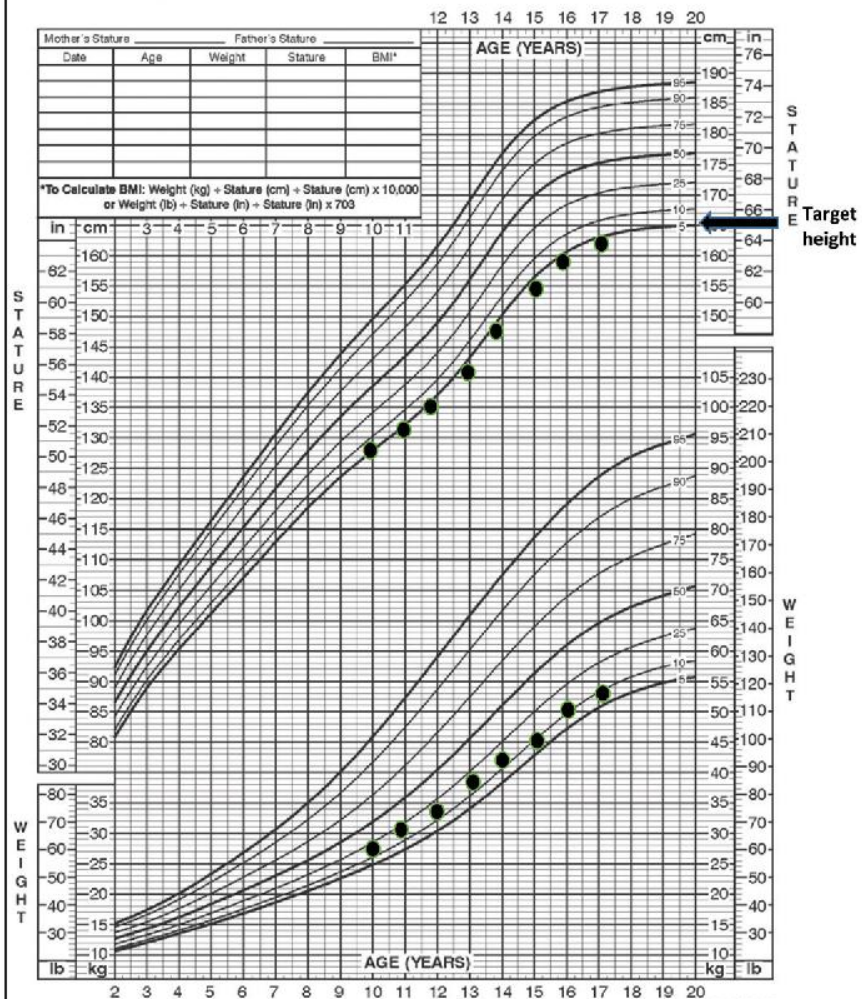


Figure courtesy of the US Centers for Disease Control and Prevention.

Published May 30, 2000 (modified 11/21/00).  
 SOURCE: Developed by the National Center for Health Statistics in collaboration with  
 the National Center for Chronic Disease Prevention and Health Promotion (2000).  
<http://www.cdc.gov/growthcharts>



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Feature	Familial short stature	Constitutional Delay
Parents' stature	Small (one or both)	Average
Parents' puberty	Usual timing	Delayed
Birth length	Normal	Normal
Growth (0 to 2 years)	Normal	Normal to slow
Growth (puberty)	Normal	Slow
Bone age	Normal	Delayed
Timing of puberty	Normal	Delayed

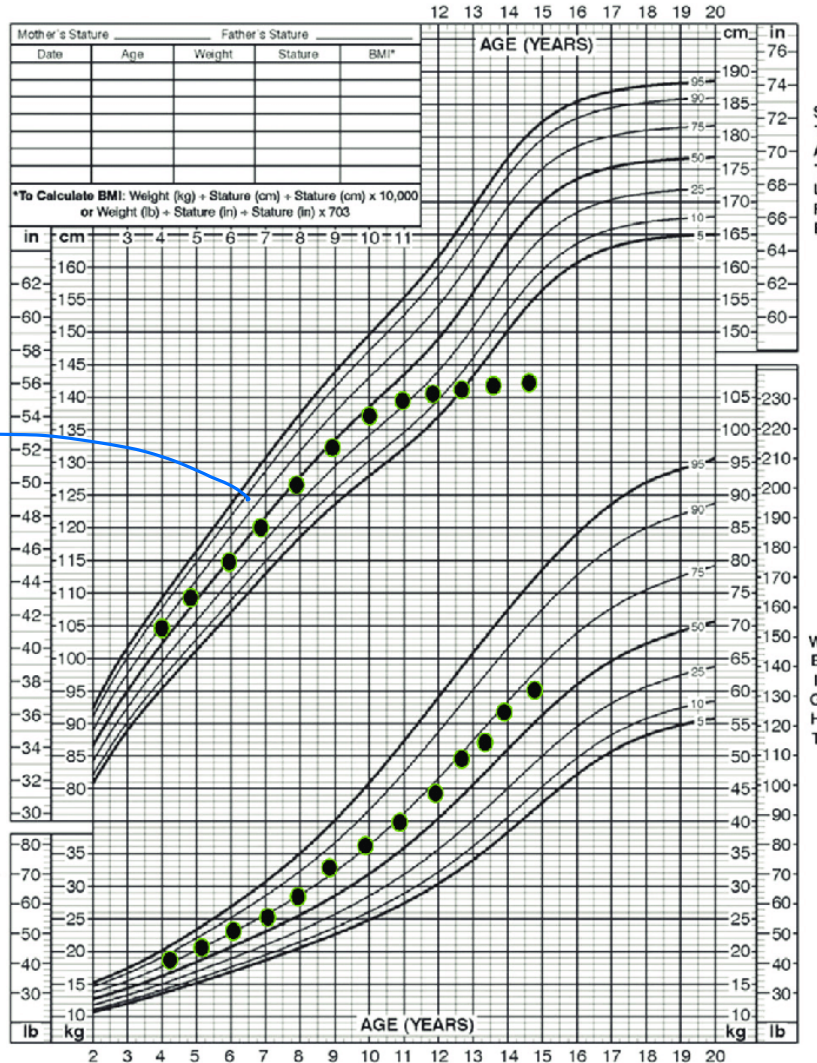
*Constitutional delay: Bone age delayed up to 2 years  
(more should make you suspect pathology like a systemic illness)*



2 to 20 years: Boys  
 Stature-for-age and Weight-for-age percentiles

NAME \_\_\_\_\_

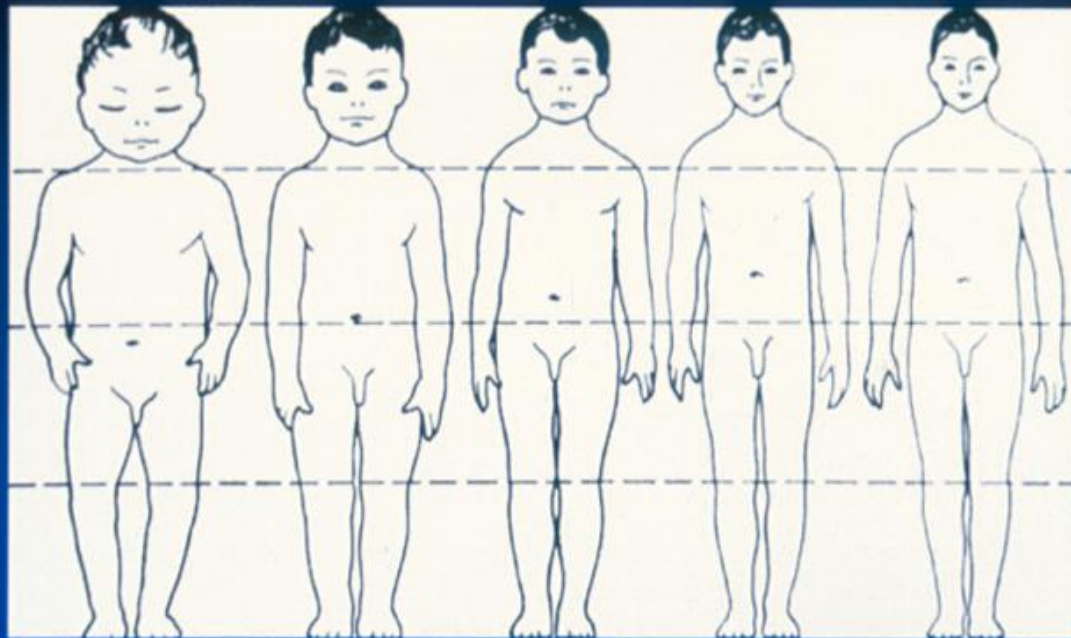
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Normal height than sudden deceleration: suspect systemic illness + acquired Gil deficiency



## Stature divided into quarters



newborn

2

5

15

adult

Age (years)

# Body proportions

- The proportions of the body change during fetal and postnatal growth. The most commonly used descriptors of body proportions are the ratio of the upper body segment to the lower body segment and the ratio of arm span to height.
- Approximate normal ratios of US/LS :
  - Birth — 1.7
  - 3 years — 1.33
  - 5 years — 1.17
  - 10 years — 1.0
  - >10 years — <1.0
- ↑ in children with rickets, achondroplasia, and Turner syndrome (because of decreased limb length) .
- ↓ in Marfan syndrome (because of increased limb length).

↓  
Lower segment: Symphysis pubis to floor

# Achondroplasia

Normal trunk  
+  
Short limbs



# Endocrine disorders

- Growth hormone deficiency and resistance cause (congenital or acquired)
- Thyroxine deficiency
- Cortisol excess
- Precocious puberty
- Idiopathic short stature

# Evaluation of the child with short stature - history

- History of growth pattern and previous measurements
- Parental heights, puberty
- Birthweight, postnatal hx
- General health
- Psychosocial situation

# Evaluation of the child with short stature - auxology

- Accurate measurement of child and parents
- Sibling measurement in selected cases
- Plot height and weight
- Plot mid parental height and target range
- Bone age by single observer

# X-ray of left hand and wrist for bone age assessment



# Evaluation of the child with short stature - examination

- General appearance and nutrition
- Body proportions
- Dysmorphic features
- Systemic examination including heart
- BP
- Pubertal status
- Fundi *-if you suspect a central cause*



# Investigation of short stature

- Low threshold for karyotype in girls with short stature.
- Short stature screening
- Further short stature investigation

# Short stature screening

- FBC and film, ESR, TTG  
a'bodies, LFT, Blood gas
- Karyotype *Added if child < 3yr or has poor nutritional status*
- TFT, IGF-1, IGFBP3, prolactin, cortisol  
*Screening of GH levels*
- Creatinine, Lytes, calcium and phosphate
- Urinalysis and culture

# Further investigation of short stature

- Endocrine stimulation testing
  - ITT/arginine/clonidine/glucagon
- Pituitary imaging
- Genetic evaluation – clinical, laboratory
- Skeletal survey

# Conditions where growth hormone therapy is recommended

- Growth hormone deficiency
- Turner syndrome/Noonan Syndrome/SHOX
- Prader-Willi syndrome.
- SGA with no catch up by 4 years
- Chronic renal insufficiency.
- ISS ??

*(even tho some of these there is no GH deficiency)*

# Treatment of GH deficiency

## Recombinant human growth hormone (RHGH)

- Daily bedtime subcutaneous injections of aqueous solutions of biosynthetic (recombinant) GH
- Follow up q 3 months

**THANK YOU**