PSYCHOMOTOR RETARDATION IN CHILDHOOD

Causes, diagnosis, management

CHILD DEVELOPMENT



- Child development- abilities acquired by child from birth to 5 yr
- Psychomotor development skills acquired by the child until its level of cognitive function is advanced enough that they can be assessed independently from the motor development.

PSYCHOMOTOR DEVELOPMENT

- The term "psychomotor development" refers to the first 2-3 years of age.
- Developmental delay is a group of symptoms, not the diagnosis.
- That means slow skill acquisition.

DEFINITION



The delay is when a child does not reach the milestones in expected time, taking into account the two deviations from mean.

MILESTONES

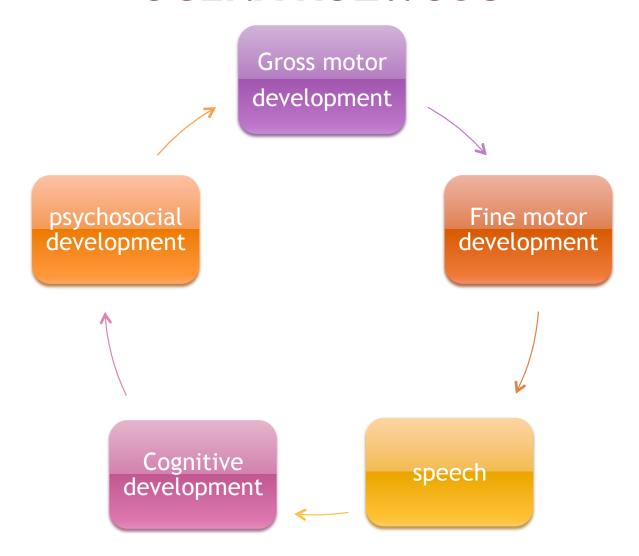


- This is the most important skill development
- It always refers to the age of the child, and preterm infants to the adjusted age

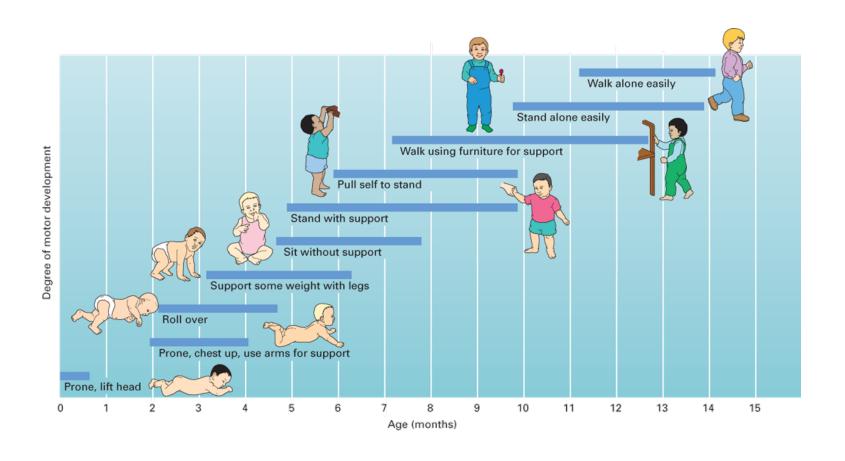
MILESTONES

- Median age when half the population acquire a skill; serves as a guide to normal pattern of development
- Limit age when a skill should have been acquired; further assessment is indicated if not achieved, it is determined as two deviations from the mean
- In the limited age many children with LBW, ELBW develop

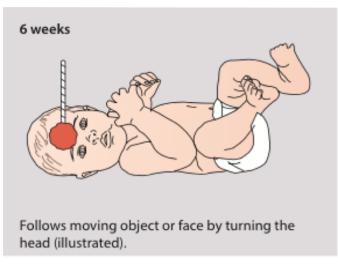
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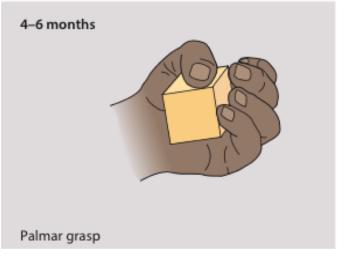


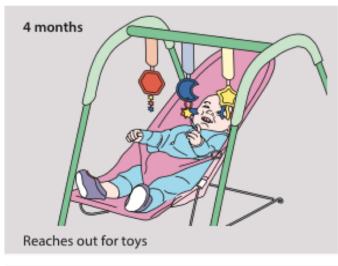
MILESTONES- GROSS MOTOR DEVELOPMENT



MILESTONES- VISION, FINE MOTOR DEVELOPMENT

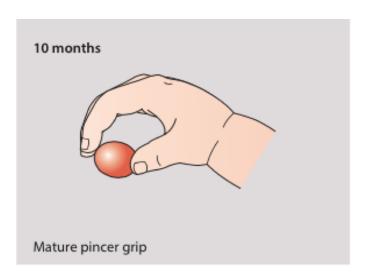


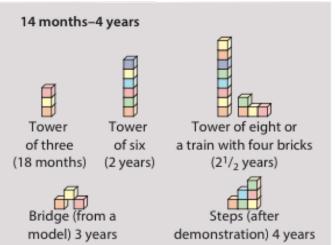




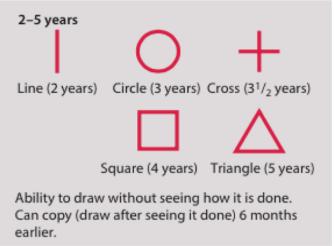


MILESTONES- VISION, FINE MOTOR DEVELOPMENT

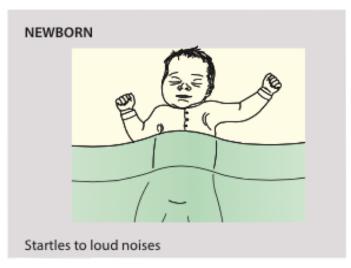






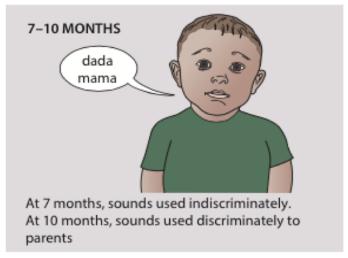


HEARING, SPEECH AND LANGUAGE







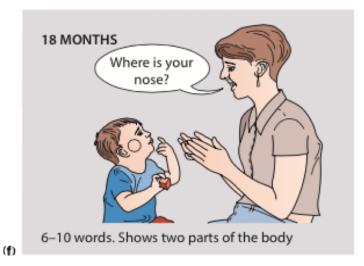


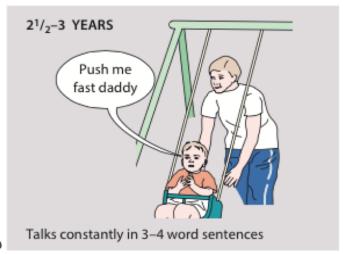
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HEARING, SPEECH AND LANGUAGE









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MILESTONES OF SPEECH DEVELOPMENT

| Language, Speech milestones | | |
|-----------------------------|---|--|
| Age | Milestone | |
| 1 months | Alerts to sound | |
| 3 months | Coos | |
| 4 months | Laugh loud | |
| 6 months | Monosyllables | |
| 9 months | Bisyllables | |
| 12 months | 1-2 words with meaning | |
| 18 months | 8-10 words vocabulary | |
| 2 years | 2-3 words sentence, use pronouns "I", "me", "you" | |
| 3 years | Ask questions; knows full name and gender | |
| 4 years | Says song or poem; tells stories | |
| 5 years | Asks meaning of words | |

PSYCHOSOCIAL DEVELOPMENT, BEHAVIOUR



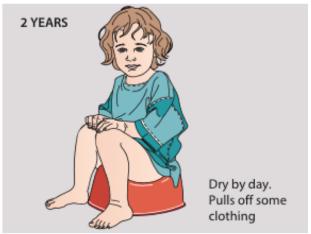


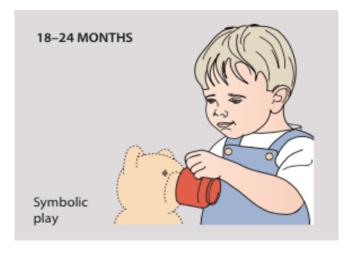


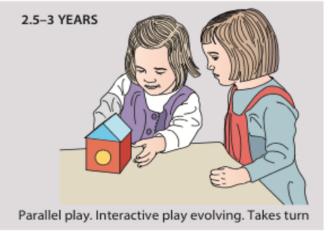


PSYCHOSOCIAL DEVELOPMENT, BEHAVIOUR









PSYCHOSOCIAL DEVELOPMENT-MILESTONES

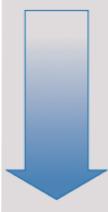
| Social and adaptive milestones | | |
|--------------------------------|---|--|
| Age | Milestones | |
| 2 months | Social smile | |
| 3 months | Recognizes mother; anticipates feeds | |
| 6 months | Recognizes strange/ stranger anxiety | |
| 9 months | Waves 'bye-bye' | |
| 12 months | Comes when called; plays simple ball game | |
| 15 months | Jargon | |
| 18 months | Copies parents in task | |
| 2 years | Asks for food, drink, toilet; pulls people to show toys | |
| 3 years | Shares toys; knows full name and gender | |
| 4 years | Plays cooperatively in a group; goes to toilet alone | |
| 5 years | Helps in household tasks; dresses and undresses | |

MILESTONES

| Age | Gross Motor | Visioon&Fine Motor | Hearing, Speech&Language | Social, emotional&behavioural |
|-----------------|----------------------|--|---|---|
| Newborn | Flexed posture | Fixes and follows face | Stills to voice Startles to loud noise | Smiles by 6 weeks |
| 7 months | Sits without support | Transfer objects from hand to hand | Turns to voice, Polysyllabic babble | Finger feeds Fears strangers |
| 12 months | Stands independently | Pincer grip (10 months), Points | 1 -2 words Understand name | Drinks from the cup Waves |
| 15-18 months | Walks independently | Immature grip of pencil, Random scribble | 6 -10 words Points to four body parts | Feeds self with a spoon Beginning to help with dressing |
| 2½ years | Runs and jumps | Draws | 3 -4 word sentences Understands two joined commands | Parallel play Clean and dry |

RED FLAGS

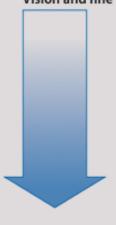
Gross motor development



- Acquisition of tone and head control
- Primitive reflexes disappear
- Sitting
- Locomotor patterns
- · Standing, walking, running
- · Hopping, jumping, peddling

| Gross motor | Limit ages |
|---|---------------------------------------|
| Head control Sits unsupported Stands independently Walks independently | 4 months 9 months 12 months 18 months |

Vision and fine motor development



- Visual alertness, fixing and following
- · Grasp reflex, hand regard
- Voluntary grasping, pincer, points
- Handles objects with both hands, transfers from hand to hand
- Writing, cutting, dressing

| Fixes and follows 3 months | |
|---|--|
| visually Reaches for objects Transfers Pincer grip 6 months 9 months 12 months | |

RED FLAGS

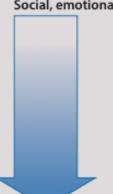
Hearing, speech and language development



- · Sound recognition, vocalisation
- Babbling
- Single words, understands simple requests
- · Joining words, phrases
- Simple and complex conversation

| Hearing, speech and language | Limit ages |
|--|--|
| Polysyllabic babble Consonant babble Saying 6 words with meaning Joins words 3-word sentences | 7 months 10 months 18 months 2 years 2.5 years |

Social, emotional, behaviour development



- · Smiling, socially responsive
- Separation anxiety
- Self-help skills, feeding, dressing, toileting
- · Peer group relationships
- Symbolic play
- · Social/communication behaviour

| Social behaviour | Limit ages |
|--|---|
| Smiles Fear of strangers Feeds self/spoon Symbolic play Interactive play | 8 weeks 10 months 18 months 2–2.5 years 3–3.5 years |



- The universal definition and classification of developmental delays is not established.
- The development level lower than the average may be present in one or more areas.
- The global developmental delay it is a significant delay in at least two of its areas



- Development, regardless of its speed, it can be harmonious or disharmonious
- Disharmonious development of individual skills is often observed in healthy children

Delays in a very narrow range (e.g. only manual skills or hand-eye coordination) is called specific development deficits efficiency





It is proposed that "global delay development" replace more diagnostic the term "early developmental disorders": persistent, significant limitations in at least two areas in children under 5 years, which currently can not be given other diagnose.

Developmental screening allows to capture global or fragmentary developmental delays in most cases.

In problematic/ borderline cases standard diagnosis will be more certain if it is established on several sources:

- standardized interview
- direct observation
- psychological examination
- speech evaluation

- There is no universal rating scale which assess the development
- The most commonly used is the scale of Denver
- There are standardized tests to evaluate the infants and toddlers: the scale of the development of the infant Griffiths or Bailey

Psychomotor delay is suspected when the development is slower then mean:

- First 6 months of life- more then 1 month
- □ 6-12 months of life more then 2 months
- □ 1-1,5 years- more then 3 months

Diagnosis should be made at least in age of 9 years old.



INCIDENCE

- Approximately 25% children in outpatient clinic is observed developmental delay
- 5-10% is diagnosed with some neurodevelopmental disorders
- Most of these "grows" from their problems spontaneously or with a little help

INCIDENCE

Global developmental delay and associated with it intellectual disability occures at least 3% of children in the group to 5 years old.



DELAY AND REGRESION

Regress - the loss of acquired skills

It can occur in a child previously properly developing and in children, the development of which was already delayed



REGRESION- CAUSES

| epilepsy |
|---|
| progessive hydrocephalus |
| proliferative process CNS |
| Vascular diseases |
| Infetious diseases |
| Inflammatory diseases |
| Autoimmunologic diseases (HIV, SSPE- subacute sclerosing panencephalitis, ADEM-Acute Disseminated Encephalomyelitis) |
| Drugs side effects |
| Psychiatric disorders |
| autism |

CAUSES AND DIAGNOSIS OF THE DEVELOPMENTAL RETARDATION

- retardation may be either isolated or present with the other disorders
- the reasons for the delay of development is impossible to determine in 40-60% of children



CAUSES AND DIAGNOSIS OF THE DEVELOPMENTAL RETARDATION

- family history (3 generations)
- pregnancy and perinatal period
- evaluation of the current development
- evaluation of the results of basic research
- pediatric study
- neurological examination

Help find the cause of about 30% of delays development

CAUSES AND DIAGNOSIS OF THE DEVELOPMENTAL RETARDATION

- the primary reason for the delay of development is often masked by prematurity and / or hypoxic-ischemic encephalopathy
- always have to take into account the impact of the toxic factors to the fetus - alcohol, nicotine, lead, phenylalanine in the mother with undiagnosed, untreated phenylketonuria

The greatest chance of success gives diagnostic team work:

pediatrician
neurologist
psychologist
clinical geneticist
neuroradiologist
specialist in metabolic diseases



LABORATORY TESTS

Blood tests:

karyotype, fragile X

Complete blood acount

Creatine kinase

TSH/fT4

lactates, amino acids, ammonia

VLCFA, carnitine, homocysteine

toksoplasmosis, CMV

rubella, HIV

Urine test:

- CMV
- organic acids
- orotic acid
- mucopolysaccharides

Skin test in Wood lamp

DIAGNOSTIC SCHEME

When the initial diagnosisis no possible to make - based on history and physical examination:

First line:

- karyotype
- fragile X
- telomeres
- MLPA
- Urea, electrolytes
- creatine kinase
- Lead
- Thyroid hormone
- Complete blood count
- ferritin
- Biotynidaza

DIAGNOSTIC SCHEME

Second line

Neuroimaging: abnormal head circumference focal seizures Metabolic disorders: Family history blood relationship regress organomegaly family features

Second line

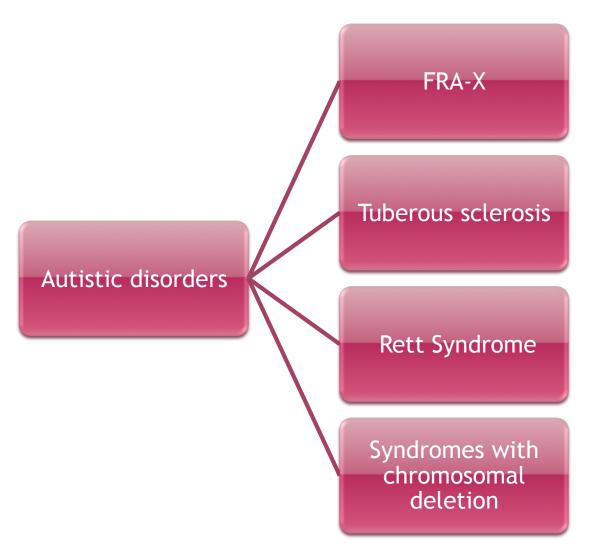
Genetics:

- dysmorphy
- Abnormal growth
- Hearing and vision disorders
- Behavioural disorders
- Family history

EEG:

- Speech delay/ speech regression
- seizures

GENETIC SYNDROMES WITH DEVELOPMENTAL DELAY



GENETIC SYNDROMES WITH DEVELOPMENTAL DELAY

Mikrocefalia

- Zespół Retta lub mutacja MECP2
- Zespół Angelmanna
- Zaburzenia Rett-like (mutacje CDKL5 i FOXG1)
- Zespoły z delecją chromosomów
- Zespół Pitta-Hopkins
- Zespół Mowata-Wilsona
- Zespół SLO
- Zespół Cornelii de Lange
- Zespół Cohena

Makrocefalia

- Zespół Sotosa
- FRA-X
- Zespół Costello
- Zespół Lujana
- PTEN
- Mutacje w RAB39B i oPHN1

SYMPTOMS SUGESTING METABOLIC DISEASE OR NEURODEGENERATIVE DISEASE

Neurologic disorders

- hypotonia
- convulsions
- Ataxia
- paroxysmal disorders of the concoiusness
- extrapyramidal disorders

Irregulatories in biochemistry tests:

- acidosis
- hyperammonemia
- hyperuricaemia
- relapsing hypoglycemia
- hipertransaminaze mia, cholestasis

SYMPTOMS SUGESTING METABOLIC DISEASE OR NEURODEGENERATIVE DISEASE

Physical disorders

- Lack of appetite
- Vomiting

Organomegaly

- hepatomegaly, splenomegaly
- cardiomiopathy

Eyeballs

- Retinopathy
- cataract
- corneal opacity

The winding, rough, brittle hair and

 impairment deep, significant laxity of the skin, light skin

SYMPTOMS SUGESTING METABOLIC DISEASE OR NEURODEGENERATIVE DISEASE

Bone anomalies

- Dysostosis multiplex
- chondrodysplazja
- severe bone pain

Skin rashes, ichthyosis, and

 Mental impairment, epilepsy

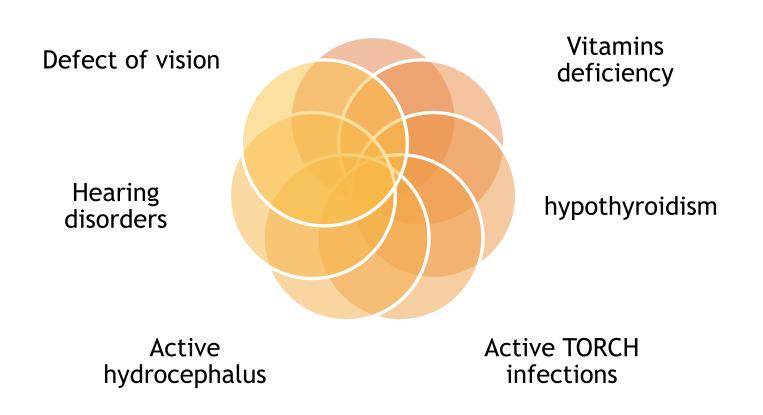
Dysmorphism features

- thickening face features
- features like Down syndrome
- features like FAS



In any case of delayed development it is necessary to use the symptomatic treatment or/and causal treatment.





THERAOY

Treatment of epilepsy

Inflamatory and autoimmunological diseases treatment

Inborn errors of metabolism (e.g. PKU, Homocystinuria, galactosemia, biotinidase deficiency, Vit.B1 deficiency)



- Children are targeted for rehabilitation (early intervention / early intervention program development).
- 2. In children at risk of developmental disorders early intervention is associated with improved cognitive development in infancy and preschool.
- The most important in the intensive health care is family.
- Children with a slight developmental delay can be observed by a pediatrician.
- 5. Parents is granted briefing care during meetings with physiotherapist, speech therapist, psychologist.





Early intervention should always be considered in children exposed to sensory deprivation for social reasons:

poor environmental conditions abandonment depression in the mother



- 1. In children moderately and significantly delayed early intervention is needed and effective at the right time and in a manner adapted to the needs and abilities of the child.
- 2. Early intervention is not always immediate.
- Critical periods for the development of most of the functions continues throughout childhood.
- 4. Only the processes of vision and hearing are relatively short (one year old).



We rehabilitate child, only if it requires it.

The appearance of the function (ability) decides the period of overproduction of synapses in the brain region, which responds for it.

Stimulating skills before the period overproduction of synapses does not make sense.

It may even be harmful, because it interferes with the natural compensatory mechanisms and influence on the for the process brain plasticity in wrong way.



A prerequisite for the existence of plastic processes is the presence of connections between neurons (sending and receiving centers).

If in some region of the brain connections not arise or be destroyed irreversibly (porencefalic pit), and the brain does not produce alternative routes, the early intervention will be ineffective.



REFFERENCES AND SOURCES

- 1. Standardy postępowania diagnostyczno-terapeutycznego w schorzeniach układu nerwowego u dzieci i młodzieży pod red. B. Steinborn
- 2. Illustrated Textbook of Paediatrics, 4th ed., T.Lissauer, G.Clayden
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