Short stature

Rasha Odeh

MB,BS.MRCPCH

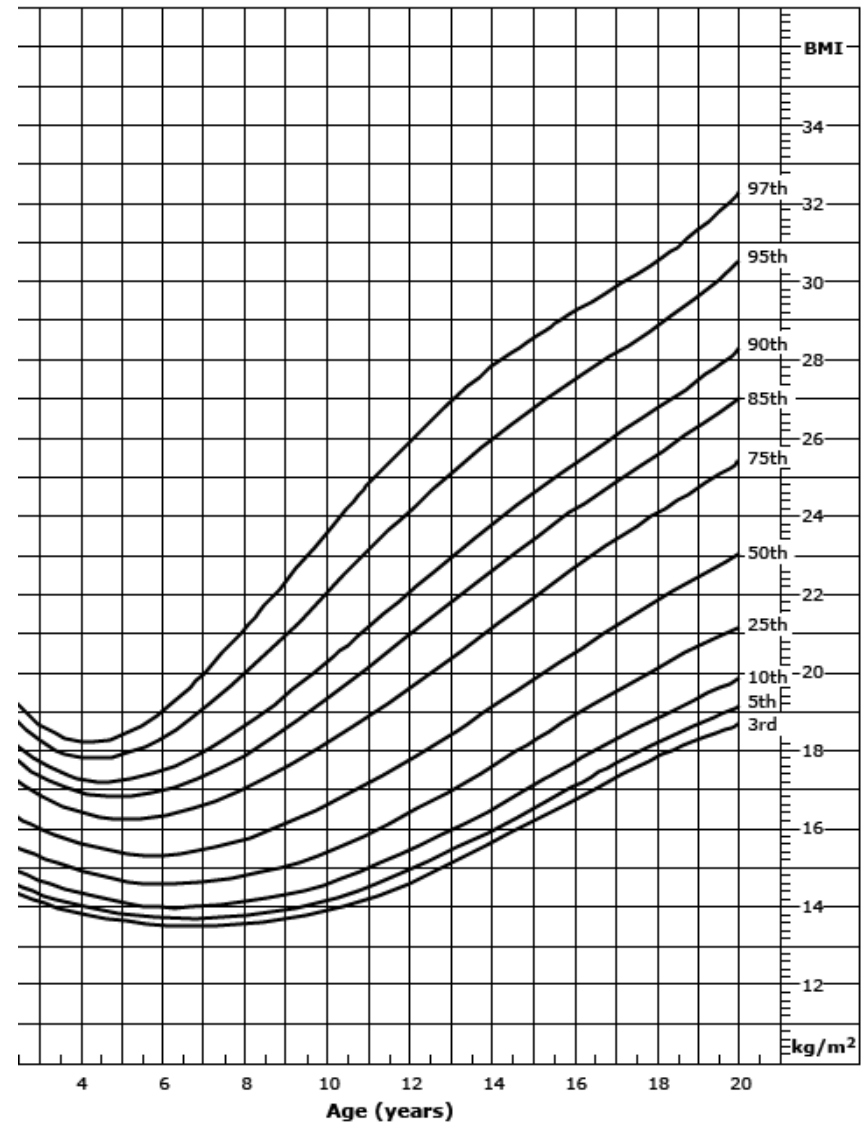Assistant professor of pediatric endocrinology

University of Jordan

Growth

- Growth in childhood, one of the most fascinating, complex and dynamic biological processes, is tightly controlled and regulated.

- remains one of the most useful of all indices of public health and economic well being, specifically in developing countries and countries in transition

Growth Parameters

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

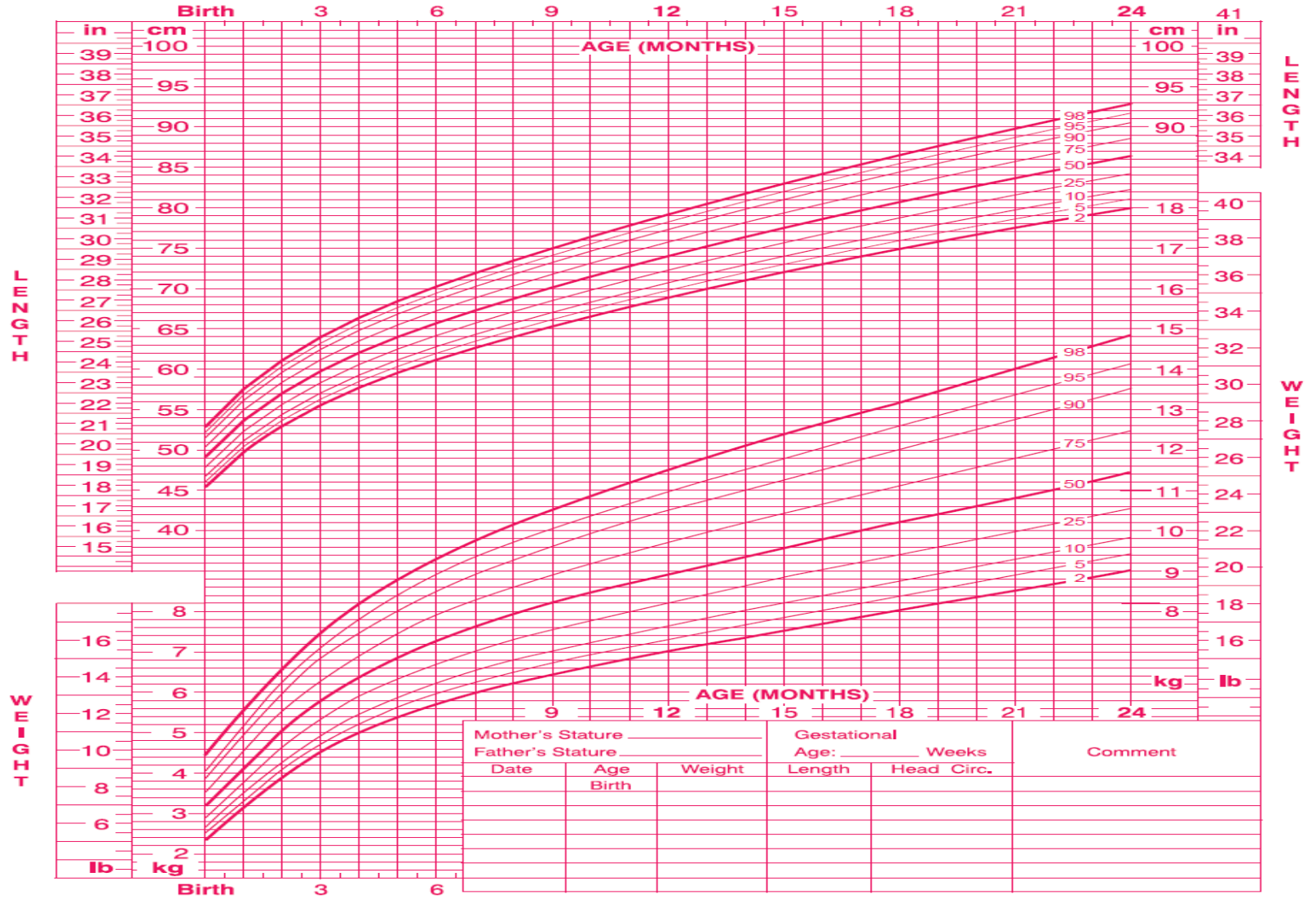overweight: BMI>85th<95th percentile
Obese: BMI>95th percentile

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

NAME _____

RECORD # _____

Mother's Stature _____ Father's Stature _____

Date	Age	Weight	Stature	BMI*

*To Calculate BMI: Weight (kg) ÷ Stature (cm) ÷ Stature (cm) x 10,000
or Weight (lb) ÷ Stature (in) ÷ Stature (in) x 703

AGE (YEARS)

STATURE

WEIGHT

SAFER · HEALTHIER · PEOPLE™

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

NAME _____

RECORD # _____

AGE (MONTHS)

Birth 3 6 9 12 15 18 21 24

LENGTH

WEIGHT

Mother's Stature _____		Gestational	
Father's Stature _____		Age: _____ Weeks	Comment
Date	Age	Weight	Length	Head Circ.
	Birth			

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

NAME _____

RECORD # _____

Outline

- Normal Growth and puberty

- Causes of short stature

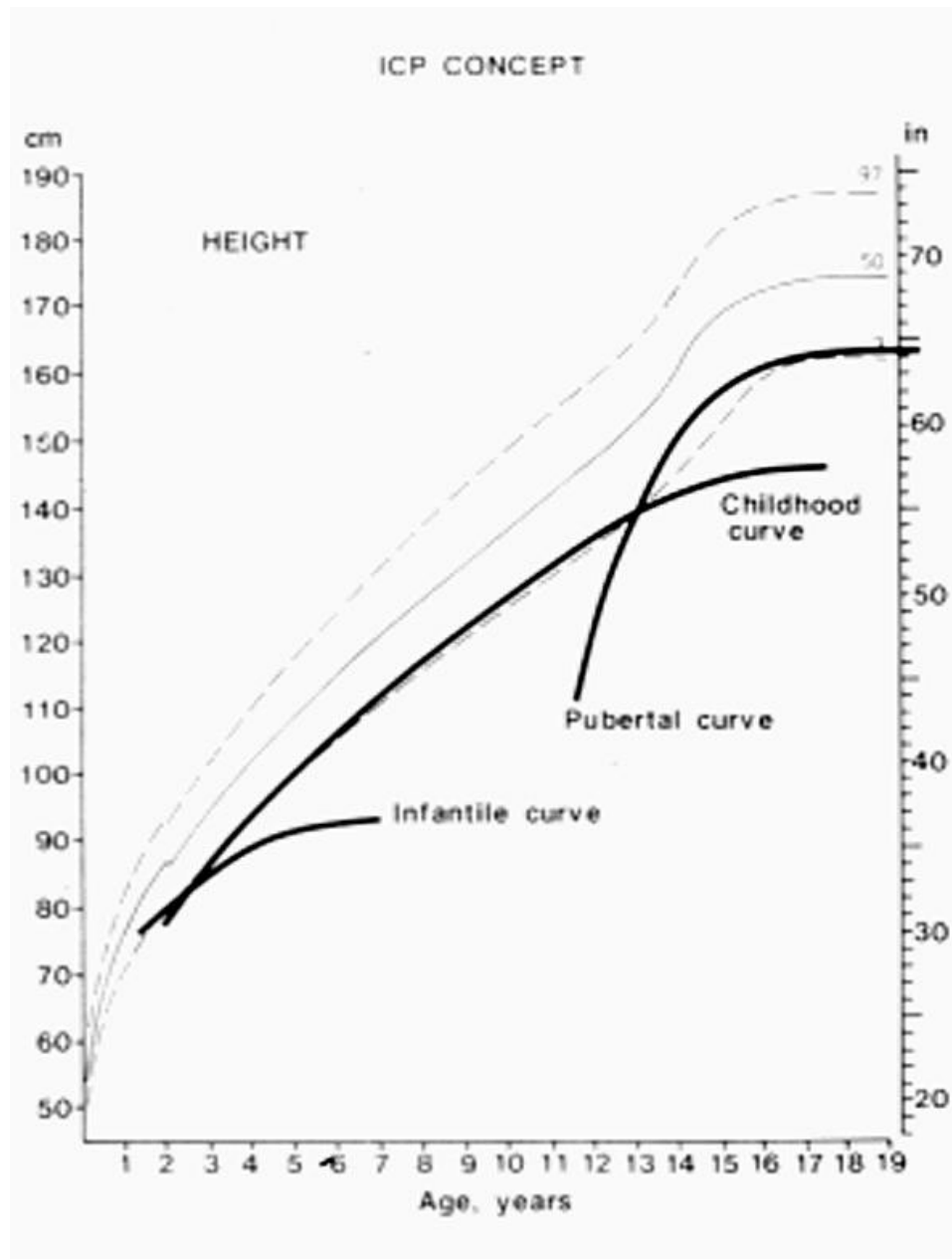- Approach to a short child

The ICP concept of growth:

Infancy

Childhood

Puberty

A mathematical model breaking down linear growth from birth to adulthood into 3 components that reflect the different hormonal phases of the growth process.

Karlberg J. Acta Paediatr Suppl 1989; 350: 70-94

ICP Concept

ICP CONCEPT

HEIGHT

Childhood curve

Pubertal curve

Infantile curve

Age, years

ICP Concept - Infantile Phase

- Rapid period of growth from 0 - 2 years

- Continuation of the rapid but decelerating intra-uterine growth phase

- Depends on factors such as nutrition, insulin and the insulin-like growth factors.

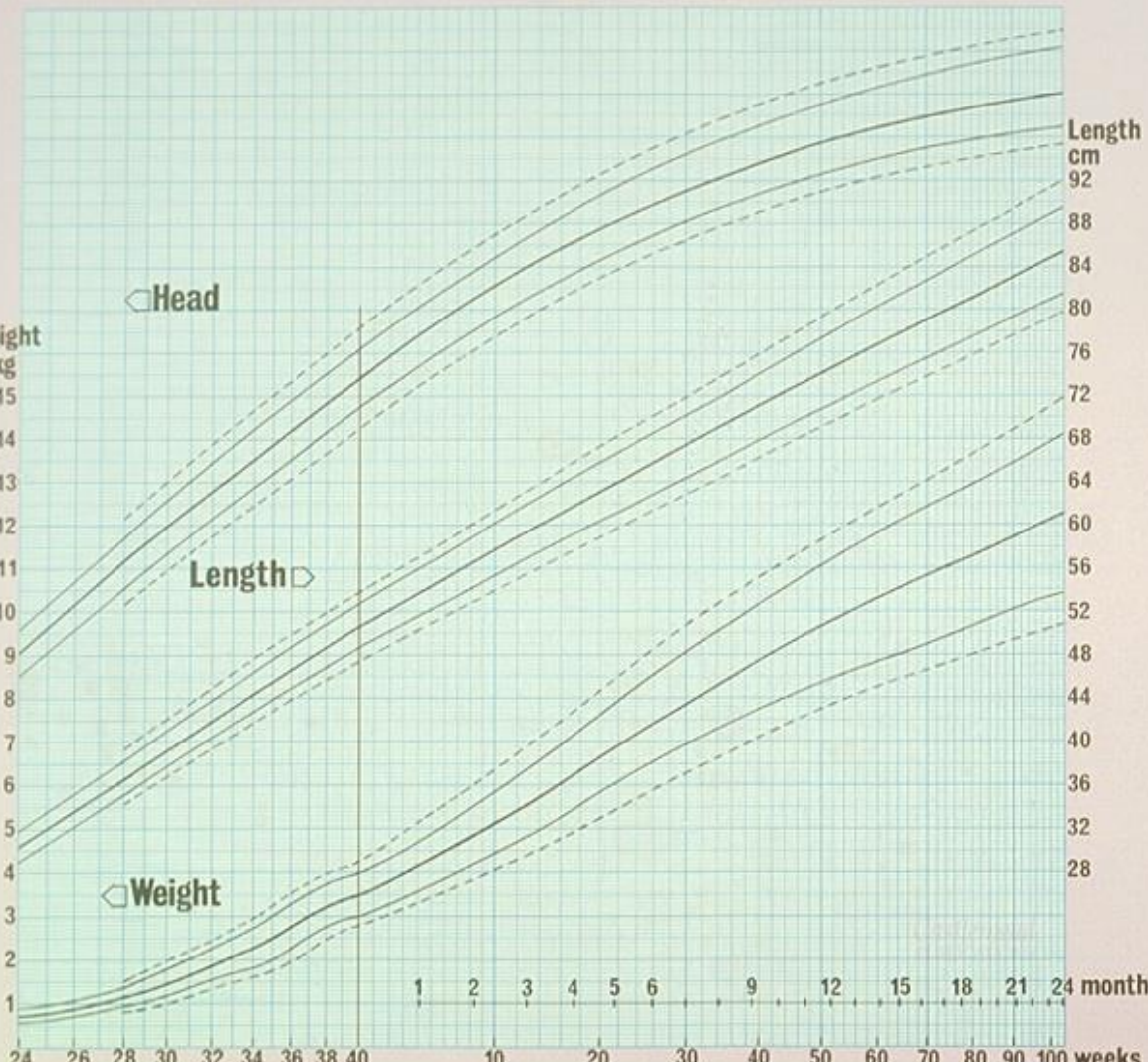- Largely independent of growth hormone

ICP Concept - Infantile Phase

- In general, birth size reflects intra-uterine environment, e.g. nutrition

- Birth size, therefore, has relatively poor correlation with final height.

- From birth to age 2, infant "tracks" on to his/her genetic centile, related to parental heights

Healy Mj etal. Aberdeen growth study.. Arch Dis Child. 1956;31(159):372.

Girls

Head cm: 52, 50, 48, 46, 44, 42, 40, 38, 36, 34, 32, 30, 28, 26, 24, 22, 20

Weight kg: 15, 14, 13, 12, 11, 10, 9, 8, 7, 6, 5, 4, 3, 2, 1

Length cm: 92, 88, 84, 80, 76, 72, 68, 64, 60, 56, 52, 48, 44, 40, 36, 32, 28

◁Head
Length▷
◁Weight

1 2 3 4 5 6   9   12   15   18   21   24 months

24 26 28 30 32 34 36 38 40   10   20   30   40 50 60 70 80 90 100 weeks

Reg. No.
Surname

Forename

Date of Birth   / /
Expected Date of Delivery (EDD)   / /

Treatment/Notes

Key to Centiles
------------------- 97th
_____ 90th
━━━━━━━━━ 50th
_____ 10th
------------------- 3rd

ICP Concept - Childhood Phase

- Long phase of growth from 2 ~ 12 years (onset of pubertal growth)

- Slower, slightly decelerating curve

- Dependent on growth hormone & thyroxine

- Healthy child will stay on childhood (genetic) centile until puberty

ICP Concept - Pubertal Phase

- From around age 12 (time of puberty) to final height

- Dependent on the sex steroid estrogen   which is aromatised from testosterone and which causes an increase in growth hormone secretion

- Resulting growth acceleration is limited by fusion of the epiphyses (due to strogen in both sexes)

ICP Concept - Pubertal Phase

- Variable growth phase in terms of :
  age of onset • *duration* • *intensity*

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

Pubertal (Tanner) staging

Stages of breast development versus height velocity in girls

B1 – prepubertal	Prepubertal (childhood phase)
B2 – breast budding	Increase in height velocity
B3 – breast mound	Peak height velocity B2-3
B4 – angle between areola and breast mound	Slowing of height velocity; onset of periods (menarche)
B5 – adult breast shape	Adult height

Genital stages versus height velocity in boys

G1 – prepubertal scrotum and penis, testicular volume <4 ml	Prepubertal – childhood phase
G2 – prepubertal penis, enlargement of scrotum, testes ≥ 4 ml	Still in childhood phase, therefore height velocity slower than in G1
G3 – penis begins to enlarge	Increase in height velocity
G4 – penis lengthens and broadens, testes ~12-15 ml	Peak height velocity
G5 – adult, testes ~ 15-20 ml	Significant further growth followed by final height

# Height velocity graph showing different growth patterns in boys (solid line) and girls (dotted line)
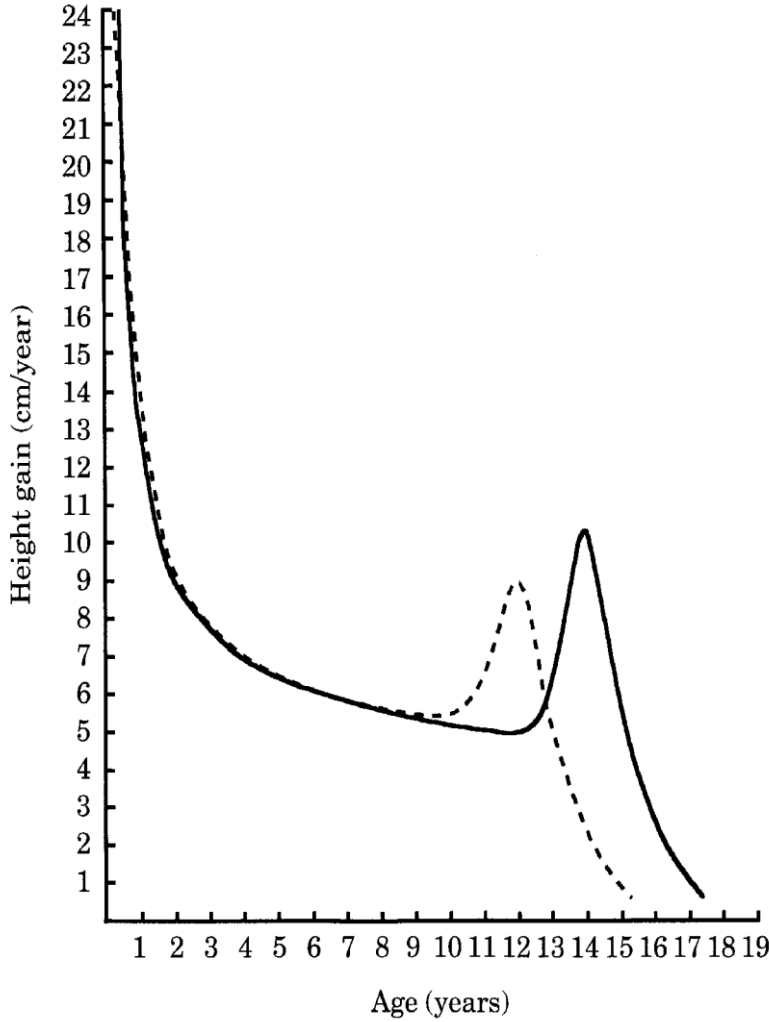
Mid-parental height and target range - girls

- Plot Mother's height on growth chart

- Plot Father's height *minus 13* cm

- Plot the mid-parental height (MPH)

  (the mean of these two measurements)

- Plot the target range (MPH ± 8.5 cm)

Mid-parental height and target range - boys

- Plot Father's height on growth chart

- Plot Mother's height *plus* 13 cm

- Plot the mid-parental height (MPH)

  (the mean of these two measurements)

- Plot the target range (MPH $\pm$ 8.5 cm)

Height, (cm)

F 180

M 162.5 + 12.5 = 175

MPH 177.5

Target range 169 — 186
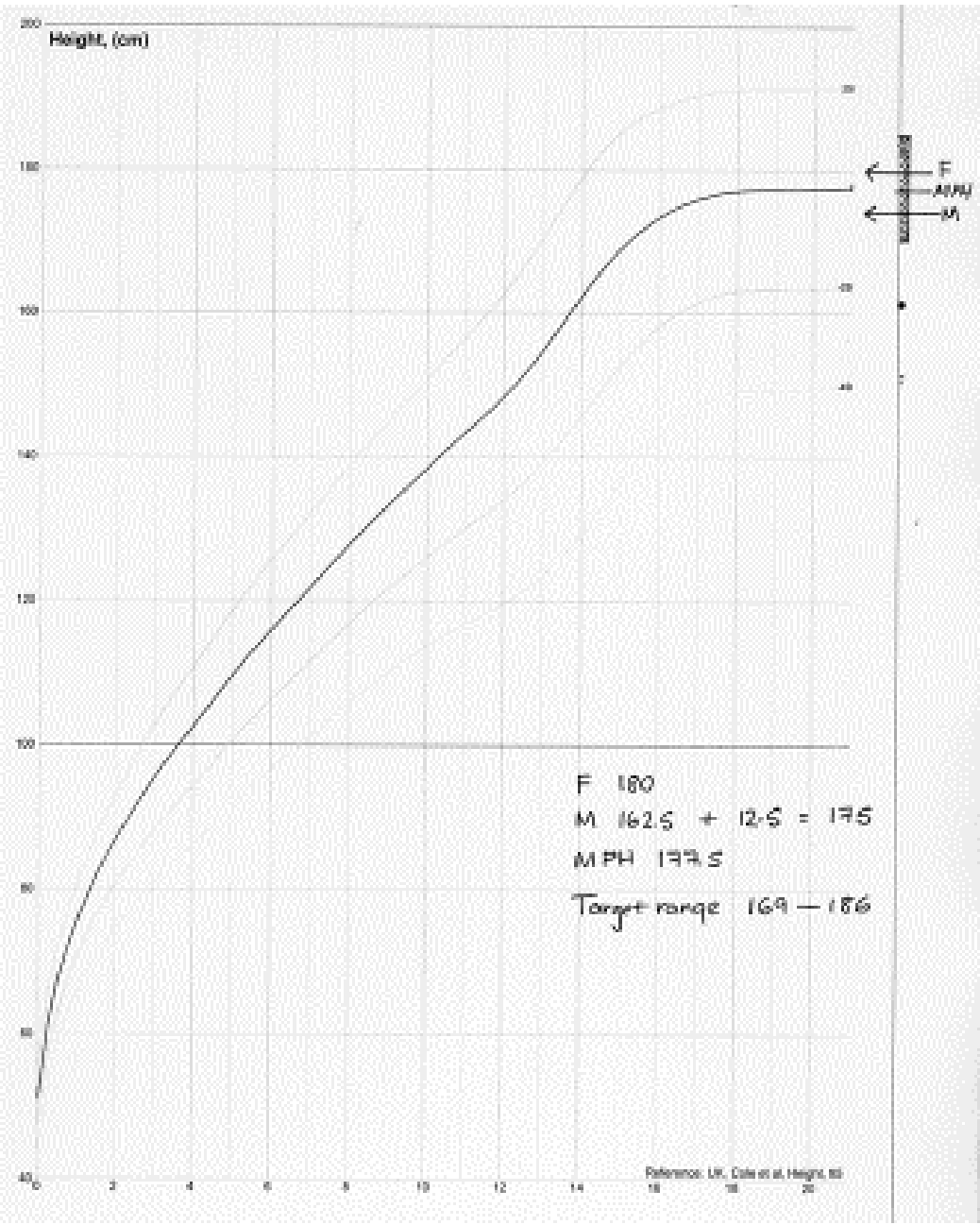
Essentials for growth assessment

- Reliable, accurate measuring equipment

- Accurate measurement of child & parents

- Calculation of decimal age and height velocity

- Careful plotting of growth chart

- Calculation of mid-parental height (MPH) and target range

- Pubertal staging

- Bone age assessment

Measurement of Height

- Reliable, accurate measuring equipment
- Accurate measurement of child & parents !
- Frankfurt plane
- Stretched height
- Measurement of sitting height in selected cases
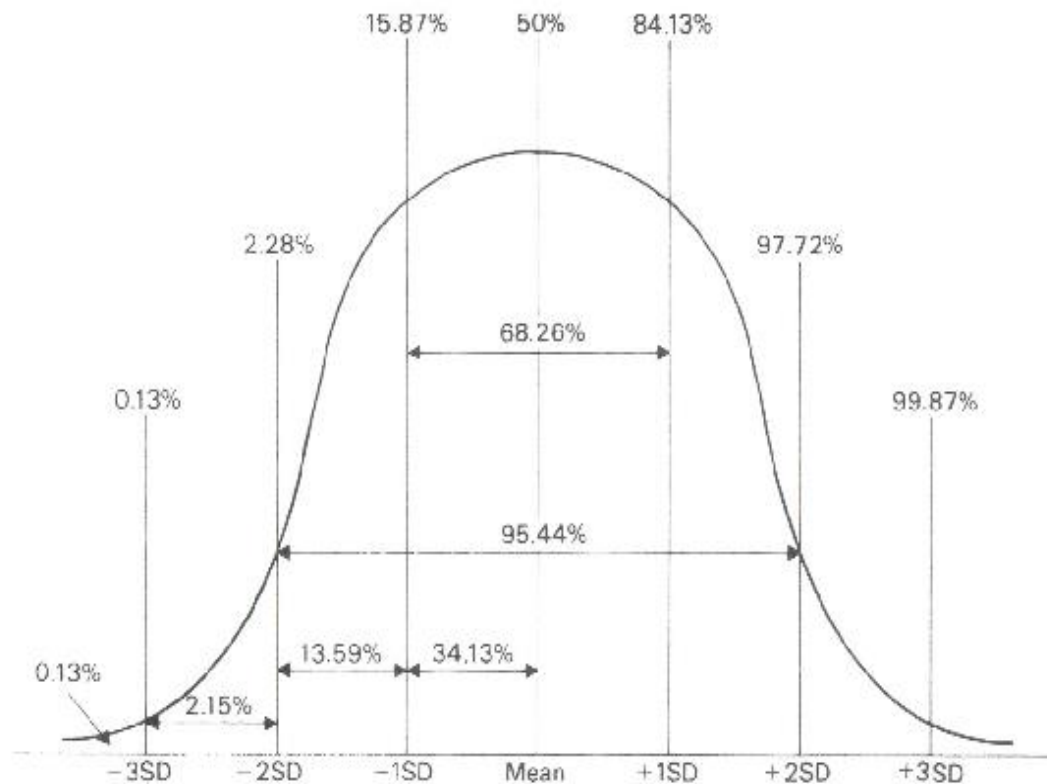
Techniques of measurement
Length

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?
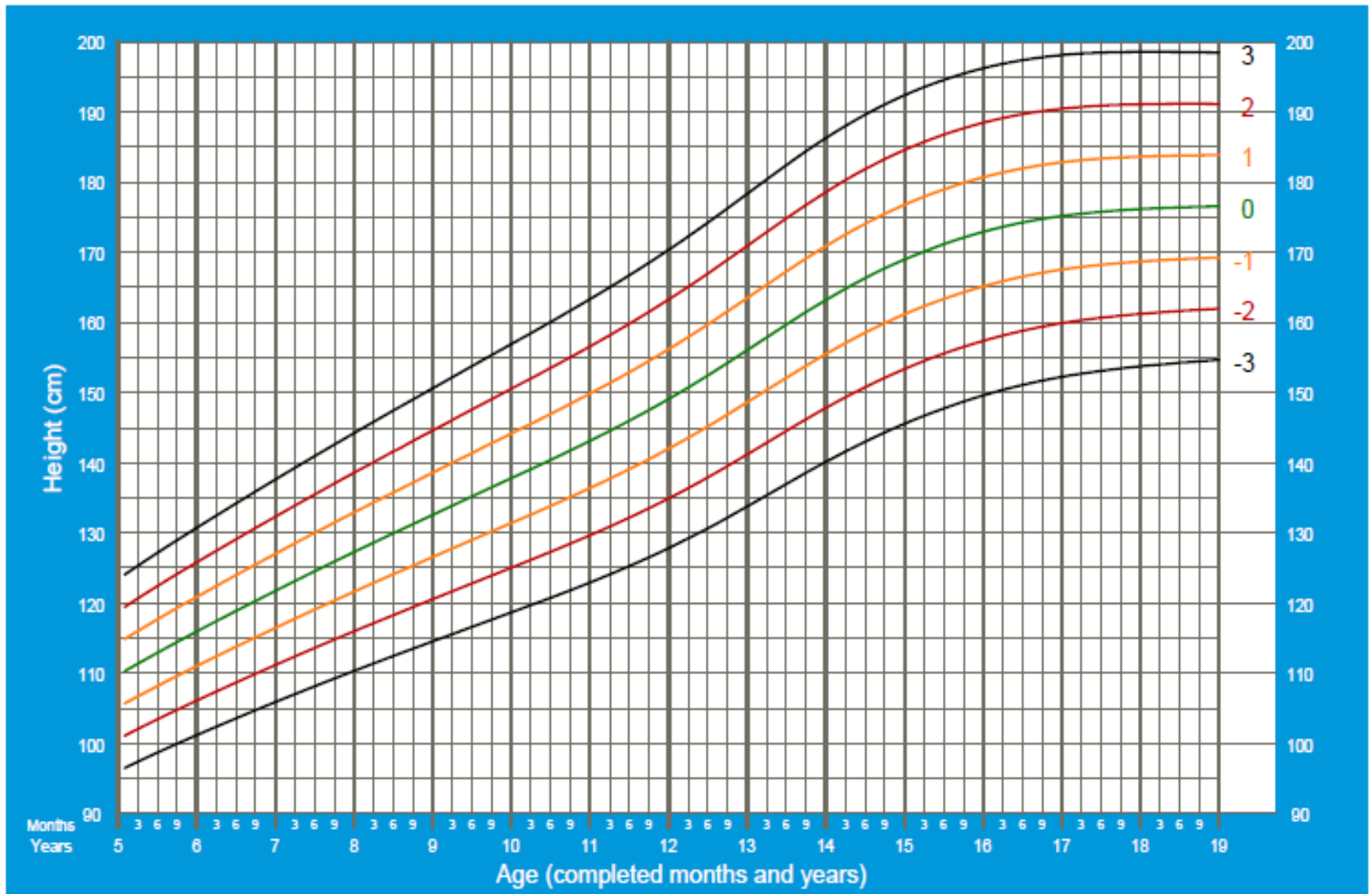
Is the child short?

Relationship between centiles and standard deviations for values with a Gaussian distribution
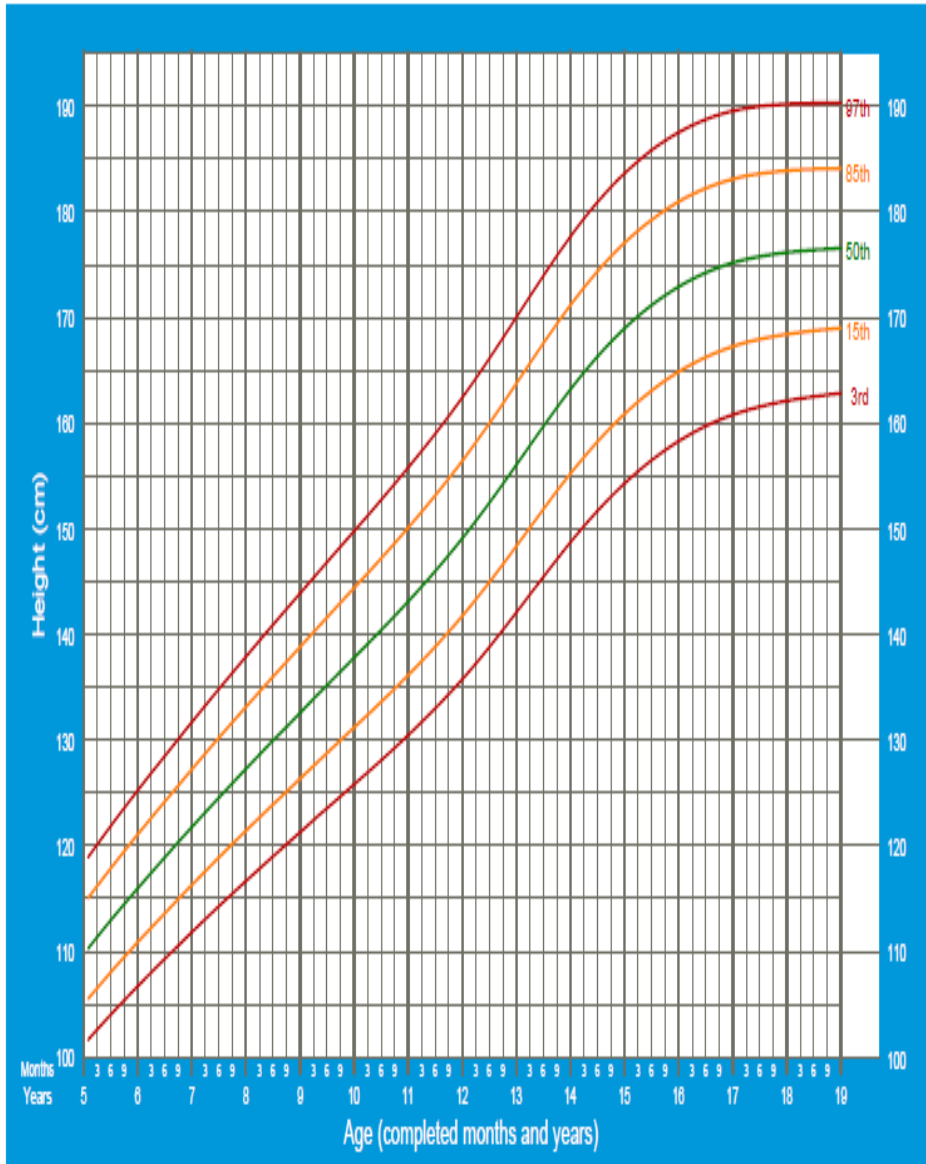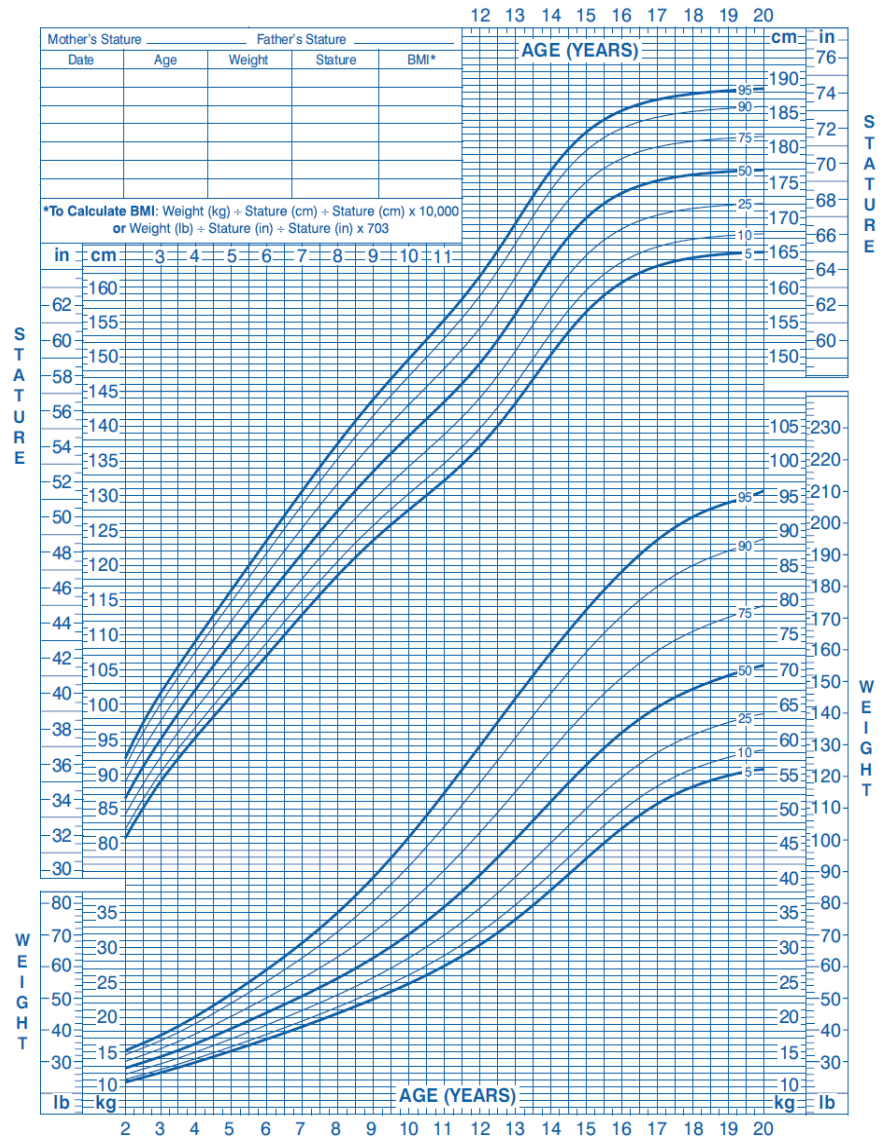
Height-for-age BOYS

5 to 19 years (z-scores)

World Health
Organization

Height (cm) / Age (completed months and years)

2007 WHO Reference

Height-for-age BOYS

5 to 19 years (percentiles)

World Health Organizatio

- 97th
- 85th
- 50th
- 15th
- 3rd

Height (cm)

Age (completed months and years)

Months 3 6 9 3 6 9 3 6 9 3 6 9 3 6 9 3 6 9 3 6 9 3 6 9 3 6 9 3 6 9 3 6 9 3 6 9 3 6 9 3 6 9

Years 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19

2007 WHO Reference

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

NAME _____

RECORD # _____

Mother's Stature _____ Father's Stature _____

Date	Age	Weight	Stature	BMI*

*To Calculate BMI: Weight (kg) ÷ Stature (cm) ÷ Stature (cm) x 10,000
or Weight (lb) ÷ Stature (in) ÷ Stature (in) x 703
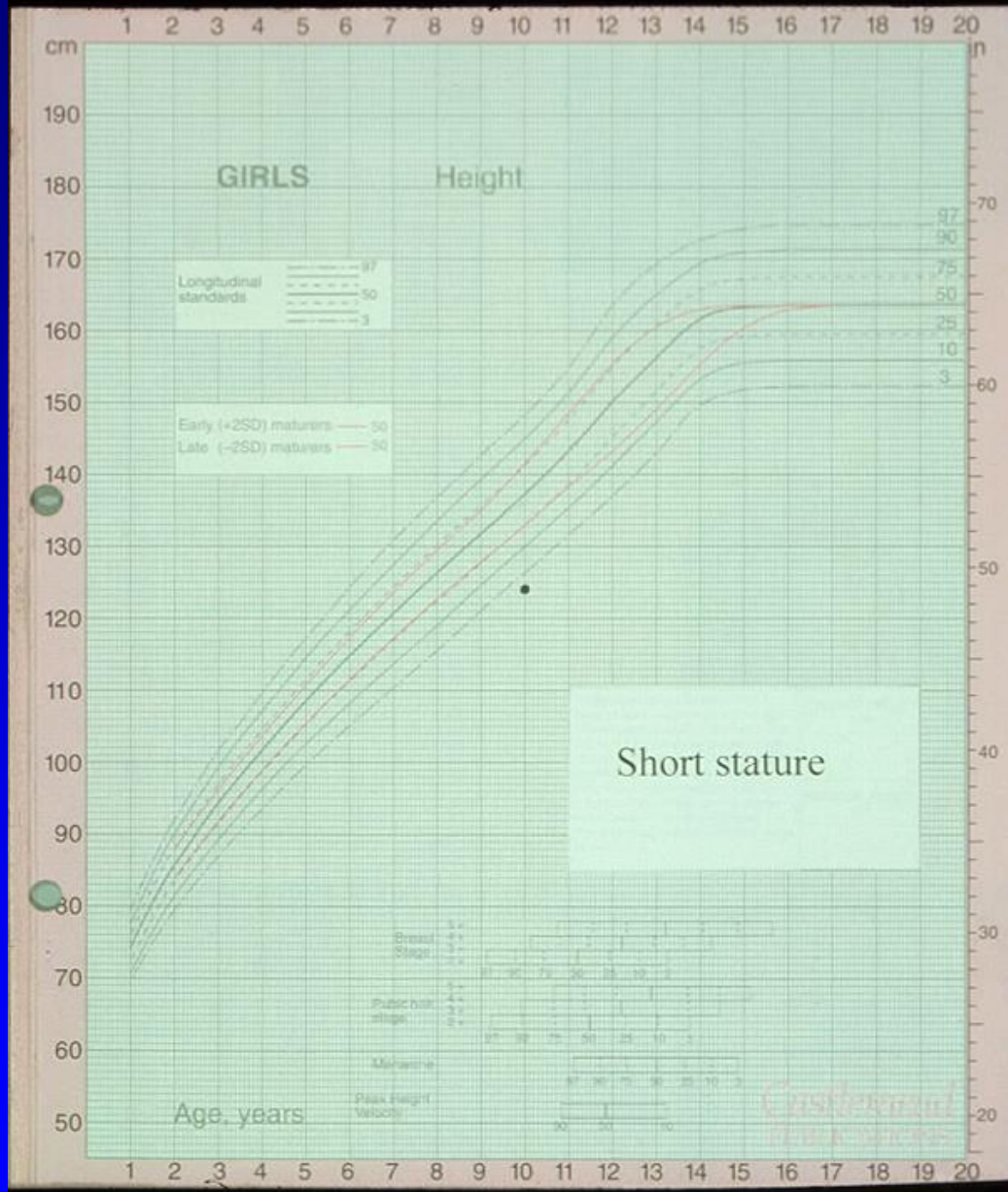
AGE (YEARS)

STATURE

WEIGHT

AGE (YEARS)

CDC

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Short stature:
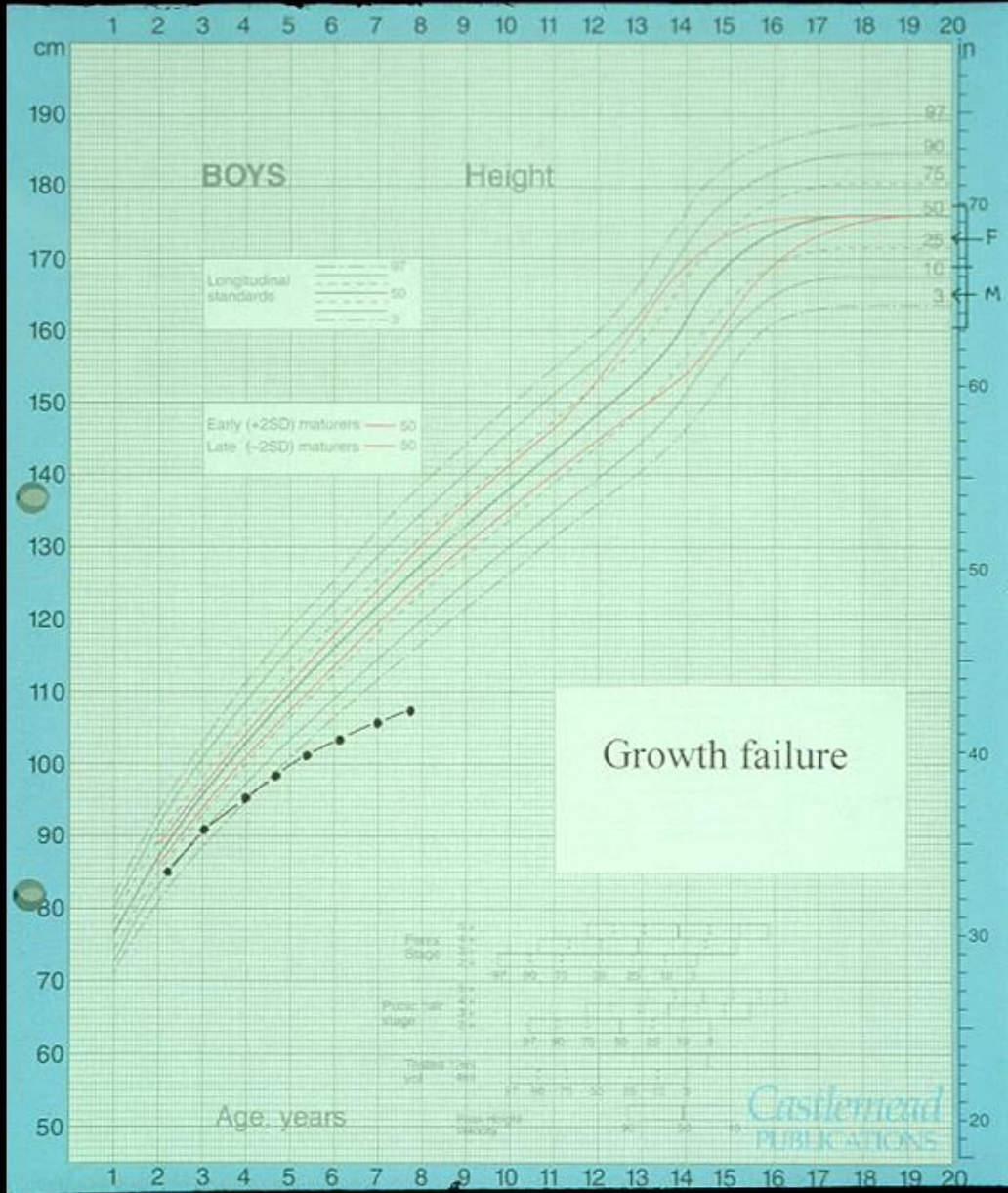
Height below 2 standard deviations from the mean. (3rd centile)
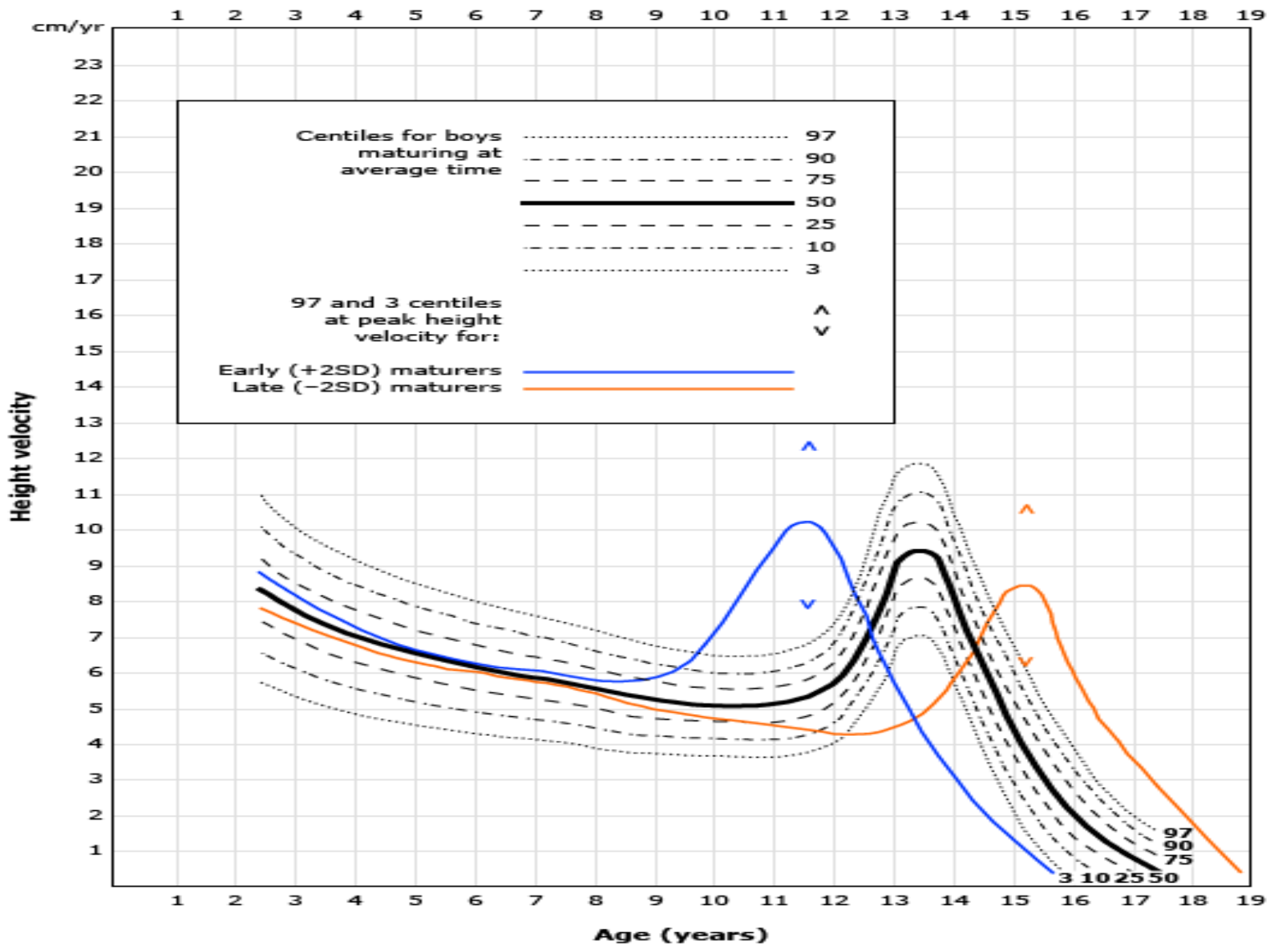
Is the child's height velocity (HV) impaired?

Growth failure:

• Failure to maintain a <u>HV </u>that is appropriate for age and maturity.

• HV should be calculated in (cm/year) using accurate measurements of height and an interval between measurements of at least six months

For children two years and older, **growth failure is likely** if:

- The height-for-age curve has deviated downwards across two major height percentile curves.

- Or, if the child is growing slower than :

- Age 2-4 years: HV less than 5.5 cm/year

- Age 4-6: HV less than 5 cm/year

- Age 6 years to puberty:

- HV less than 4 cm/year for boys

- HV less than 4.5 cm/year for girls

Figure: Height velocity centiles for boys (cm/yr) by Age (years)

Legend:
- Centiles for boys maturing at average time: 97, 90, 75, 50, 25, 10, 3
- 97 and 3 centiles at peak height velocity for: ∧ ∨
- Early (+2SD) maturers (blue)
- Late (−2SD) maturers (orange)

Failure to thrive/weight faltering:

Term usually applied to infants and pre-school children, denoting failure to gain weight at an appropriate rate

What is the child's likely adult height?

- Many methods, different accuracies but in general it depends on:

1. Mid parental height (target height)
2. Bone age : Greulich & Pyle, Tanner Whitehouse…

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

Causes of Short Stature

- Normal (genetic and constitutional delay)
- Small for gestational age
- Dysmorphic syndromes
- Skeletal dysplasias
- Chronic disease
- Endocrine
- Dire social circumstances!
- Idiopathic short stature

Normal genetic short stature

- Child short and normal

- Parent(s) short and normal

- Bone age not delayed

- Child destined to become short adult

Familial (genetic) short stature
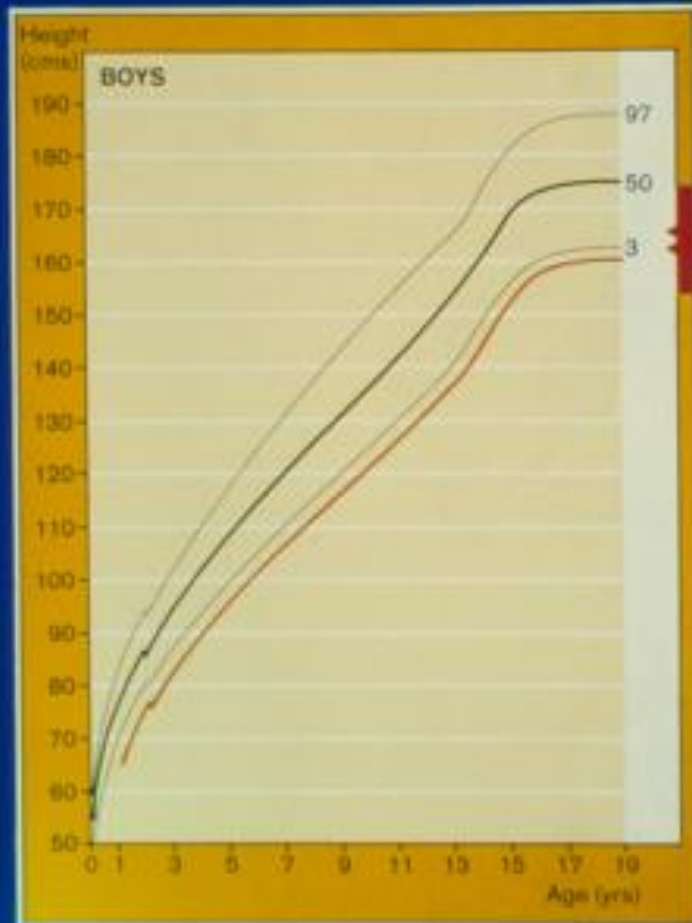
Constitutional delay

- Child short, normal but looks younger than chronological age
- Parent(s) not short, but may have been so in childhood
- Bone age delayed
- Late puberty and catch-up growth
- Final height usually in lower half of target range
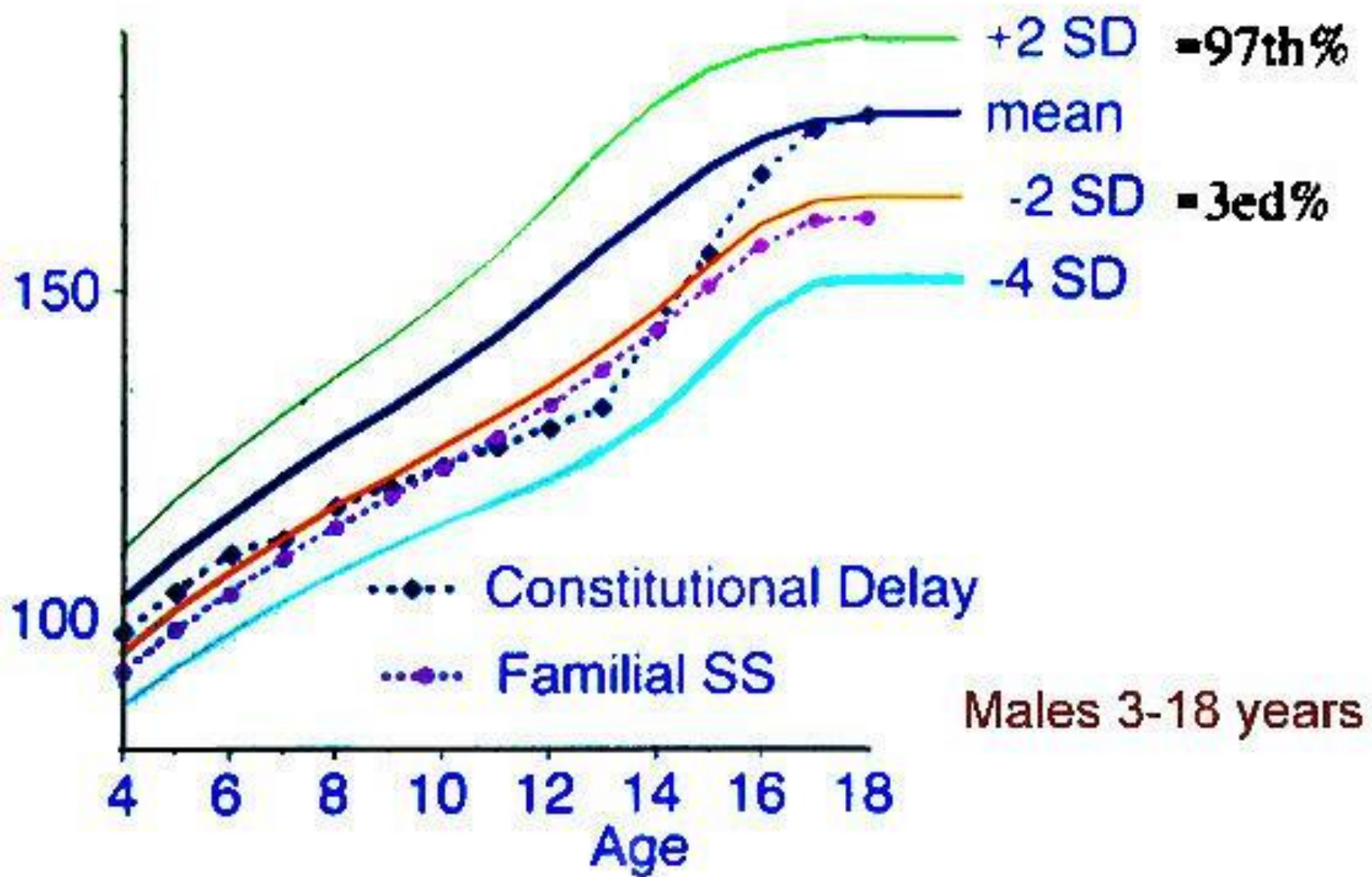
Intrauterine growth retardation (IUGR)

- **Asymmetric (HC>L>Wt)**
  3rd trimester malnutrition; catch-up growth during infancy; normal childhood growth and final height

- **Symmetric (HC=L=Wt)**
  prolonged intrauterine malnutrition; catch-up growth in infancy partial/absent; short stature in childhood despite normal growth rate; short adult

- Most infants born small for gestational age (SGA) experience catch-up growth by two years of age
- About 10 % of SGA infants, particularly those born with more severe SGA, do not experience catch-up growth to reach the normal range by two years of age.

Dysmorphic syndromes

- Turner
- Noonan
- Silver Russel
- William
- Prader-Willi

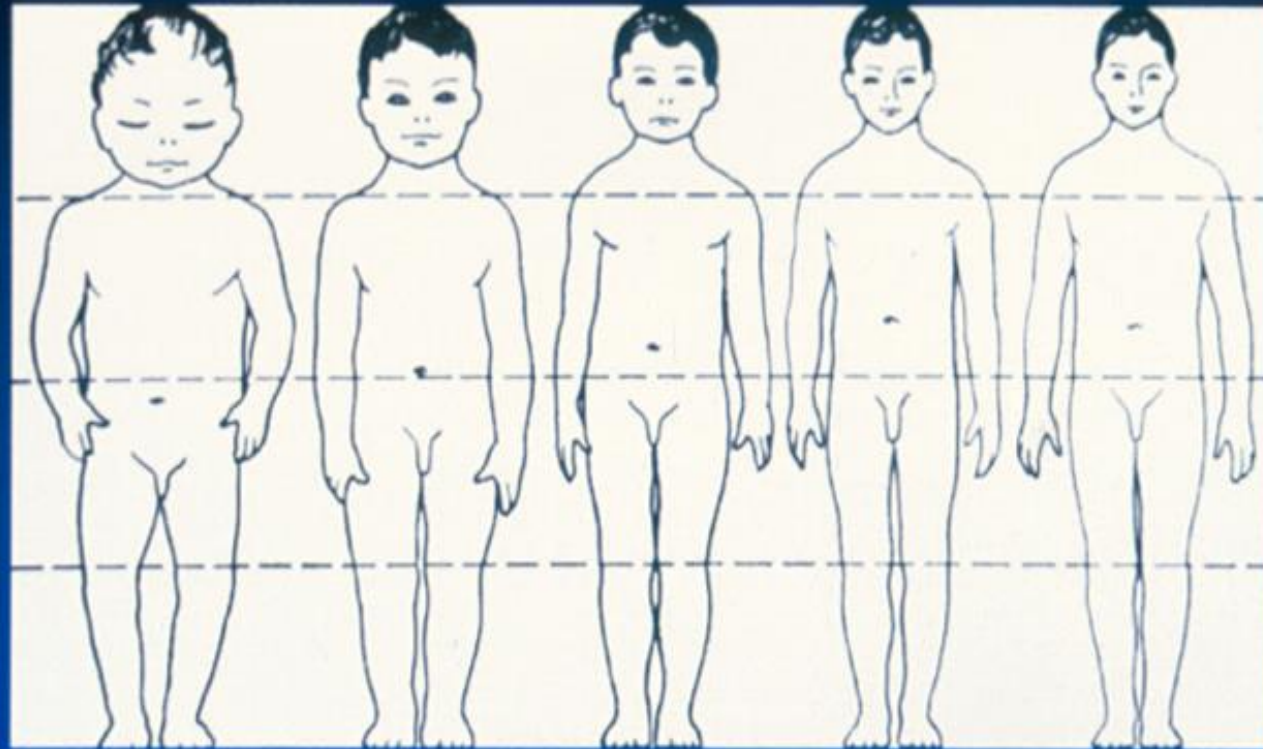# Skeletal dysplasia

- Frank skeletal abnormality affecting

  - epiphyses (epiphyseal dysplasia)

  - metaphyses (metaphyseal dysplasia)

  - spine (spondyleal dysplasia)

  in various combinations (e.g. spondylo-epiphyseal dysplasia)

- Bone age tends to be advanced

- Adult height SDS < Childhood height SDS

Stature divided into quarters

newborn 2 5 15 adult

Age (years)

Achondroplasia

Chronic systemic disease

Many different types

Any chronic condition (e.g. GI, cardiac, respiratory, renal, metabolic, CNS, joints) can cause short stature and/or slow growth

Chronic renal disease and GI disorders (e.g. coeliac disease) can be silent, and present with short stature.

Social and emotional short stature

- Children from poor communities tend to be shorter than those from affluent areas

- Severe short stature with growth failure may result from emotional abuse/neglect (psychosocial deprivation)

Endocrine disorders

- Growth hormone deficiency and resistance

- Thyroxine deficiency

- Cortisol excess

- Precocious puberty

- Idiopathic short stature

Growth hormone

- anterior pituitary protein secreted by somatotrophs .

- antagonizes insulin action, promotes lipolysis in fat, and augments protein synthesis.

- Secretion is pulsatile

- Stimulated be sleep and exercise, inhibited by free fatty acids

The Growth Hormone / IGF Axis

FORNIX (CORPUS)
FORNIX (CRUS)
CHOR
OF 3R
US CALLOSUM
ANTERIOR COMMISSURE
PREOPTIC RECESS
OPTIC CHIASM
INFUNDIBULUM
PITUITARY GLAND
CEREBRAL PEDUNCLE
PONS
TUBER CINEREUM
MAMMILLARY BODY
INTERPEDUNCULAR FOSSA
OCULOMOTOR NERVE (III)
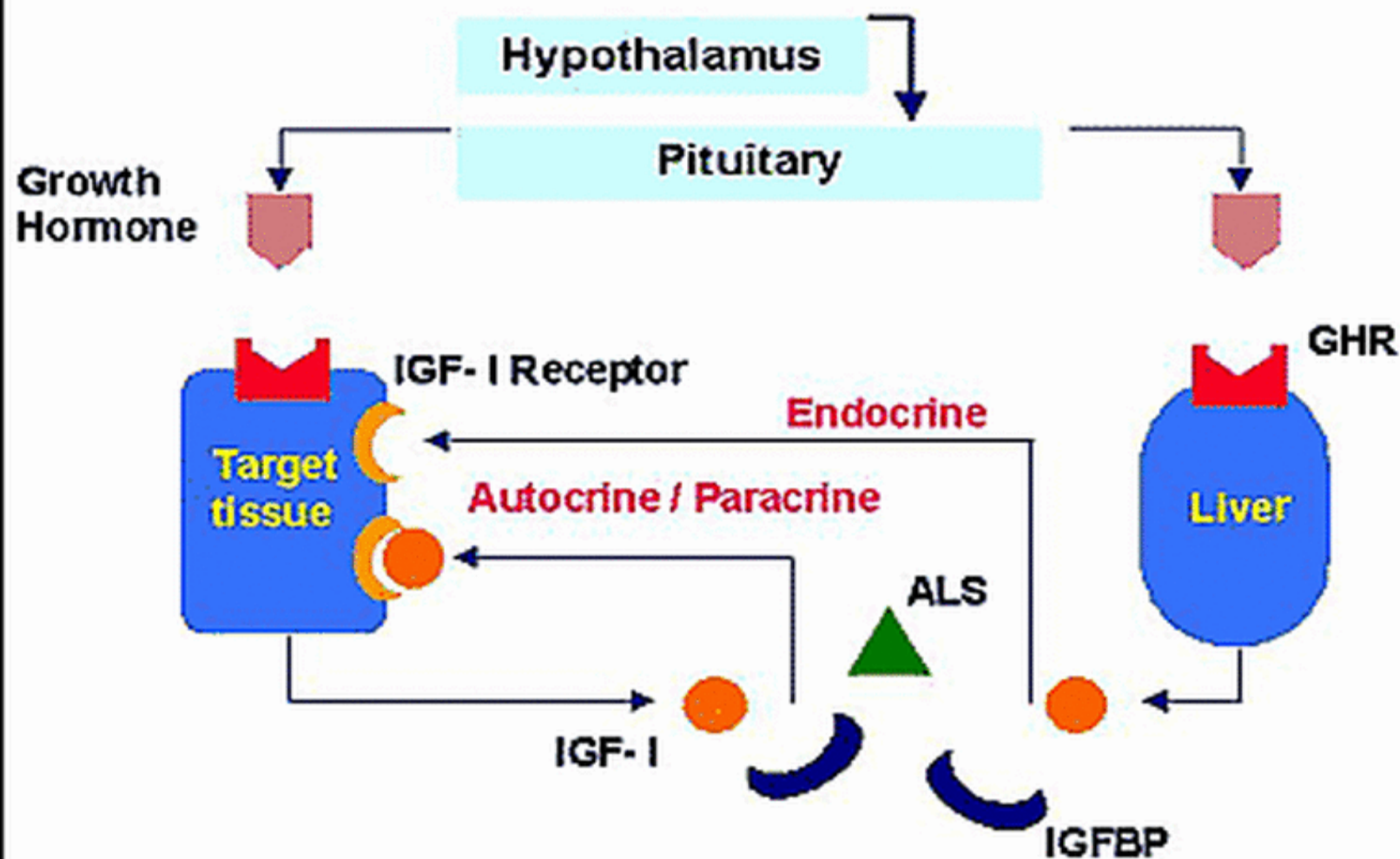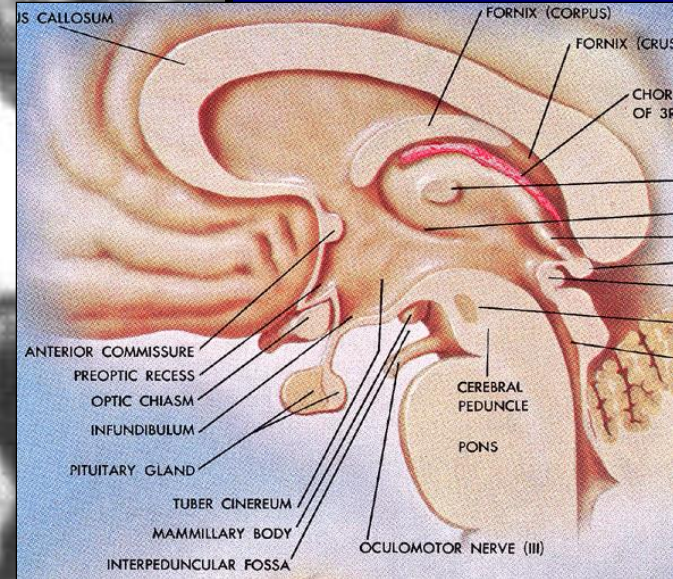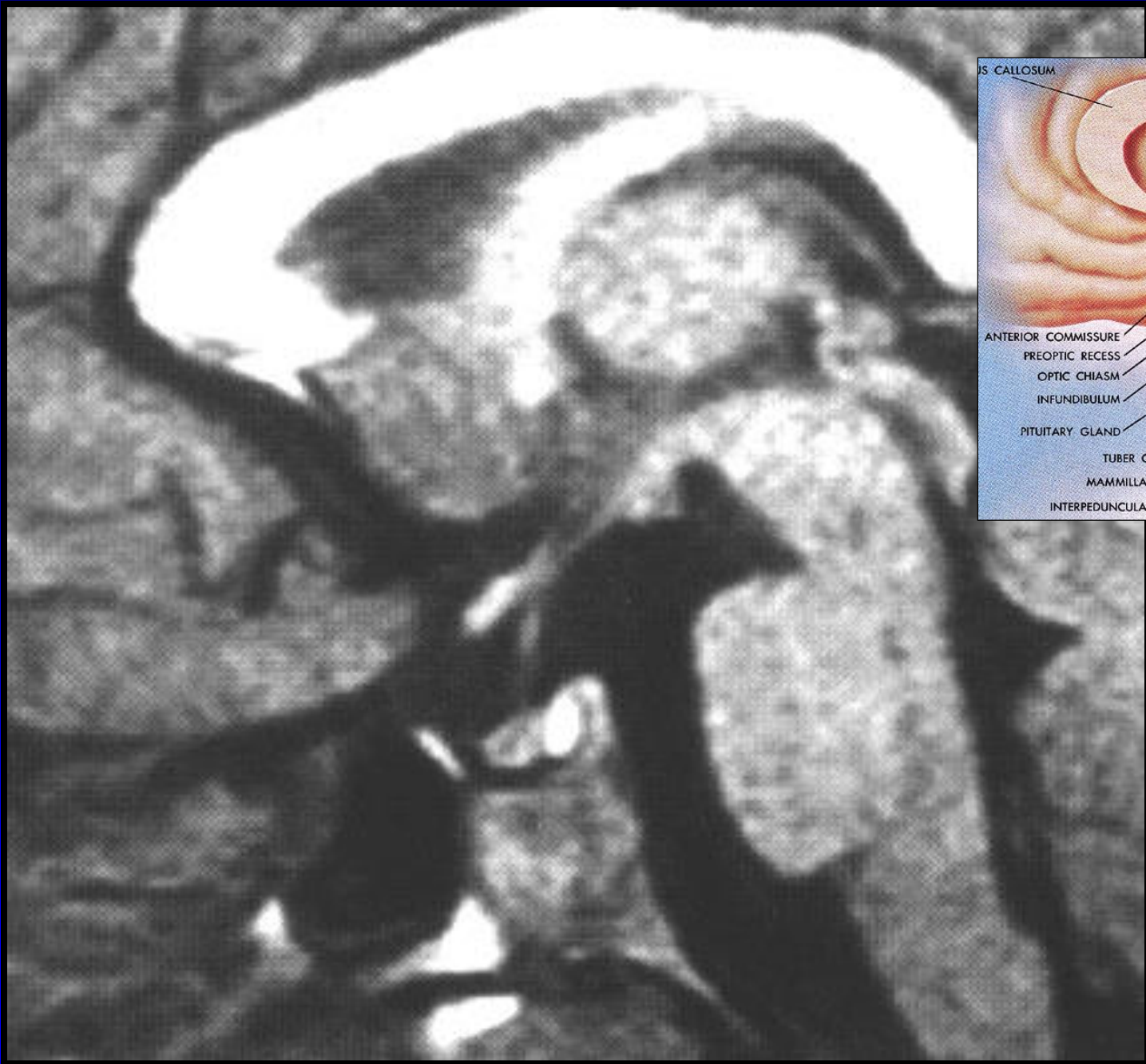
L.J. Abernethy

Classification of growth hormone deficiency

- cause (congenital or acquired)

- nature (true permanent deficiency or functional/temporary insufficiency)

- severity (complete, severe or partial)

- whether isolated or part of multiple anterior pituitary deficiency

Causes of hypopituitarism

CONGENITAL

- Idiopathic
- Genetic defect

  (i) isolated GH deficiency (AR, AD, X-linked)

  (ii) multiple AP deficiency (eg PROP-1)

  (iii) Prader-Willi synd
- Midline defects (e.g. septo-optic dysplasia)

ACQUIRED

- Tumours (e.g. craniopharyngioma, optic glioma)
- Surgery
- Cranial radiotherapy (e.g. for medulloblastoma)
- Granulomatous disease (e.g. Langerhans cell histiocytosis)

Idiopathic short stature

- Definitions vary according to different centres

- One definition is significant short stature (< -2.5 SD) not attributable to normal familial short stature or constitutional delay or to other causes of short stature

- Heterogenous condition which may include partial GH resistance, mild skeletal dysplasias etc.

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 (nice dots!)
- Sitting height and leg length if disproportion suspected
- Plot mid parental height and target range
- Bone age by single observer

Evaluation of the child with short stature - examination

- General appearance and nutrition
- Body proportions
- Dysmorphic features
- Systemic examination including heart
- BP
- Pubertal status
- Fundi

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
- TFT, IGF-1,IGFBP3, prolactin, cortisol
- 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

Low Incidence of Pathology Detection and High Cost of Screening in the Evaluation of Asymptomatic Short Children

Stephanie Sisley, MD[1],*, Marcela Vargas Trujillo, MD[2],**, Jane Khoury, PhD[3], and Philippe Backeljauw, MD[1]

Treatment

- Treat the cause.

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 ??

Treatment of GH deficiency

<u>Recombinant</u> human growth hormone (RHGH)

- Daily bedtime subcutaneous injections of aqueous solutions of biosynthetic (recombinant) GH

- Follow up q 3 months

norditropin® nordilet® 5mg/1.5 ml

mg	Dose	IU
0.07	1	0.21
0.13	2	0.39
0.20	3	0.60
0.27	4	0.81
0.33	5	0.99
0.40	6	1.20
0.47	7	1.41
0.53	8	1.59
0.60	9	1.80
0.67	10	2.01
0.73	11	2.19
0.80	12	2.40
0.87	13	2.61
0.93	14	2.79
1.00	15	3.00
1.07	16	3.21
1.13	17	3.39
1.20	18	3.60
1.27	19	3.81
1.33	20	3.99
1.40	21	4.20
1.47	22	4.41
1.53	23	4.59
1.60	24	4.80
1.67	25	5.01
1.73	26	5.19
1.80	27	5.40
1.87	28	5.61
1.93	29	5.79

norditropin® nordilet® 10mg/1.5 ml

mg	Dose	IU
0.13	1	0.39
0.27	2	0.81
0.40	3	1.20
0.53	4	1.59
0.67	5	2.01
0.80	6	2.40
0.90	7	2.79
1.07	8	3.21
1.20	9	3.60
1.33	10	3.99
1.47	11	4.41
1.60	12	4.80
1.73	13	5.19
1.87	14	5.61
2.00	15	6.00
2.13	16	6.31
2.27	17	6.81
2.40	18	7.20
2.53	19	7.59
2.67	20	8.01
2.80	21	8.40
2.93	22	8.79
3.07	23	9.21
3.20	24	9.60
3.33	25	9.99
3.47	26	10.41
3.60	27	10.80
3.73	28	11.19
3.87	29	11.61

norditropin® nordilet® 15mg/1.5 ml

mg	Dose	IU
0.20	1	0.60
0.40	2	1.20
0.60	3	1.80
0.80	4	2.40
1.00	5	3.00
1.20	6	3.60
1.40	7	4.20
1.60	8	4.80
180	9	5.40
2.00	10	6.00
2.20	11	6.60
2.40	12	7.20
2.60	13	7.80
2.80	14	8.40
3.00	15	9.00
3.20	16	9.60
3.40	17	10.20
3.60	18	10.80
3.80	19	11.40
4.00	20	12.00
4.20	21	12.60
4.40	22	13.20
4.60	23	13.80
4.80	24	14.40
5.00	25	15.00
5.20	26	15.60
5.40	27	16.20
5.60	28	16.80
5.80	29	17.40

Clicks into mg

Dose in mg

Differs according to the indication of growth hormone treatment

Side Effects of GH Treatment

- Metabolic effects (hyperglycemia)
- Antibodies to growth hormone
- Progression of pre-existing scoliosis
- Slipped capital femoral epiphysis
- Pseudotumor cerebri

- Transient gynecomastia
- Increased growth and pigmentation of nevi
- Carpal tunnel syndrome
- Edema / Arthralgia
- Hypothyroidism
- Second neoplasms
Meningioma
- Pancreatitis

Diagnostic Approach in Children with Short Stature

Wilma Oostdijk[a]   Floor K. Grote[a]   Sabine M.P.F. de Muinck Keizer-Schrama[b]
Jan M. Wit[a]

[a]Department of Pediatrics, Leiden University Medical Center, Leiden, and [b]Department of Pediatrics, Erasmus MC – Sophia Children's Hospital, Rotterdam, The Netherlands

Supplement

Etiologies and Early Diagnosis of Short Stature and Growth Failure in Children and Adolescents

Alan D. Rogol MD, PhD [1] ⊠, Gregory F. Hayden MD [2]

Thank You