Child Development

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Objectives

➤To outline the link between embryology and clinical practice...examples from 2 systems

- >To define child development and its assessment
- >To outline the factors affecting development
- ➤To use the framework model to rehearse developmental assessment
- >To look at abnormal development



>The overall context

Teratogenesis – causes and effect

Recap embryology of nervous and urogenital systems

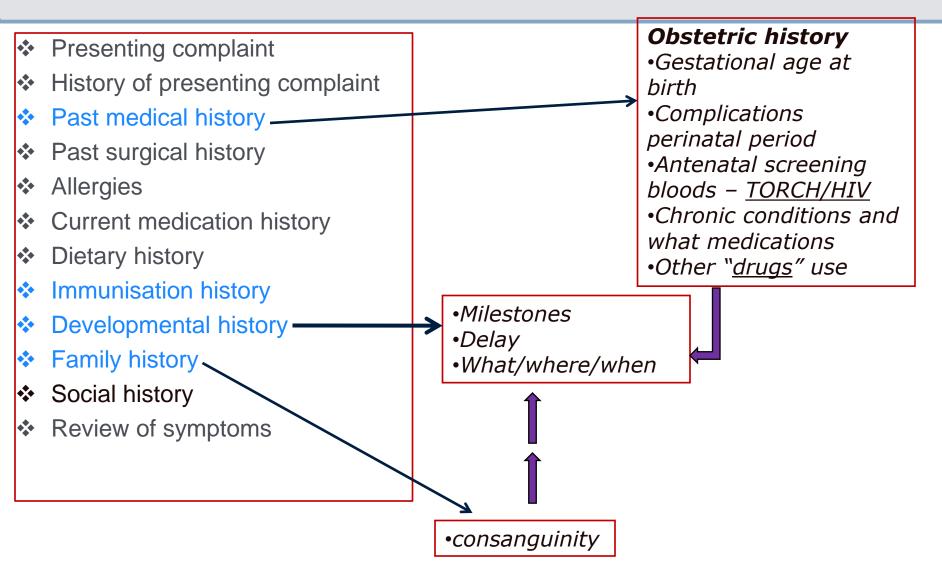
Specific examples outlining prevention and treatment

Renal development and examples

- Genital development and examples
- Developmental milestones
- >Abnormal development

The overall context

The paediatric history



Causes-teratogenesis

- Infectious agents-TORCH, HIV, syphilis, varicella, hepatitis
 Physical agents-X-rays and hyperthermia
- Chemical agents-phenytoin, valproate, lithium, amphetamines, ACE inhibitors, alcohol, cocaine
- ≻Hormones-androgens
- Maternal diabetes

T=Toxoplasma O=Other R=Rubella C=Cytomegalovirus H=Herpes



≻Aplasia → Absent

≻Hypoplasia → Underdeveloped

≻Dysplasia → Abnormal shape

≻Hyperplasia — Too big

➢ Ectopia → Abnormal position

The urogenital system

The renal system

Abnormal function
Abnormal position
Abnormal drainage

The genitalia

≻Ambiguity

Recap.....renal development

≻Mesonephric duct→ureteric bud

>Ureteric bud dilates → renal pelvis, and splits → major calyces

≻Each calycx→consecutive divisions→minor calyces and renal pelvis

>Thus, ureteric bud \rightarrow ureter, renal pelvis, calyces and collecting tubules

Abnormal function

Ducts surrounded by undifferentiated tissue

➤Ureteric buds fail to branch

≻Nephrons do not develop

>Failure to differentiate \rightarrow poor renal development and function

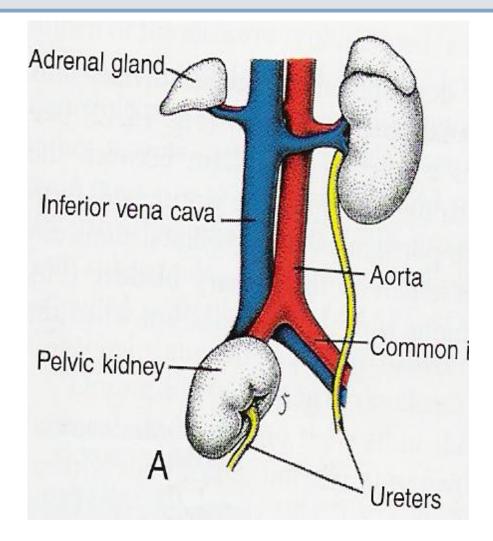
>This leads to reduced fetal urine \rightarrow amniotic fluid \rightarrow hypoplastic lungs

Abnormal function

- •Potter's syndrome
 - ➢Bilateral renal agenesis
 - ≻Anuria
 - Oligohydramnios
 - Hypoplastic lungs
 - ≻Other defects
 - Characteristic features
 - ≻Lethal

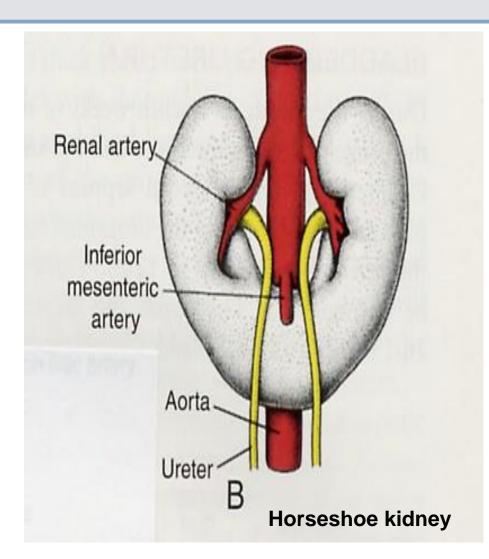


Abnormal position



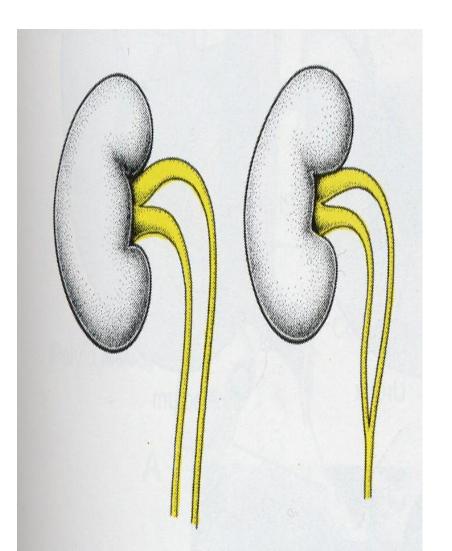
Abnormal formation

•Failure of ascent of kidneys→fusion of lower poles→horseshoe kidney



Abnormal drainage

- •Ureteric buds split
- •Splitting partial or complete
- •Extra ureter may insert abnormally
- •May lead to (recurrent) urine infections and reflux→renal scarring→hypertension



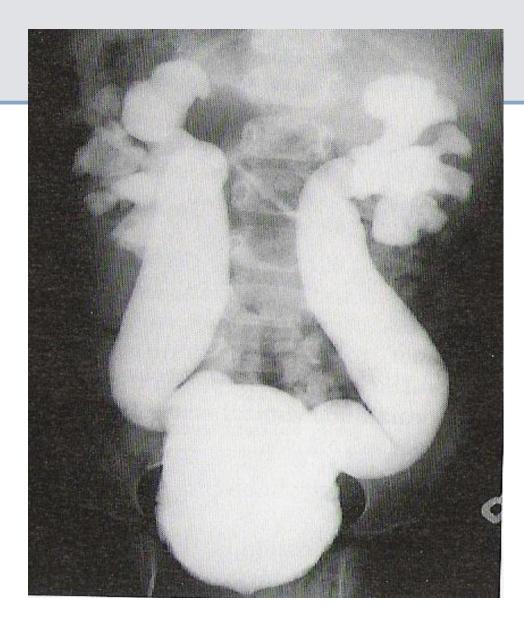
Gross dilatation of

pelvic calcyces

•ureters

•bladder

as a result of severe vesicoureteric reflux



The consequences

- Vesicoureteric reflux
- •Recurrent urinary tract infections
- Renal scarring
 Renal hypertension
 Renal failure

Recommended imaging schedule for infants younger than 6 months

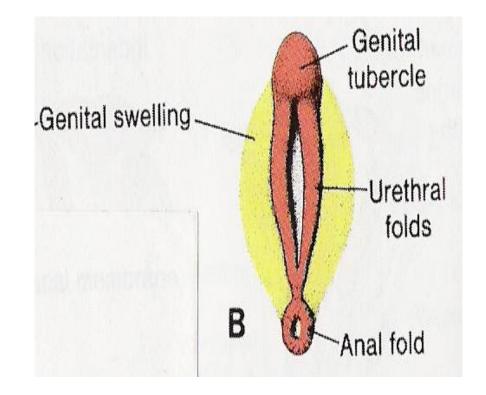
Test	Responds well to treatment within 48 hours	Atypical UTI ^a	Recurrent UTI ^a
Ultrasound during the acute infection	No	Yes ^c	Yes
Ultrasound within 6 weeks	Yes ^b	No	No
DMSA 4–6 months following the acute infection	No	Yes	Yes
MCUG	No	Yes	Yes

Recommended imaging schedule for infants and children 6 months or older but younger than 3 years

Test	Responds well to treatment within 48 hours	Atypical UTI ^a	Recurrent UTI ^a
Ultrasound during the acute infection	No	Yes ^c	No
Ultrasound within 6 weeks	No	No	Yes
DMSA 4-6 months following the acute infection	No	Yes	Yes
MCUG	No	No ^b	No ^b

Recap....development of genitalia

- Indifferent at early stage
- In males, androgens are required → elongation of the genital tubercle to become the phallus
- •Urethral folds close over urethral plate to become penile urethra
- •Absence of SRY→ovary
- Presence of SRY→testis

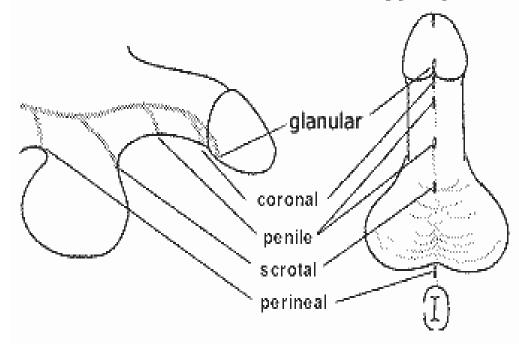


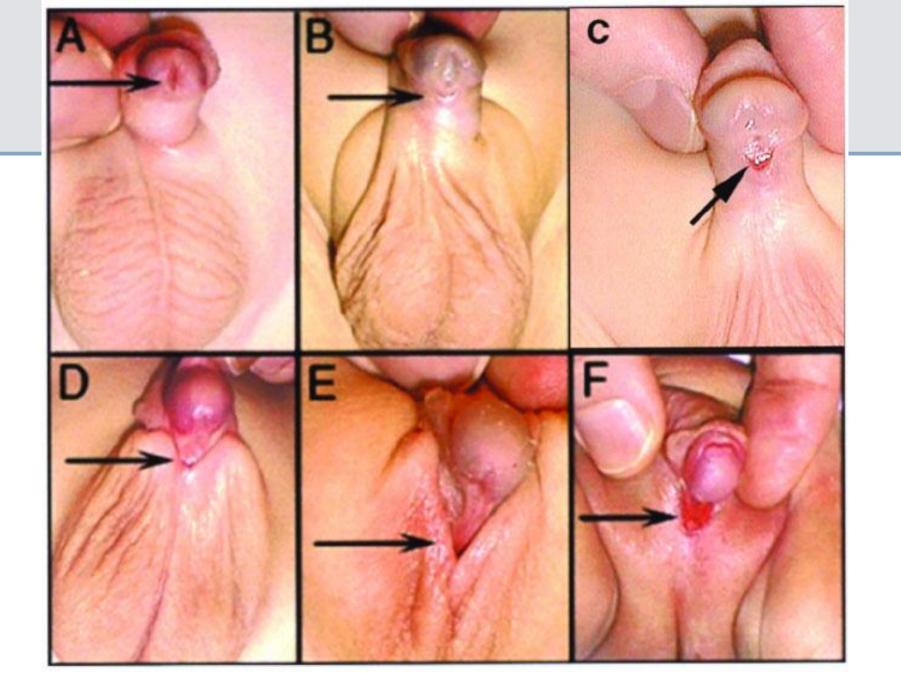
Hypospadias

Incomplete fusion of the urethral folds

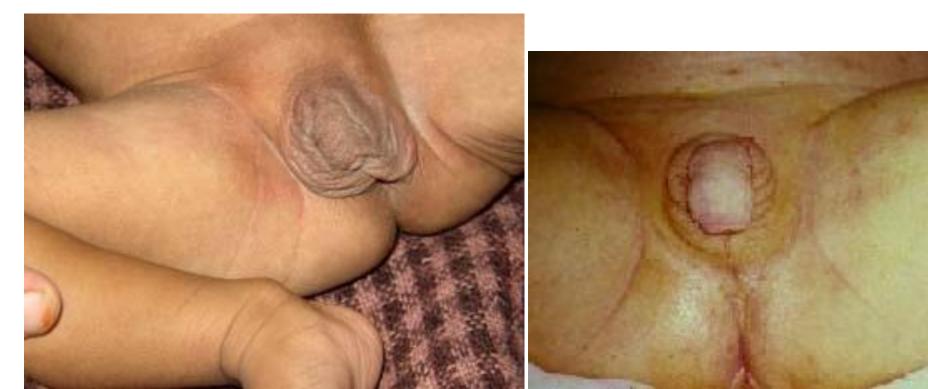
•Openings of the urethra occur along the inferior aspect of the penis

Location of Urethral Meatus in Hypospadias







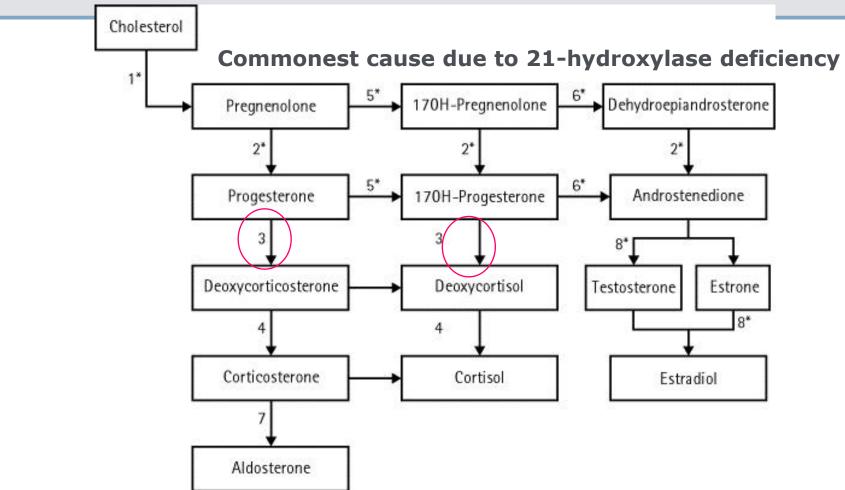


Ambiguous genitalia

Numerous causes

- Point mutations or deletions of the SRY gene
- Biochemical abnormalities resulting in imbalance in steroid hormones
- End-organ or hormone receptor resistance

Biochemical abnormalities



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Hormonal and clinical consequences

Decrease aldosterone

Decrease cortisol

Increase oestradiol

•salt losing crisis and shock in males

over-virilisation in females
 >psychological implications of appearance and future reproductive potential

•Other causes may result in hypertension

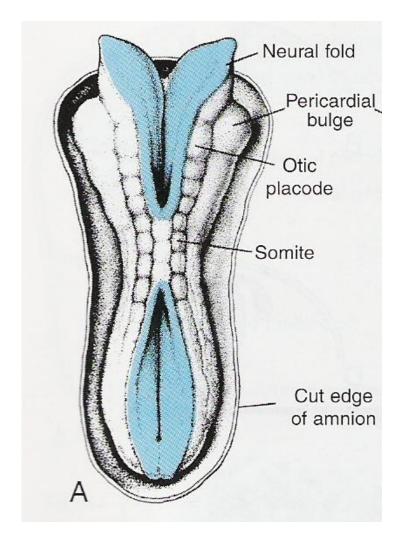
 Management ⇒ mineralocorticoids ± glucocorticoids ± surgery

CNS embryology

•Begins in 3rd week

•Neural plate elevates to form neural folds

•Neural folds fuse to form neural tube

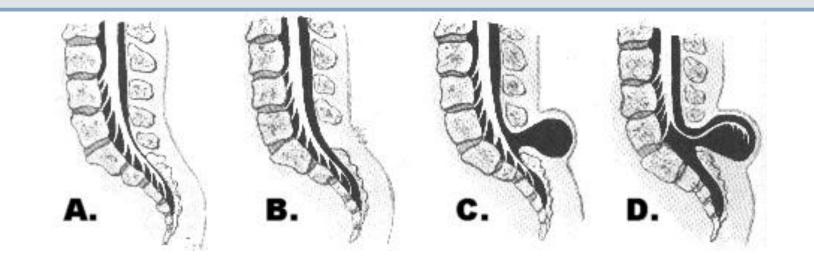


When things go wrong:



Failure of fusion→neural tube defects
Anencephaly-failure of rostral fusion
Spina bifida-failure of caudal fusion
AEDs and genetics are risk factors

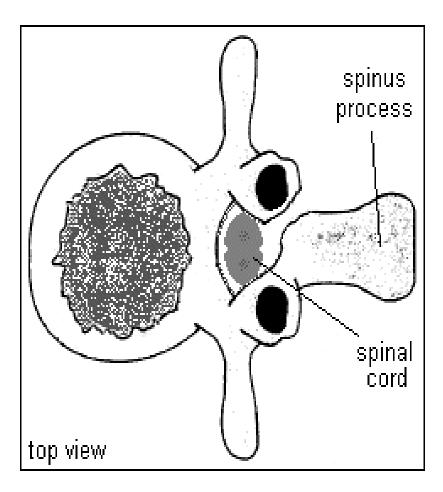
Spina bifida



- A: Normal spine
- B: Spina bifida occulta
- C: Meningocele } Spina bifida
- D: Myelomenogocele } cystica

Spina bifida occulta

- •Defect in vertebral arches covered by skin
- •Does not usually involve underlying neural tissue
- •Occurs in lumbosacral region
- •Usually marked by a patch of hair or dimples overlying the defect





Sacral dimples



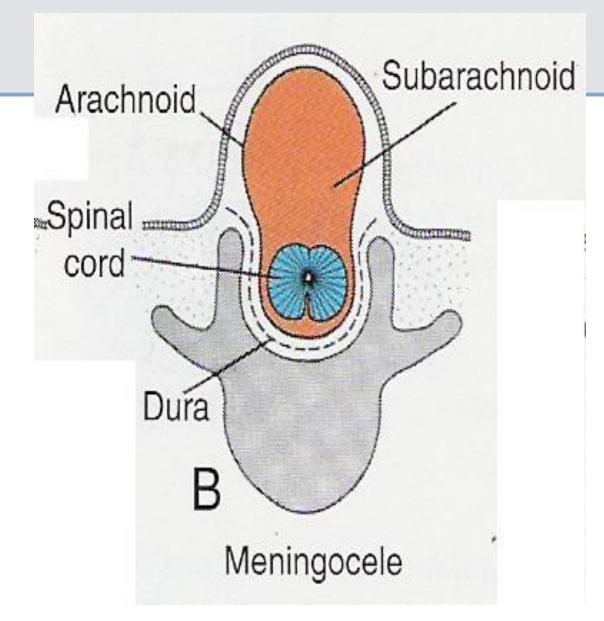
Spina bifida cystica

Severe defect

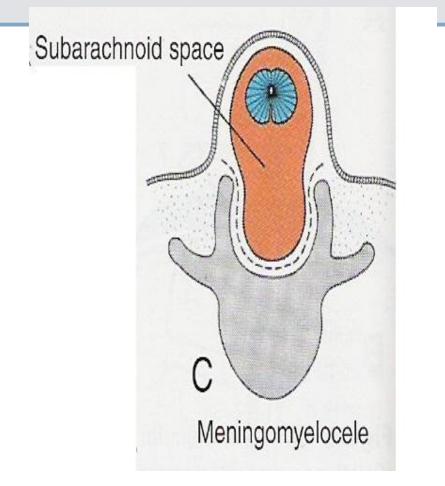
Meninges and/or spinal cord protrude through vertebral arch defect to form cyst-like sac

➤Most lie in lumbosacral region

➢ Result in neurological deficits









Complications of spina bifida cystica

Neurogenic bowel • incontinence Neurogenic bladder-

- recurrent infections
- incontinence
 Lower limb paralysis

Fractures

Joint contractures

Developmental deformities

-learning impairment

-hydrocephalus

(2° to meningitis)

Gastrointestinal Hearing Respiratory Gastro-oesophageal reflux Respiratory infections Conductive or sensorineural hearing impairment Oromotor incoordination Aspiration of food or saliva Chronic lung disease Constipation Sleep apnoea Vision Neurological Squint Urogenital Epilepsy Impaired visual acuity Urinary tract infection Visual field deficits Delay in establishing continence Cerebral palsy Unstable bladder Orthopaedic Vesico-ureteric reflux Nutrition Hip subluxation/dislocation Neuropathic bowel and bladder Poor weight gain Fixed joint contractures Failure to thrive Dynamic muscle contractures Painful muscle spasm Spinal deformity Behaviour Osteoporosis/fractures Organic or reactive **Common medical problems** Sibling behaviour Specialist health visitor Parental distress Help coordinate multidisciplinary and multiagency care Advice on development of play Feeding Language development or local authority schemes Speech development e.g. Portage Dietician Advice on feeding and nutrition voice synthesizers Social worker/ Social services Advice on benefits: disability,

mobility, housing, respite care, voluntary support agencies Day nursery placements Advocate for child and family Register of children with special needs

Psychologist (clinical and educational)

Cognitive testing Behaviour management Educational advice

Paediatrician

Assessment, investigation and diagnosis Continuing medical management Coordination of input from therapists and other agencies - health, social services, education

Child Development Service

Aspiration pneumonia

Microcephaly/ hydrocephalus

Speech and language therapist

AAC (augmentative and alternative communication) aids e.g. Makaton sign language, Bliss symbol boards,

Occupational therapist

Eye-hand coordination ADL (activities of daily living) feeding, washing, toileting, dressing, writing Seating Housing adaptations

Physiotherapist

Balance and mobility Postural maintenance Prevention of joint contractures, spinal deformity Mobility aids, orthoses

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Diagnosis, prevention & treatment

Diagnosis

> Prenatally by ultrasound > α -fetoprotein (AFP) in maternal serum and amniotic fluid

Prevention

➢Folic acid from 2 months prior to conception reduces risk

Treatment

Surgery in utero at 28 weeks gestationIndividual management of complications



>The overall context

➤Teratogenesis – causes and effect

Recap embryology of nervous and urogenital system

Illustrated how embryology can aid understanding of clinical scenario

CHILD DEVELOPMENT – DEVELOPMENTAL FRAMEWORK

Outline

- ➤ Where does it all fit in?
- Definition of development
- Domains of development
 - Gross motor and posture
 - Fine motor and vision
 - Language and hearing
 - Social and behaviour
- Assessment of development

Where does it all fit in?

Cellular and molecular biology ↓ Fertilisation and embryogenesis ↓ Growth and <u>development</u>

What is development?

Global impression of a child which encompasses growth, increase in understanding, acquisition of new skills and more sophisticated responses and behaviour

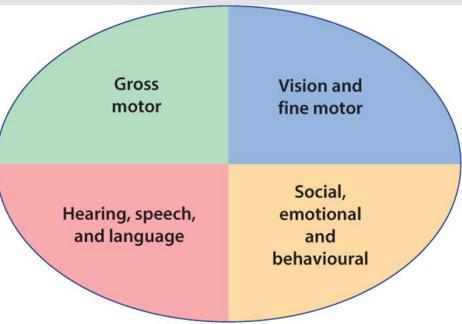
It serves to endow the child with increasingly complex skills in order to function in society

What influences development

- >genetic factors
- >biochemical
- environmental influences
 internal milieu
 maternal illness
 teratogens

Domains of development

- Gross motor and posture
- Fine motor and vision
- Language and hearing
- Social and behaviour



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Developmental norms are called <u>milestones</u>-they describe recognised patterns of development that children are expected to follow

Gross motor and posture

Looks at posture and execution of large movements

- Standing
- Walking
- Running
- Kicking a ball
- Climbing stairs
- Peddling a tricycle



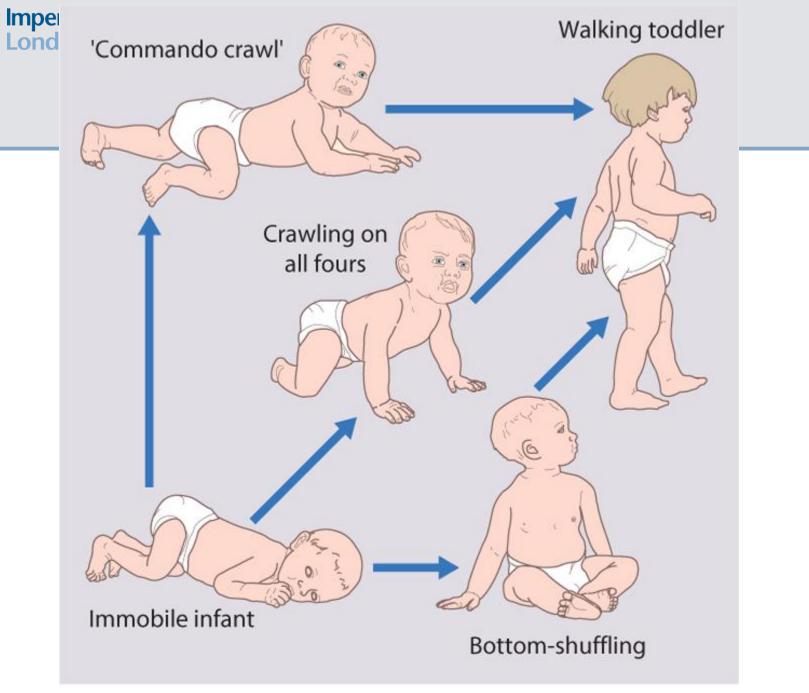
Gross motor

Develops in a cephalo-caudal direction

➤ Lying mobile ⇒ standing/walking

 \succ Flexion \Rightarrow extension

ightarrow Rolling \Rightarrow sitting \Rightarrow crawling \Rightarrow walking



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Primitive reflexes

Are protective and have a survival value

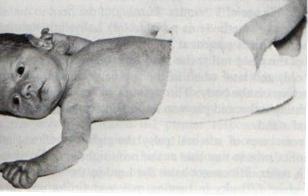
Promote proper orientation

Promote postural support and balance

Primitive reflexes

Stepping







Moro

These should disappear by 4-6 months

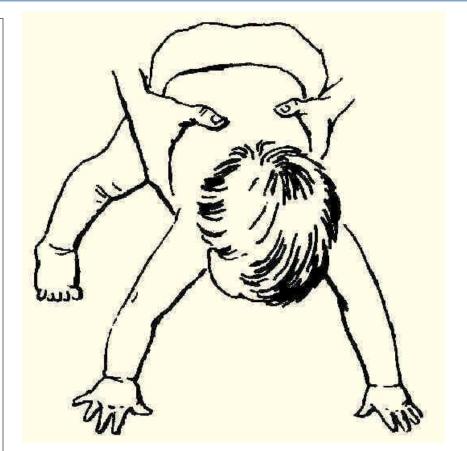
Grasp





Protective or righting responses

Have to develop before can attain motor development Head righting Downward parachute by 4-6 months Forward parachute by 7-9 months



Forward parachute



Gross motor development (median ages)



Limbs flexed, symmetrical postures

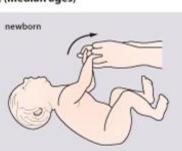




Raises head to 45°



12 months



Marked head lag on pulling up





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Fine motor and vision

Looks at hand function and hand-eye coordination

- ➢Also tends to give some information of cognitive function
 - Holding objects
 - Picking up objects
 - Pointing
 - Waving
 - Throwing/catching

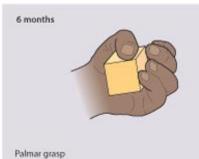




Vision and fine motor (median ages)

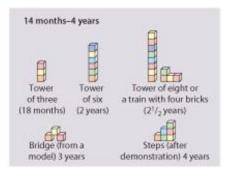


Newborn – follows face in midline. Follows moving object or face by turning the head (illustrated).



10 months

Mature pincer grip

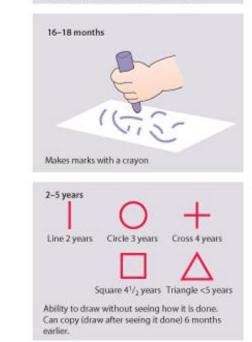


4 months

Reaches out for toys

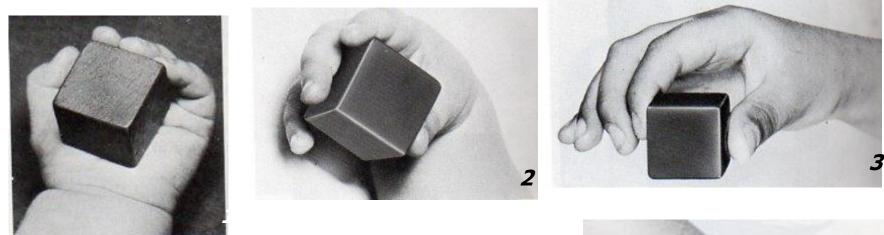


Transfers toys from one hand to another

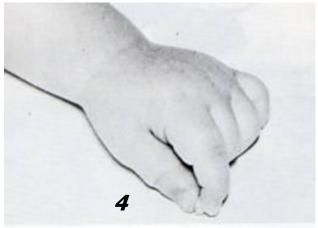


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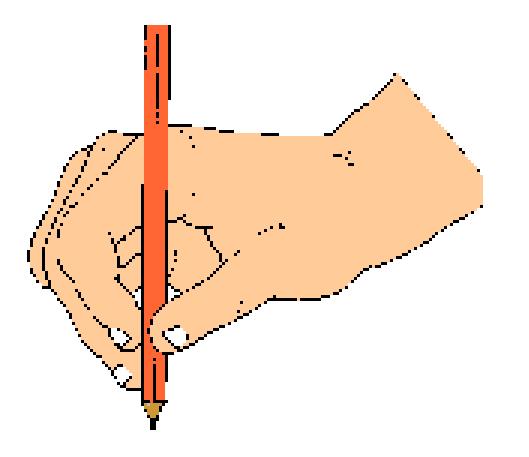
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Palmar grasp = 6 m
 Intermediate grasp = 8 m
 Mature grasp = 12 m
 Pincer grasp = 12 m



Tripod grasp

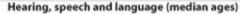


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Language and hearing

Normal hearing is needed for normal speech

- Babbling
- Words
- Sentences
- Social communication





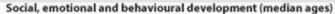
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Social, behaviour and play

- Looks at interaction with others and self care skills
- Stranger awareness
 Play
 Feeding
 Toileting
 Dressing







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Parallel play. Interactive play evolving. Takes turn

clothing

g

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Age	Gross motor control	Vision and fine motor	Language and hearing	Social and daily living skills
2-4 months	Head steady in sitting	Follows object through 180°	Squeals with pleasure	Smiles
5-8 months	Sits without support	Passes cube hand to hand	Turns to soft voice Baba/Gaga babble (up to 10 months)	Feeds self biscuit
9-14 months	Stands with support	Neat pincer grasp of raisin	Mama or Dada specifically	Indicates needs by gesture
12–16 months	Walks well alone	Stack of two cubes (up to 21 months)	Three words (up to 21 months)	Drinks from a cup
15-24 months	Walks up steps	Scribbles spontaneously	Points to one body part	Removes garment
21-36 months	Jumps on the spot	Draws vertical line in imitation	Uses plurals and phrases	Puts on clothing Plays tag with other children
3-4 ¹ / ₂ years	Balances on one foot for 5 s	Copies a ladder Draws a face	Understands cold, tired and hungry Asks 'Wh' questions	Separates from mother

Developmental Milestones Table

AGE	Gross Motor	Vision & Fine Motor	Speech & Language	Personal Social
6 weeks	Pull to sit: Head lag and rounded back. Ventral Suspension: Head momentarily in same plane as body. Prone: Pelvis high but knees no longer under abdomen. Chin raised occasionally.	Fixates and follows to 90 °	Vocalising by 8/52 Quiets to sound. Startles to sound.	Smiles responsively.
3 months	 Pull to sit: Only slight head lag. Head occasionally bobs forward. Ventral Suspension: Head above plane of body. Prone: Pelvis flat. Lifts head up 45° - 90°. 	Hand regard. Follows object from side to side (180°) Hands held loosely. Grasp object placed in hand. Not reaching out.	Squeals with delight. Turns head to sound.	Laughs.
5 months	Pull to sit: NO head lag and sits with straight back. Lying supine : Feet to mouth.	Reaches for objects. Plays with toes.		Mouthing.
6 months	Pulls to sit: Lifts head off couch in anticipation. Sits with support. Bears full weight on legs. Prone: Supports weight on hands with chest and upper abdomen off couch. Rolls prone to supine.	Palmar grasp of cube with ulnar approach. Moves head and eyes in all directions. No squint (after 4 months).		
7 months	Sits with hands on couch for support. Rolls from supine to prone.	Feeds self with biscuits. Transfers objects form hand to hand. Rakes at pea.	Babbling in single syllables. Babbling in combined syllables at 8 months. Distraction Test.	Stranger anxiety.

AGE	Gross Motor	Vision & Fine Motor	Speech & Language	Personal Social
9 months	Sits steadily. Leans forward but cannot pivot. Stands holding on. Pulls self to stand.	Inferior pincer grasp. (Scissors grasp).	Localises sound at 3 feet, above and below the ear level.	Feeds with spoon occasionally. Looks for fallen toys. Understands "NO!"
10 months	Crawls on abdomen. Pull self to sit.	Index approach. Uses index finger to poke at pea. Able to let go of cube in hand.		Waves "Bye bye" Plays "Pat-a-Cake"
11 months	Creeping on all FOURS Pivoting. Cruising. Walks with 2 hands held.		ONE word with meaning.	Plays "peek-a-boo"
1 year	Gets from lying to sitting to crawling to standing. Walks like a bear. Walks with ONE hand held. Walks well (13 months). Stands alone.	Neat pincer grasp. Bangs 2 cubes. Sees and picks up hundreds and thousands.	Understands phases. (e.g. where is your shoes). 2 - 3 words with meaning. Localising sound above head.	Casting (13 months) Less mouthing. Shy.
15 months	Creeps upstairs. Stoops for toy and stands up without support. (best at 18 months)	Tower of 2 cubes. Scribbles spontaneously (15-18 months)	More words. Points to objects he wants. Continual jabber and jargon.	Takes off shoe. Feeds self with cup (able to pick up and put down) and spoon (but spills). Mouthing stops.
18 months	Gets up and down stairs holding on to rail or with one hand held. Pulls toy or carries doll. Throws ball without falling. Sits on a chair.	Tower of 3 cubes. Scribbles spontaneously. Visual test: Picture charts. Handedness (18 - 14 months).	Points to 2 - 3 body parts. Picture Cards - identify one.	Imitates housework. Toilet trained. Uses spoon well. Casting stops.

AGE	Gross Motor	Vision & Fine Motor	Speech & Language	Personal Social
2 years	Goes up and down stairs alone, 2 feet per step. Walks backwards (21 months) Runs. Picks up toy without falling. Able to throw and kick ball without falling.	Tower of 6 cubes Imitates cube of train with no chimney. Imitates straight line.	Joins 2 - 3 word in a sentence. Uses 'you' 'me' 'I'. Picture cards - Names 3 objects Points to 5. Obeys 4 simple commands. Points to 4 body parts.	Puts on shoes, socks, pants. Dry by day. Play near other children but not with them.
2 1/2 years	Jumps on both feet. Walks on tip toes.	Tower of 8. Imitates train with chimney. Holds pencil well. Imitates and	Knows FULL name and sex. Names one colour.	
3 years	Goes up stairs one foot per step. Down stairs 2 feet per step. Jumps off bottom step. Stands on 1 foot for seconds. Rides tricycle.	Tower of 9. Imitates bridge. with cubes: Copies Imitates Draw a man test. (3 - 10y)	Can count to 10 Names 2 colours. Nursery rhymes. Understands "on", "in", "under".	Unbuttons. Dresses and undresses fully if helped with buttons and advised about correct shoe. Dry by night. Plays with others.
3 1/2 years		Copies bridge.		
4 years	Goes up and down stairs one foot per step. Skips on one foot. Hops on one foot.	Imitates gate with cubes. Copies + Goodenough test 4.	Names 3 colours. Fluent conversation. Understands "in front of", "between", behind".	Buttons clothes fully. Attends to own toilet needs.
4 1/2 years		Copies gate with cubes. Copies square. Draws recognisable man and house.		



Age† (months)	Gross motor control	Vision and fine motor	Language and hearing	Social and daily living skills
3	Complete head lag*	Following with eyes	Searching for sounds with eyes	Smiling
6	Preference for one hand Persistent Moro*	Squint*	Head turn to soft voice	Interest in people
9	Sitting with support	Persistent hand regard*	Ba-ba-ba babble	Awareness of strangers
12	Pulling to stand Standing with support	Pincer grosp	Trying one or two words	Constant* mouthing
18	Walking alone	Constructive play with blocks Casting toys*	Six words Constant dribbling*	Pointing at items Finger feeding
24	Running	Turning book pages	Fifty single words	Interested in other children Helps in dressing
36	Kicking a ball	Drawing lines	2–3 word phrases Echolalia*	Active play with peers
48	Pedalling and hopping	Drawing a face	Sentences and 'Wh' questions	Imaginative play Toilet trained — day

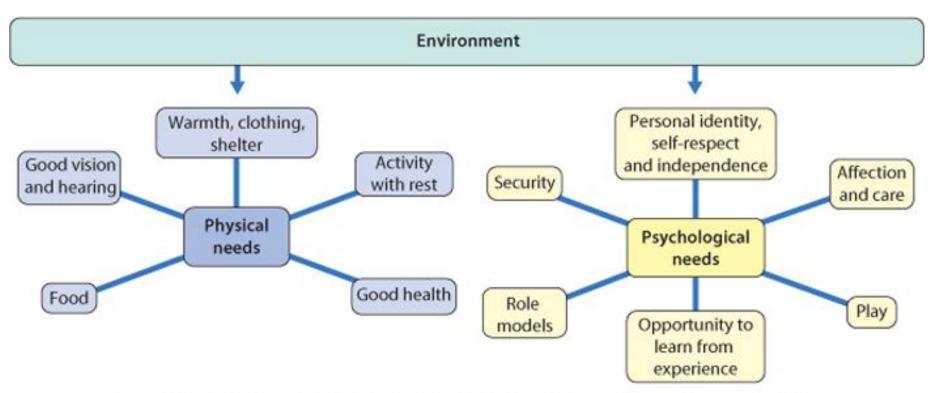
NORMAL RANGES (APPROXIMATELY 25TH CENTILE TO 90TH CENTILE)

Age	Gross motor control	Vision and fine motor	Language and hearing	Social and daily living skills
2-4 months	Head steady in sitting	Follows object through 180°	Squeals with pleasure	Smiles
5-8 months	Sits without support	Passes cube hand to hand	Turns to soft voice Baba/Gaga babble (up to 10 months)	Feeds self biscuit
9-14 months	Stands with support	Neat pincer grasp of raisin	Mama or Dada specifically	Indicates needs by gesture
12-16 months	Walks well alone (up to 21 months)	Stack of two cubes (up to 21 months)	Three words	Drinks from a cup
15-24 months	Walks up steps	Scribbles spontaneously	Points to one body part	Removes garment
21-36 months	Jumps on the spot	Draws vertical line in imitation	Uses plurals and phrases	Puts on clothing Plays tag with other children
3-4½ years	Balances on one foot for 5 s	Copies a ladder Draws a face	Understands cold, tired and hungry Asks 'Wh' questions	Separates from mother

WARNING SIGNS TO WORRY ABOUT

Age (months) [†]	Gross motor control	Vision and fine motor	Language and hearing	Social and daily living skills
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48	Pedalling and hopping	Drawing a face	Sentences and 'Wh' questions	Imaginative play Toilet trained by day

Developmental assessment....setting the scene



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Developmental assessment...1

$2-2^{1}/_{2}$ year review

Your child is 2-2¹/₂ years old and is learning many new skills, such as:

- * wanting to explore everything and be more independent
- * wanting to run and climb and always being on the go
- enjoying messy play but not sharing!
- * starting to join up words and trying to repeat words you say. Favourite words are "NO" and "MINE!"
- * enjoying books and joining in with songs and rhymes
- * liking being close to you and having cuddles and hugs
- playing with other children
- * using a spoon at mealtimes and using a feeder cup
- * starting to show an interest in potty training
- * turning from laughter to anger very quickly, which can be hard work

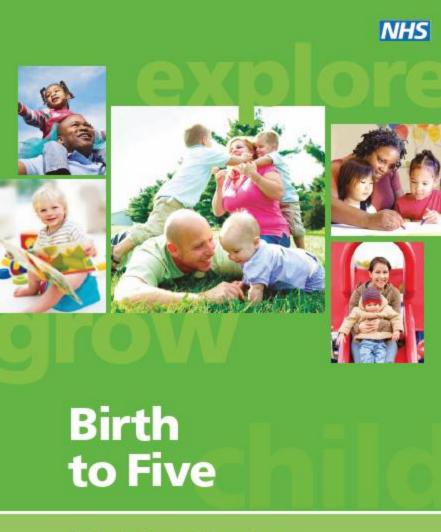
S/he has got used to tooth brushing with a fluoride toothpaste. S/he has been to the dentist.

Birth to Five gives information about what children are usually doing at this age.

Other things you may want to talk about at the review are:

- speech and language
- learning
- ★ diet
- behaviour
- safety
- your own health

You may find it helpful to write down here anything you would like to discuss at the 2-21/2 year review:



This book gives you information on:

Becoming a parent Taking care of yourself and your child Finding practical help and support

Developmental assessment...2

Observe the child

- Gross motor
- Fine motor
- Speech
- Social & play

Use play

Do not force your agenda

Give the child appropriate props

- Bricks & 'thousands'
- Pencil & paper & crayons
- Ball
- Doll
- Pictures

Developmental assessment...3

Standardised tests

- •<u>S</u>chedule <u>o</u>f <u>G</u>rowing <u>S</u>kills (0-5 years)
- Griffiths Mental Development Scales (0-8 years)

SOGS II

	Screening Da	tes		Screening Dat	es	
P/	SSIVE POSTURAL SKILLS		U	OCOMOTOR SKILLS		+
Su	pine Position		M	lovement and Balance		+
1.	Head in midline	1	22	Rolls and squirms to move about	1	-
2.	Lifts legs into vertical position and grasps		23.		2	-
* *	foot (foot regard)	2	24	Walks with hands held, taking full weight on fee	t 3	-
Ve	ntral Suspension		25.			+
3.	Head in line with body, hips semi-extended	1	26.			
4.	Head above line of body, hips and shoulders extended	2	27.	Walks well, feet only slightly apart, can turn corners and stop suddenly	6	
Pu	ll to Sit		28.	Picks up objects from floor without falling	7	
5.	Head lag on pulling, when body vertical; head held momentarily erect before falling forwards	1	29.	Runs confidently, stopping and starting with care and avoiding obstacles	8	1
6.	Little or no head lag	2	30.	Jumps taking both feet off the ground	9	
7.	Braces shoulders and pulls self up	3	31.	Walks tiptoe	10	
Siti	ing Position (supported by adult)		32.	Runs tiptoe	11	1
3.	Back curved	1	33.	Hops on one foot for 3 steps	12	1
9.	Back straight	2	34.	Heel-to-toe walking forwards (for a minimum of 4 steps)	13	1
PAS	SSIVE POSTURAL SKILLS SCORE		35.	A DESCRIPTION OF A REAL OF A DESCRIPTION OF A DESCRIPTION OF A DESCRIPTION	13	+
40	TIVE POSTURAL SKILLS		St.	airs	14	+
_						+
_	ne Position			Crawls upstairs	1	-
0.	Head sideways, resting on cheeks, buttocks high with knees flexed under abdomen, arms close to chest with elbows flexed	1	37.	1	2	+
1.	Lifts head momentarily, buttocks high	2	39.		5	+
2.	Holds up head and upper chest on forearms, with buttocks flat	3	40.	downstairs (two feet to a step) Walks alone upstairs and downstairs -	4	+
3.	Supports weight on flattened palms and extended arms	4	41.	one foot per step (adult fashion) Runs upstairs	5	+
4.	Gets into crawling position	5	LO	COMOTOR SKILLS SCORE	F	+
litt	ing Position (unsupported)				100	-
5.	Sits alone momentarily without support	1	M	ANIPULATIVE SKILLS		
6.	Sits alone for prolonged periods (at least to the count of 10)	2	Ha	and Skills		
7.	Gets into sitting position from either		42.	visitio otocci and thank tanke in	1	
	prone or supine	3		Hand regard and finger play	2	
	nding			Clasps hands and presses palms together	3	4
-	Held standing, bears some weight on feet	1		Palmar grasp	4	1
	Held standing, takes full weight on feet	2	46.		5	+
0.	Stands holding on	3	47.	 Holds two cubes – one in each hand, bringing them together 	6	
	Pulls self to stand	4	48.	Inferior pincer grasp	7	+
CT	IVE POSTURAL SKILLS SCORE		49.	 Neat pincer grasp 	8	+
			50.	 Throws toys to the floor deliberately (casting) 	9	+
			51.		10	+
		F Martine P	52.		11	+
KEY		53.		12	+	
	Stimulus material needed for this item.		54.	Contraction of the second second second	13	+
	This item contains a cognitive element.			reported in personal in 50 seconds		1

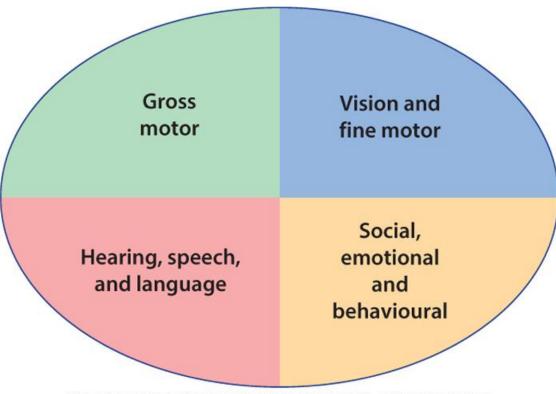
The Schedule of Growing Skills II - Record Form

Griffiths

Subscale A Locomotor		Response	Subscale B Personal-Social		Response	Subscale C Language		Response
1	Jumps off 1 step 🛛 🕞; 🗺		1	Puts away toys when encouraged to do so		1	Names 12 objects in box	
2	Static balance 1: can stand on one foot for 3+ seconds (); 122 ()		2	Gives first name		2	Picture vocabulary (12) (NB: Administer after item FIII.10)	
3	Can rise from kneeling without using hands		3	Assists with small household tasks on request		3	Defines by use (2+)	
4	Can run fast indoors or in a small outside space		4	Uses spoon and fork together, without help		4	Picture description: names 6+ objects in large picture	
5	Can stand and walk tip-toe: 6+ steps		5	Knows own gender		5	Uses 2 or more descriptive words	

Subscale D Eye and Hand Co-ordination		Response	Subscale E Performance		Response	Subscale F Practical Reasoning		Response
1	Builds a tower of 8+ bricks		1	4-squares board: 50 secs		1	Knows 'penny' or 'money'	
2	Copies a horizontal stroke		2	6-hole board: 50 secs		2	Repeats one digit (8; 2; 7)	
3	Handles scissors: tries to cut		3	Returns 9 bricks to box and puts lid on: 50 secs		3	Compares two insets for size	
4	Threads ó beads 🛛 🛈 🖬		4	Reassembles screw toy 🛛 🕞 📴		4	Repeats 2 digits (1-6; 5-3; 9-4) 🔶	
5	Copies a circle: Stage 1		5	4-squares board: 40 secs		5	Knows 'big' and 'little'	

So.....



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Developmental milestones

While looking at the video make notes on what skills each child is able to perform in each of the 4 developmental domains

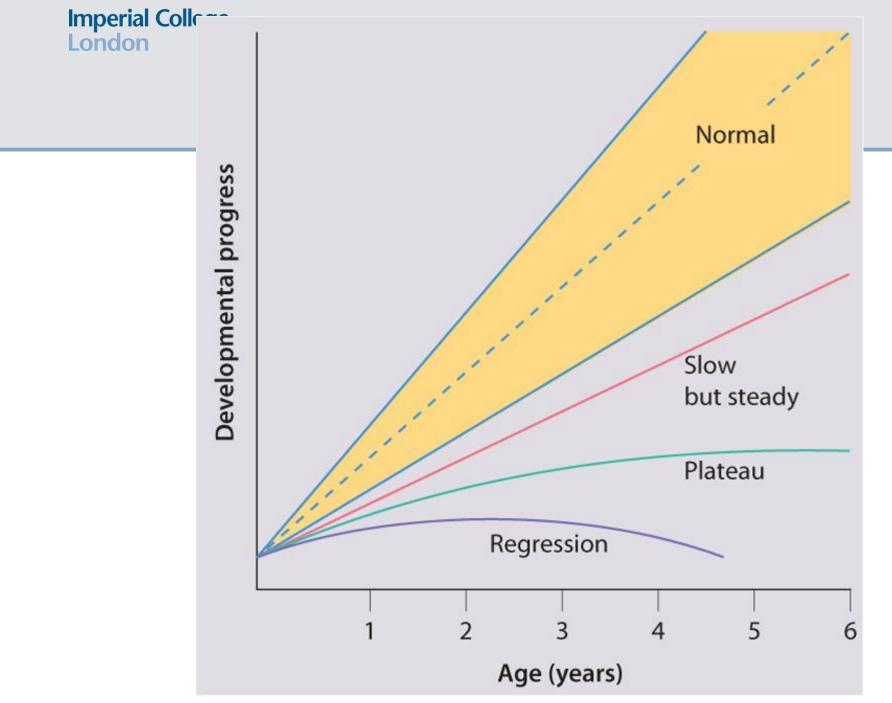
Estimate the developmental age of the child using the table of milestones

	Α	В	С
Gross motor/ posture			
Fine motor/ vision			
Language/ hearing			
Social/ behaviour			

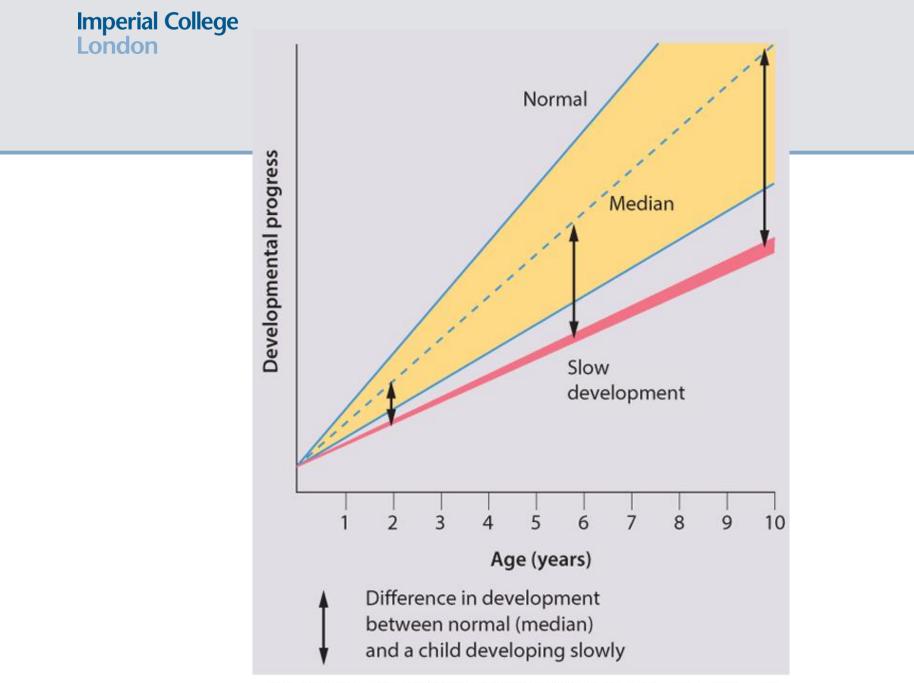
Abnormal Development

The Questions

- *What* is the problem?
- Where is the problem?
- Which key development field is worst affected?
- *Why* did things go wrong?
- When did things go wrong?
- *How* did things go wrong?
- Who is affected?
- *What* can be done?



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What is the problem?

Cerebral Palsy – a prototype of abnormal development

Definition

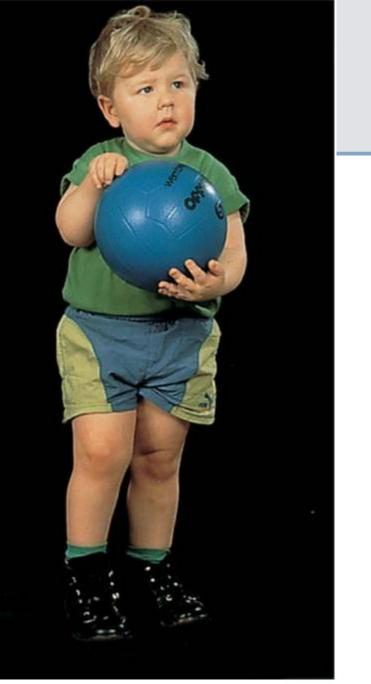
➢ is a disorder of movement and posture due to a non-progressive lesion of motor pathways in the developing brain.

➤ the clinical manifestations emerge over time, reflecting the balance between normal and abnormal cerebral maturation.

➢ is the most common cause of motor impairment in children, affecting about 2 per 1000 live births.

Other problems

- learning difficulties (about 60%)
- •epilepsy (40%)
- •squints (30%)
- •visual impairment from errors of refraction and cortical damage (20%)
- •hearing impairment (20%)
- •speech and language disorders (due to hearing loss, oromotor incoordination and learning difficulties)
- behaviour disorders
- •feeding problems
- •joint contractures, hip subluxation, scoliosis.

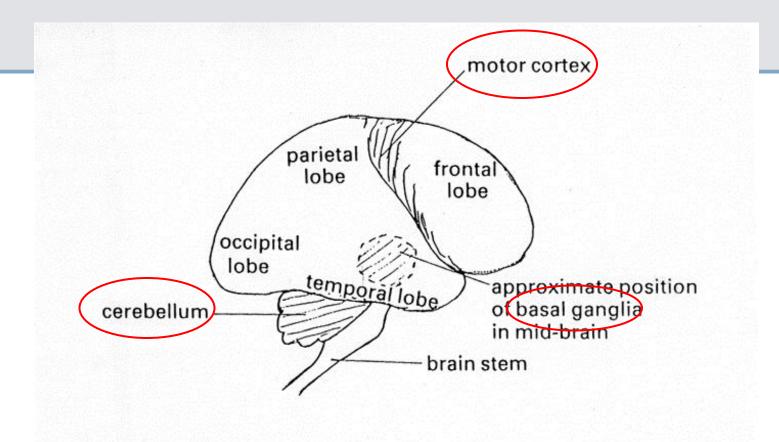


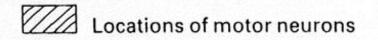


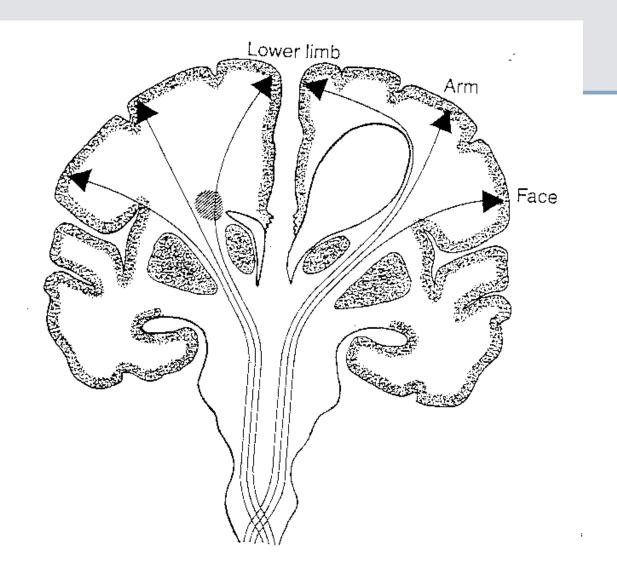
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Where is the problem?

Motor neurones







Why did things go wrong?

Genetic

Acquired

When did things go wrong?

Prenatal

Perinatal

Infancy

Early childhood

How did things go wrong?

Hypoxia-ischaemia

Hypoglycaemia

Infections

Trauma

Other



•~ 80% of cerebral palsy is antenatal in origin - genetic syndromes and congenital infection.

•Only about 10% of cases are thought to be due to hypoxicischaemic injury at birth

•About 10% are postnatal in origin.

- rise in survival of extremely preterm infants meningitis/encephalitis/encephalopathy,
- head trauma symptomatic hypoglycaemia, hyperbilirubinaemia.

Who is affected?

Babies (Preterm/Term)

Infants

Young children

What can be done?

Hearing Conductive or

Conductive or sensorineural hearing impairment

Vision

Squint Impaired visual acuity Visual field deficits

Orthopaedic

Hip subluxation/dislocation Fixed joint contractures Dynamic muscle contractures Painful muscle spasm Spinal deformity Osteoporosis/fractures

Specialist health visitor

Help coordinate multidisciplinary and multiagency care Advice on development of play or local authority schemes e.g. Portage

Dietician Advice on feeding and nutrition

Social worker/ Social services Advice on benefits: disability, mobility, housing, respite care, voluntary support agencies Day nursery placements Advocate for child and family Register of children with special needs

Psychologist (clinical and educational)

Cognitive testing Behaviour management Educational advice

Gastrointestinal

Gastro-oesophageal reflux Oromotor incoordination Aspiration of food or saliva Constipation

Urogenital

Urinary tract infection Delay in establishing continence Unstable bladder Vesico-ureteric reflux Neuropathic bowel and bladder

Common medical problems



Child Development Service

Paediatrician

Assessment, investigation and diagnosis Continuing medical management Coordination of input from therapists and other agencies - health, social services, education

Respiratory

Respiratory infections Aspiration pneumonia Chronic lung disease Sleep apnoea

Neurological

Epilepsy Microcephaly/ hydrocephalus Cerebral palsy

Nutrition

Poor weight gain Failure to thrive

Behaviour

Organic or reactive Sibling behaviour Parental distress

Speech and language therapist

Feeding Language development Speech development AAC (augmentative and alternative communication) aids e.g. Makaton sign language, Bliss symbol boards, voice synthesizers

Occupational therapist

Eye-hand coordination ADL (activities of daily living) feeding, washing, toileting, dressing, writing Seating Housing adaptations

Physiotherapist

Balance and mobility Postural maintenance Prevention of joint contractures, spinal deformity Mobility aids, orthoses

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VIDEO

(Final) Summary

- Basic review of embryology
- > Mapped how abnormalities $\rightarrow \rightarrow$ clinical scenario
- Normal development and milestones
- Framework tool for assessment
- > When development is abnormal an example
- Year 3 Introduction to Clinical Specialties

Any questions?

Acknowledgements and further reading

Dr Annie Dai, Consultant Community Paediatrician

Lissauer, T and Clayden, G. Illustrated textbook of Paediatrics. 3rd ed

Holt, K. Child development: diagnosis and management. 1991

Robertson, D.M. Practical Paediatrics. 6th ed, 2007