

Examination of hearing loss in newborns

Hearing defect

- 6 - 12 / 1000
- 1 / 1000 has a very severe hearing loss
- Higher incidence than in phenylketonuria, congenital hypothyroidisms

Hearing defect

- congenital (genetic)
- obtained (during pregnancy, childbirth or shortly after, during the life-infection, trauma, tumor ..)

Hearing defect

- **prelingually** (created before the end of the child's speech development, 6th year of age)
- **postlingually** (persons with advanced speech)

Genetic hearing loss

- several genes
(mostly gene for connexin)
- these are the different types disability of cochlea
- (Michel's, Mondini's, Scheibe's type ...)



Scheibe's dyspalsia

- Scala media disappeared
- Bony labyrinth is normal
- Dyspalsia is seen in the cochlea and saccule
- Autosomal-recessive, non syndromic

Michel's aplasia

- Complete absence of bony and membranous labyrinth
- External and middle ear may be unaffected

Mondini's dysplasia

- May be seen in many syndromes
- Only basal coil is present

Genetic hearing loss

- part of **syndromes**
- Alport syndrome (a kidneys)
- Uscher syndrome (an eye)
- Turner syndrome (45X) ..
- **incidence can be reduced**



Acquired hearing defect

- Prenatal- before birth
- Perinatal- during birth
- Postnatal- after birth

Acquired hearing defect

- "Sensitive" period is around 20 day after conception
- maternal infectious diseases (rubeolla, measles, mumps, scarlet fever, flu, toxoplasmosis in the first trimester)
- ototoxic drugs, X-ray
- maternal metabolic diseases (diabetes, uncontrolled high blood pressure ..)

Acquired hearing defect

- **premature birth** - risk neonate
- **risk increase** 20-40 / 1000
- **at birth** (neonatal prematurity, Rh-incompatibility, a difficult birth, which may cause bleeding into the inner ear or to the brain centers for the hearing, asphyxia or hypoxia of the child)
- **after birth** (mechanical ventilation longer than 4 days, severe neonatal jaundice, severe fetal infection, meningitis)

Identifying hearing loss

- **difficult**, if not profound deafness
- parents perceive deviation after several months (**2-3 years**), delayed speech, poor pronunciation, or disobedience



Identifying hearing loss

- hearing test, reaction to sound
- lack of adequate response
- undeveloping speech
- the incidence of congenital deafness in the family
- followed by an objective hearing tests

Screening

- Screen = sieve
- search method of early forms of disease, or deviations from the norm in a population through tests



TEOAE

- In the **neonatal** ward
- **1st** and **4th** day
- **quiet room**, usually after feeding
- Into ear insert a **small probe** (transmitting sounds and records the otoacoustic emissions - the sound of the probe evoked "response" of outer hair cells in the cochlea (inner ear)).

- emissions are present ("**PASS**"), confirmed the proper function of outer hair cells - probably is not present any hearing defect.
- emissions are absent ("**REFER**"), which may indicate a hearing defect.



- continual monitoring the child and his speech development
- rare and unlikely hearing disorders (**disorder of the auditory nerve**), which may begin to develop after birth
- hearing may be damaged in the **child development**

- For PASS TEOAE is needed not only proper function of outer cells in the cochlea, but also air and function of the **middle ear and ear canal**.
- REFER do not always mean (sensorineural) hearing loss.
- absence of emission (30%), it is appropriate to repeat the test after two days, or after a month.

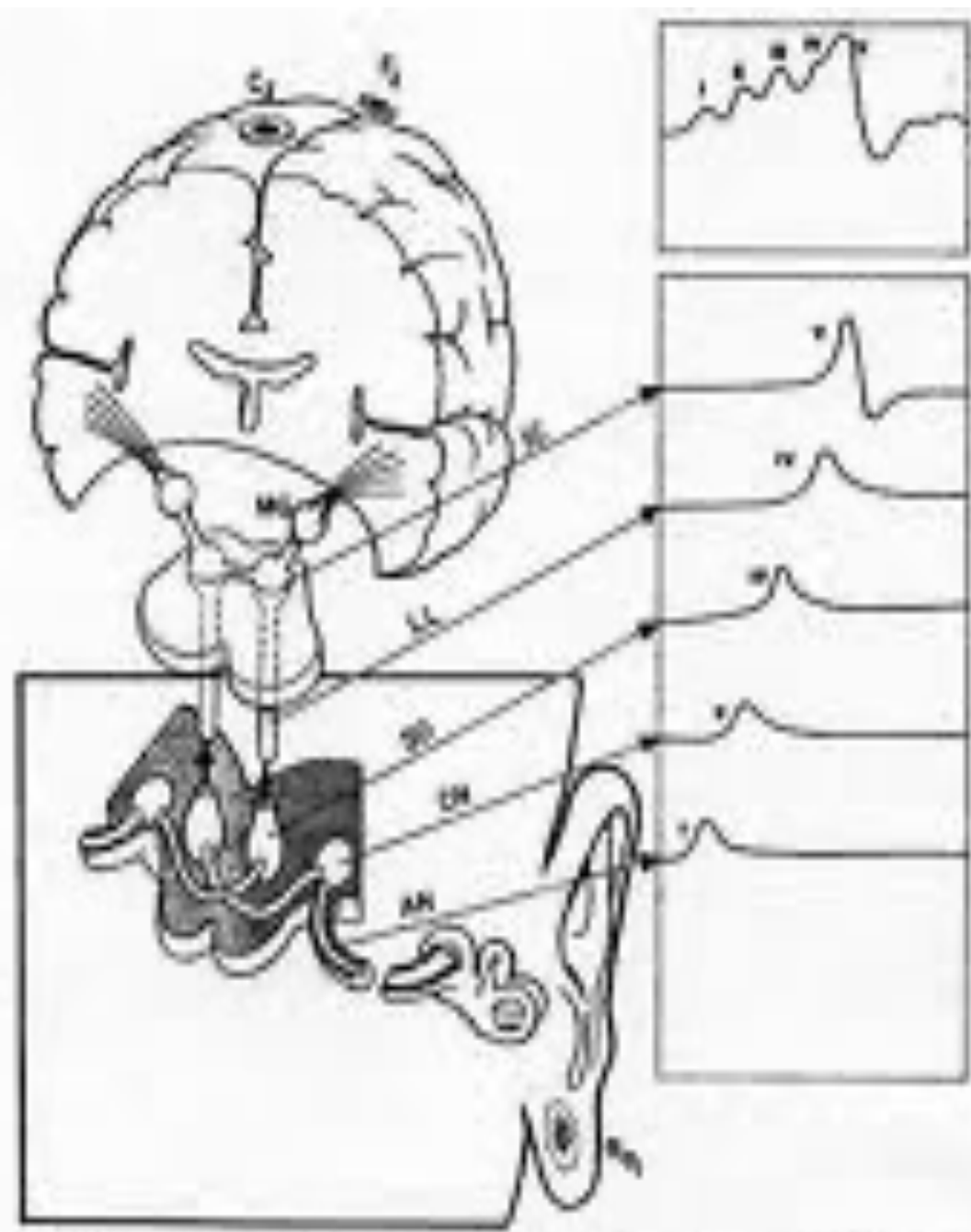
Tympanometry

- **Absence of emissions** (sebaceous plug in the ear canal, middle ear ossicles changes, fluid in the middle ear)
- perform tympanometry-**objective examination**
- examination of **eardrum compliance**, which is dependent on the ratios of the middle ear
- for the accuracy of the TEOAE result is needed **curve A**.



BERA

- Repeatedly **absented** TEOAE, Tympanometry **A**
- **suspected** hearing loss incurred in the cochlea
- BERA (Brainstem Evoked Responses Audiometers)
- **objective** audiological test
- records the **electrical potentials of the brain stem**, which arise as a brain response to sound signals coming from the surroundings
- Headphone tones are bringing the patient to which a hearing brainstem patient responds to electrical activity (recorded by the probe attached to the patient's head)



rehabilitation of hearing

- first hearing aids (from 3 months)
- on both ears
- effect is considered ideally around the 6th – 9th months of age
- inadequate compensation for hearing loss with hearing aids, cochlear implant
- „Learn to hear“

First Moravian Centre for cochlear implantation

- 19th july 2012
- It is intended for patients with severe sensorineural hearing loss
- allow patients to understand speech

First Moravian Centre for cochlear implantation

- It was established at the University Hospital St. Anna in Brno (adults) + FN Brno, KDORL (children)
- close cooperation with reputable implanting center in Vienna (Prof. univ. Dr. Wolf-Dietrich Baumgartner from Universitätsklinik für Hals-, Nasen-und Ohrenkrankheiten, Medizinische Universität Wien)

First Moravian Centre for cochlear implantation

- there is no economically challenging commute
- Brno center offers quality and subsequent operative, rehabilitation needs across the spectrum
satisfaction guarantee not only adults, but also implanted children and their parents.

1st implant child



