

PEDIATRICS

RR-8.0

Contents

General Paediatrics : Growth	1
General Paediatrics : Development and Nutrition	8
General Paediatrics : Genetics, Infections and Metabolic Disorders	16
Neonatology	25
Systemic Paediatrics : Neurology	34
Systemic Paediatrics : Pulmonology and Cardiology	41
Systemic Paediatrics : Gastroenterology, Nephrology and Endocrinology	50

GENERAL PAEDIATRICS : GROWTH

----- Active space -----

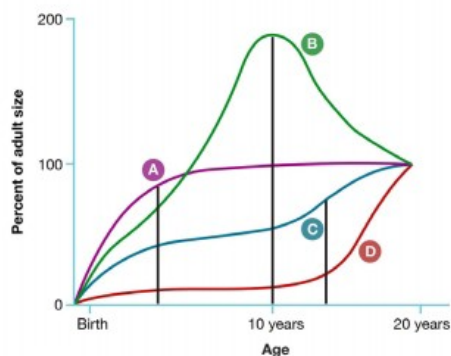
Normal Growth

00:00:34

Growth Phases :

	Period of growth		Days of life
Prenatal (Before birth)	Ovum		0 - 14 days of gestation
	Embryo		14 days - 8 weeks of gestation
	Fetus		9 weeks - Birth
Perinatal			22 weeks of gestation - 7 days after birth
Postnatal	Newborn	Early	0 - 7 days after birth
		Late	8 - 28 days after birth
	Infancy		Till 1 yr
	Toddler		1 - 3 yrs
	Preschool		3 - 6 yrs
	School age		6 - 12 yrs
	Adolescence	Early	10 - 13 yrs
		Mid	14 - 16 yrs
		Late	17 - 19 yrs

Growth Patterns :



Scammon growth curve

- A** : Brain growth (max at 2 yrs).
Parabolic.
- B** : Lymphoid growth (E.g : Tonsils).
>100% adult size.
- C** : Somatic growth.
Growth spurt after 12 yrs.
- D** : Gonadal growth

Anthropometry

00:07:06

Weight :

- Average birth weight : 2.9 Kg.
- Low birth weight (LBW) : < 2.5 Kg.

----- Active space -----

Weight loss after birth : D/t loss of excess extracellular fluid

	Weight loss (In % in 1 st week)	Regains weight by
Term	10	10 th day
Preterm	15	15 th day

Weight gain :

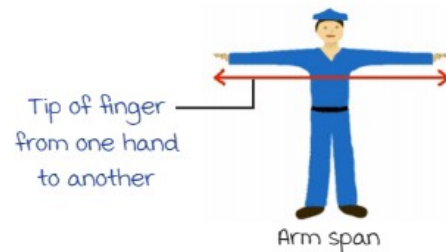
Age	↑ in weight
Till 3 months	30 g/day
Till 1 yr	400 g/month
1 - 7 yrs	2 kg/yr
>7 yrs	3 kg/yr

- Doubles : 5 - 6 months
 - Triples : 1 year
 - Quadruples : 2 years
- } After birth.

Height/Length (< 2 years) :

Arm span : Height equivalent in older children.

Comparison	Age
Arm span (< 2.5 cm) < Length	Birth
Arm span = Length	11 yrs
Arm span (> 1 cm) > Length	> 11 yrs



Increase in height :

Age	Height
At birth	50 cm
At 1 year	75 cm
At 2 years	87.5 cm (Height = Half of adult height at 18 to 24 months)
> 2 yrs - Till puberty	↑ of 6 cm/yr
Growth spurt at puberty	<ul style="list-style-type: none"> • Boys : 20 - 30 cm • Girls : 16 - 28 cm
Adult	160 - 170 cm

- Doubles (100 cm) : 4 years.
- Triples (150 cm) : 12 years.

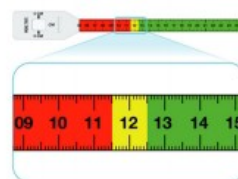
Circumference :

mid arm circumference (MAC) :

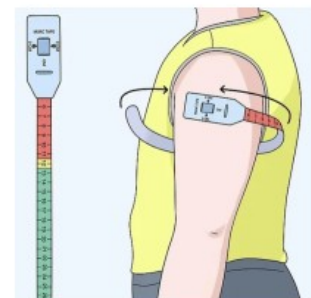
- Nutritional status assessment.
- Age : 1 - 5 years.

Shakir's tape :

- Normal : >12.5 cm
- malnutrition : 11.5 - 12.5 cm
- Severe malnutrition : <11.5 cm



Shakir's tape



Head circumference :

Brain growth assessment.

----- Active space -----

Age	Head circumference & growth rate
At birth	33 - 35 cm (34 cm)
First 3 months	2 cm/month (40 cm)
3 - 6 months	1 cm/month (43 cm)
6 months - 1 yr	0.5 cm/month (46 cm)
1 - 2 yr	2 cm/year (48 cm)
At 12 yrs	Adult value : 52 cm

WHO Growth Chart, Dentition & Puberty

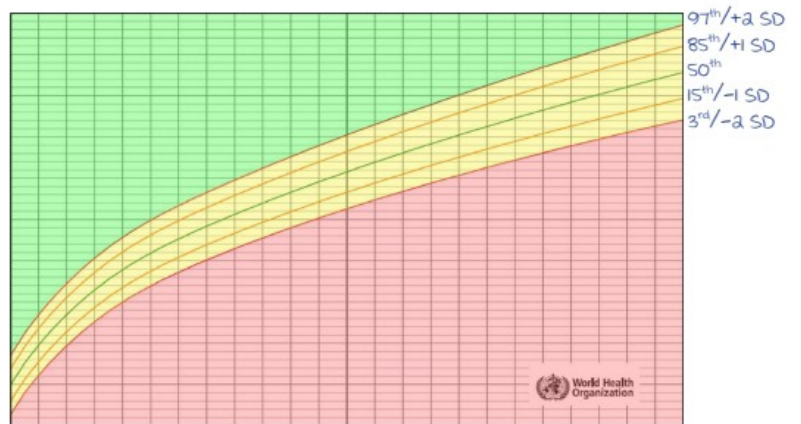
00:24:25

WHO Growth Chart :

Colours :

- Pink : Girls.
- Blue : Boys.

- Tall stature (>97th centile/>+2 SD).
- Normal height.
- Short stature (<3rd centile/<-2 SD).



Dentition :

	1° dentition (Temporary)	2° dentition (Permanent)
Total no. of teeth	20	32
1 st tooth to erupt	Lower central incisor	1 st molar
Time of eruption	6 months	6 yrs

mixed dentition :

- Both temporary & permanent teeth.
- Age : 6 - 12 yrs.

Supernumerary teeth :

- Additional teeth.
- m/c : In b/w 2 central incisors.

Delayed dentition : Non-appearance of teeth by 13 months.

Cause :

- Idiopathic (m/c).
- malnutrition.
- Genetic syndromes (Down, Turner).
- Hormonal deficiency (GH, thyroid).

----- Active space -----

Puberty Changes :

Females (8 - 13 years) :



males (9 - 14 years) :



Sexual maturity rating (SMR) : Tanner's staging.

Stage 1 : Prepubertal.

Stage 3 : Growth spurt in girls.

Stage 4 : Growth spurt in boys.

Stage 5 : Adult-like.

Short Stature

00:36:21

Height/age : < 3rd percentile or < -2 SD.**Causes :**

- Normal variants (m/c than pathological causes).
 - Constitutional delay : m/c cause of short stature & delayed puberty.
 - Familial short stature.

mid parental height : Child's genetic potential.

$$\frac{\text{Father's height} + \text{mother's height}}{2} \pm 6.5$$

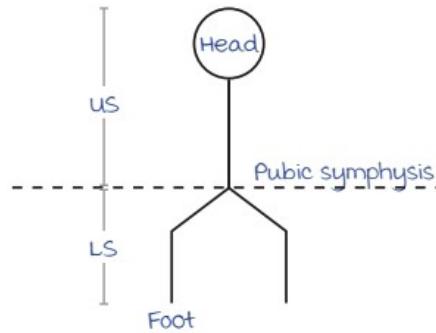
- For boys : + 6.5
- For girls : - 6.5

Constitutional delay vs familial short stature :

	Constitutional delay	Familial short stature
Adult height	Normal	Short
Puberty	Delayed	Normal for age
Bone age		
Parent's height	Normal	Short

US:LS Ratio :

Age	US : LS
At birth	1.7 : 1
3 yrs	1.3 : 1
10 yrs	1 : 1
>10 yrs	0.9 : 1



US : upper segment
LS : Lower segment

----- Active space -----

Approach to short stature :

US : LS ratio is normal for age : Proportionate short stature	US : LS ratio is abnormal for age : Disproportionate short stature	
	Short trunk (US)	Short limbs (LS)
<ul style="list-style-type: none"> • Normal variants • Chronic malnutrition (Stunting) • GH deficiency 	<ul style="list-style-type: none"> • Spondylo-epiphyseal dysplasia • Hemivertebrae • mucopolysaccharidosis (MPS) • Pott's disease : TB spine 	<ul style="list-style-type: none"> • Achondroplasia • Rickets • Congenital hypothyroidism

Anomalies of Head Size & Growth

00:45:55

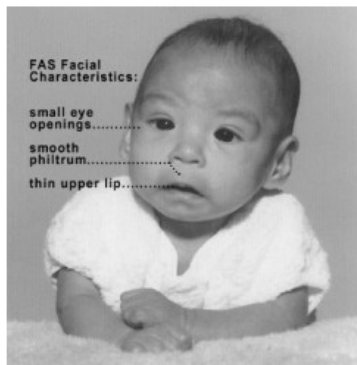
MICROCEPHALY

Causes :

1°/genetic microcephaly	2° microcephaly
<ul style="list-style-type: none"> • Developmental anomalies • Genetic defects (Trisomies) 	<ul style="list-style-type: none"> • Prenatal : maternal TORCH infections, teratogenicity • Perinatal : Birth asphyxia • Postnatal : Infections, trauma

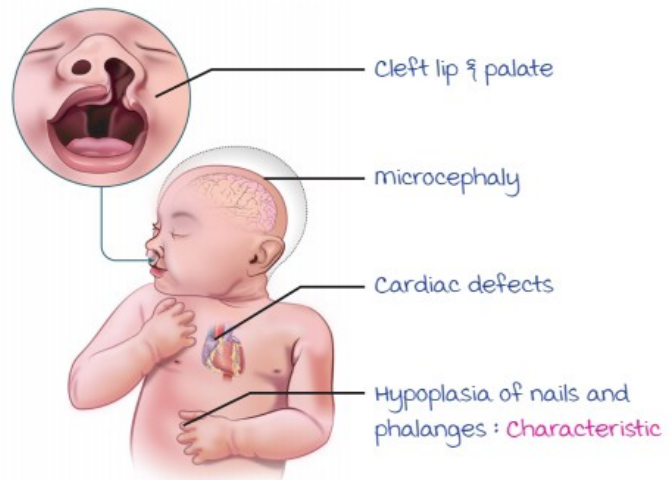
Fetal Alcohol Syndrome :

VSD : m/c heart disorder.



Fetal Hydantoin Syndrome :

VSD : m/c heart disorder.



----- Active space -----

Rett's Syndrome :

- X-linked dominant disorder.
 - Defect : **MECP2** defect.
 - Clinical features :
 - microcephaly
 - Developmental regression
 - **Stereotypes** : Hand wringing movements
 - Speech defects, ataxia
- } Symptoms appear after 1 yr
(Normal at birth).

MACROCEPHALY

Causes :

- Hydrocephalus : \uparrow CSF.
 - MPS
 - Leukodystrophies :
 - Alexander disease
 - Canavan disease
 - Thalassemia
 - Osteogenesis imperfecta
- } megalencephaly.
- } \uparrow Bony component.

Congenital Hydrocephalus :

Normal CSF :

- Adults : 150 mL.
- Infants : 50 mL.

Etiopathogenesis :

Anomalies \rightarrow Obstruction in CSF pathway.**Aqueductal stenosis** (m/c) : Narrowing b/w 3rd & 4th ventricle.

Clinical features :

1. macrocephaly :
 >2 cm/month \uparrow in head circumference.
2. Bulging fontanelle.
3. Congested/prominent scalp veins.
4. Sunset appearance : visible upper sclera
(D/t downward rotation of eyeball).
5. Cracked pot resonance : D/t \uparrow pressure.

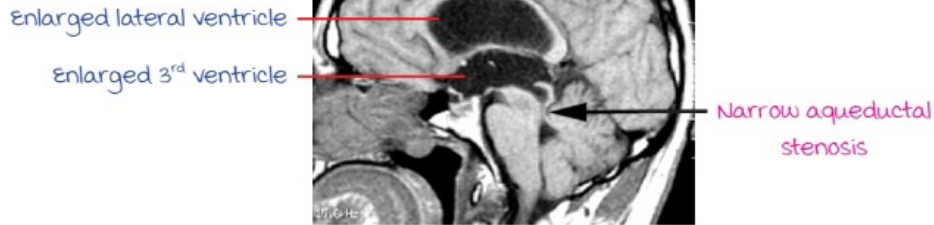


Features of macrocephaly

----- Active space -----

Investigation :

- Postnatal MRI :



- Intranatal USG :
 - Anomalies detected at 2nd trimester.
 - Fetal ventriculomegaly (m/c cause : Aqueductal stenosis).

management :

- ventriculo-peritoneal shunt (m/c) : CSF shunting.
- Endoscopic 3rd ventriculostomy.

Disorders of Puberty

01:04:04

Abnormal onset of 2^o sexual characteristics :

	males	Females
Delayed puberty	> 14 yrs	> 13 yrs
Precocious puberty	< 9.5 yrs	< 8 yrs

Delayed Puberty :

Cause :

Constitutional delay (m/c).

Central causes	Peripheral causes
<ul style="list-style-type: none"> • CNS abnormalities (Pituitary/hypothalamus) : Tumours, trauma, infiltration in pituitary. • Syndromes : Kallman syndrome (Anosmia). 	<ul style="list-style-type: none"> • Genetics (Gonadal defects) : <ul style="list-style-type: none"> - Turner : Streak ovaries. - Klinefelter } Cryptorchidism. - Noonan }

Precocious Puberty :

Central causes (Pituitary, hypothalamus) : Gonadotropin dependent (↑ LH, FSH)	Peripheral causes : Gonadotropin independent
<ul style="list-style-type: none"> • Tumours : <ul style="list-style-type: none"> - Hypothalamic hamartoma (m/c overall & m/c in boys). - Glioma. • Idiopathic (m/c in girls). 	<ul style="list-style-type: none"> • Functional testicular tumors (↑ testosterone) : Seminoma, germinoma. • Autonomously functioning ovarian cyst : McCune Albright syndrome. <ul style="list-style-type: none"> - Precocious puberty (D/t ↑ estrogen). - Polyostotic fibrous dysplasia. - Cafe-au-lait macules.