Hearing loss

Overview

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Not "deafness"

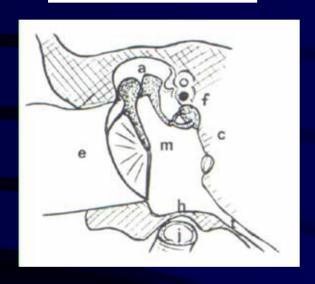
• Deaf is a total lack of hearing

• Deafness has connotations of discrimination

Word "deaf" frightens people

Hearing loss (HL) - classification

Conductive



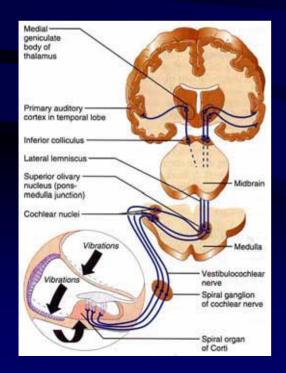
any interruption of passage of acoustic energy between pinna and oval window (CHL)

Sensorineural

Hearing loss (HL) - classification

Conductive

• Sensorineural



any damage to the organ
 of Corti or VIII nerve
 pathway or auditory cortex
 (SNHL)

Hearing loss (HL) - classification

• Can be "mixed"

conductive & sensorineural

Testing

• Multitude of testing possibilities

- clinical (in room with patient)

- audiological (by audiologist or audiometrician)

Testing

- Clinical
 - tuning forks

simple, quick and accurate means of differentiating between CHL & SNHL and between true HL and malingering

Tuning forks

- T fork large so that rate of decay is not rapid
- 512 Hz (or 256 Hz)
- Broad base applied to bone

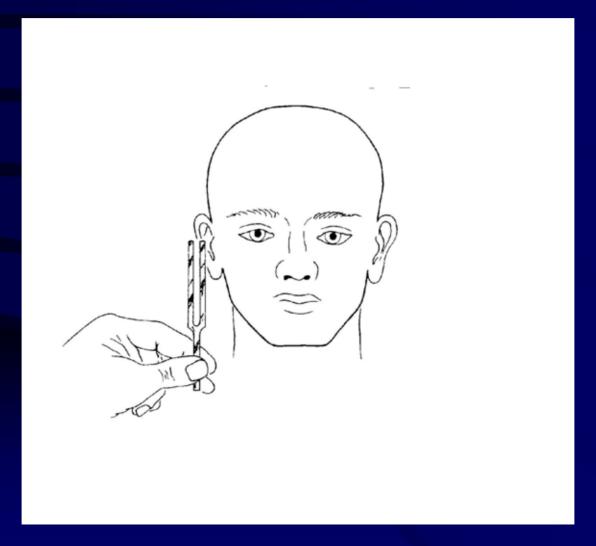
- If frequency too low felt more than heard
- If frequency too high dissipates too fast

Rinne

 Tests air conduction – much more efficient than BC

- + Rinne
- - Rinne
- false Rinne

Tuning fork - Rinne



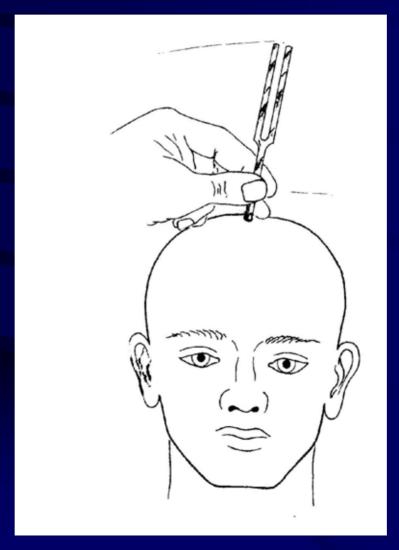
Weber

Assymetrical conductive loss

• Very sensitive in compliant patient

• If CHL unilateral as little as 5 dB detected

Tuning fork - Weber

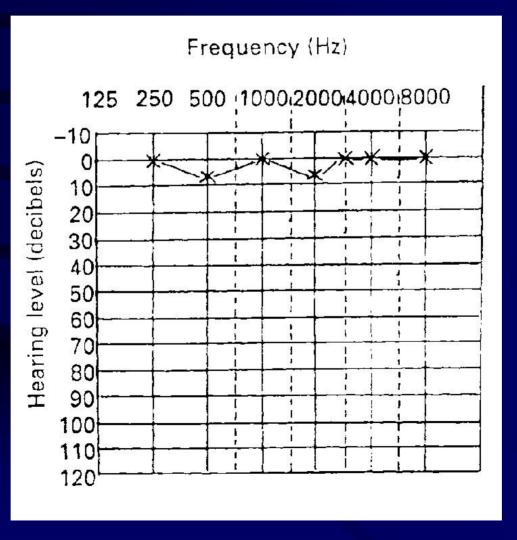


Testing

- Clinical
 - free-field speech testing

Simple, rapid, accurate to within 3 dB of true threshold of speech reception threshold

Normal audiogram



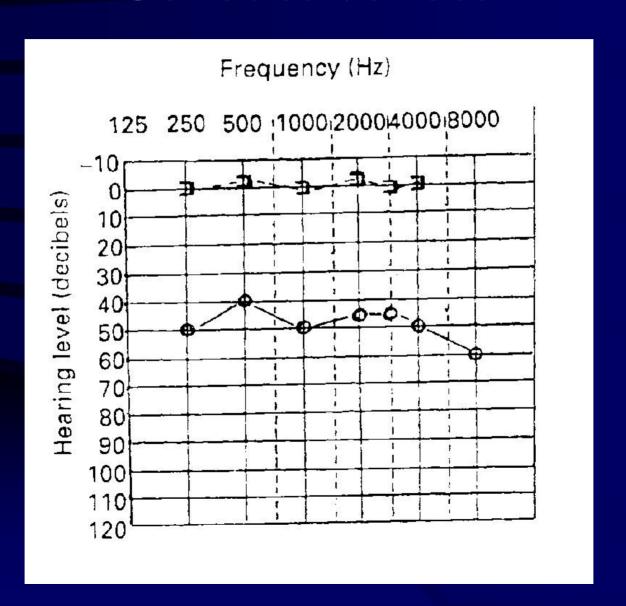


Conductive HL

 Congenital – many syndromes affecting the development of ext. ear from 1st branchial cleft and 1st & 2nd branchial arches

(eg Treacher-Collins, Pierre-Robin, Crouzon's, Apert's)

Conductive loss



Congenital CHL classification

Hereditary

- EA or ME
- present at birth (syndromes)
- appearing in childhood (osteogenesis imperfecta)
- Predisposing disorders
 - cystic fibrosis, cleft palate,
 Down syndrome
- Miscellaneous
- congenital cholesteatoma
- fibrous dysplasia

Congenital CHL

- Minor aplasia EAC narrow
 - pinna normal or minor deformity
 - ossicular fixation

Congenital CHL

- Major aplasia microtia
 - EAC atresia
 - ossicular fixation

Congenital CHL

- Major aplasia / atresia EAC atretic
 - tympanic cavity small
 - cochlear abnormal

Congenital CHL - management

 Congenital loss may be suspected at birth (eg family history or syndromal)

- Refer to Paediatric ENT or Audiologist for:
 - investigation
 - appropriate surgery
 - appropriate rehabilitation

Conductive HL

Acquired

- OME/ effusions
- foreign body / wax
- perforation
- ossicular damage
- otitis externa / furuncle
- fracture
- EAC neoplasm benign
 - malignant
- atresia and stenosis

Acquired CHL - management

From history and examination,
 diagnose and treat appropriately

Sensorineural HL

Congenital – many syndromes involve the
 VIII nerve to varying degrees

(eg Pendred's, Usher's Waardenbur's)

Congenital SNHL classification

- Hereditary HL present at birth HL alone
 syndrome + HL
 - HL appears in childhood
 - HL alone
 - syndrome + HL
- Secondary to intrauterine event
 - Infections (rubella, CMV, syphilis)
 - Ototoxic drugs (aminoglycosides, diuretics, cytotoxics, salicylates, quinine, anticonvulsants)

Congenital SNHL classification

Secondary to intrauterine event

- Metabolic disorders (diabetes mellitus)

Perinatal disorders (hypoxia,
 hyperbilirubinaemia,
 premature delivery,
 low birth weight)

Congenital SHL

- Variable uni- or bilateral
 - mild
 - moderate
 - severe
 - profound

Congenital SHL - management

Prevent

Depends on cause

Depends on extent

• Depends whether uni- or bilateral

Congenital SHL - management

• Hearing amplification - if appropriate

Alternative means of communication

- lip reading

- sign language

Cochlear implant

- if appropriate

Congenital SHL - management

• Family support and advice

Sensorineural HL

Acquired

- trauma sharp/blunt
- acoustic blast / noise
- barotrauma window rupture
- surgery
- infective labyrinthitis
- syphilis
- Meniere's
- presbycusis
- ototoxicity
- CVA

Acquired SHL - management

• Prevent

Depends on cause

Depends on extent

• Depends whether uni- or bilateral

Acquired SHL - management

• Hearing amplification - if appropriate

Alternative means of communication

- lip reading

- sign language

Cochlear implant

- if appropriate

Acquired SHL - management

• Family support and advice