

SCREENING for HEARING LOSS in NEONATES

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INTRODUCTION

Congenital permanent hearing impairment

- common neurosensory handicap in newborn & children
- prevalence differs from country to country
 - estimated prevalence internationally, 1.1 – 1.5 per 1 000 live births
- more prevalent than commonly screened medical conditions
 - hypothyroidism (25/100 000)
 - haemoglobinopathies (13/100 000)
 - phenylketonuria (7/100 000)
- no Malaysian data - estimated 840 babies annually
 - 30% cases associated with other neurodevelopmental conditions, mental retardation
 - 10-20 times higher in high risk babies than in general population

INTRODUCTION

- carried out technology assessment
- to determine
 - accuracy of screening test
 - effectiveness of screening
 - effectiveness of early detection
 - cost-effectiveness of a newborn hearing screening programme
 - types of audiologic interventions
 - organisational implications

TECHNICAL FEATURES

Types of tests...

- **Otoacoustic emission (OAE)** - integrity of inner ear
- **Auditory brainstem response (ABR)** - identifies normal cochlear function but abnormal VIII cranial nerve function
- **Automated ABR (AABR)** - similar mechanism to ABR, but fully automated; allows variety of trained personnel to use it

Accuracy of screening tests

- **OAE** – quick, easy, inexpensive, referral rate 34-44%, sensitivity 78-99%, specificity 90-99%
- **AABR** – slower (8-20min), > complicated, > expensive, referral rate 0.2-2.5%, sensitivity 96-99%, specificity 99-100%

Screening test...

- **to reduce false positives 2 step approach**
 - OAE done twice
 - OAE followed by ABR
 - AABR for those failing 1st test
- **Accuracy of tests affected by** - child crying, middle ear effusion, external ear debris

EFFECTIVENESS OF SCREENING

- Lack of overt physical signs - identification is late without screening programme
- Age of detection
 - 42.0 months [UK, 1997]
 - 3 years [National Institute of Health Panel, 1993]
 - 14.5 months [National Institute of Health Panel, 1997]
 - 48.0 months [National University Malaysia 1990]
(high risk children – 41 months; children without risk – 53 months)

results...

SCREENING UNIVERSAL vs HIGH RISK

- Most authorities endorse **universal screening programme**
 - National Institute of Health, USA; Task Force on Newborn and Infant Hearing, American Academy of Pediatrics; US Joint Committee on Infant Hearing; Center of Disease Control and Prevention, US; Canadian Task Force on Preventive Health Care; UK National Screening Committee
- **US Preventive Services Task Force** - insufficient evidence to recommend for or against Universal Newborn Hearing Screening
- **Swedish HTA agency** - insufficient cost effective data on universal screening

High risk screening

- Yield of screening & proportion of true positive results substantially higher
- may miss up to 50% of 'low risk' infants with congenital hearing loss

ORGANISATIONAL ASPECTS

PRE-REQUISITES FOR A SCREENING PROGRAMME

- Well coordinated & closely knit services provision
- Availability of diagnostic services
- Availability of diagnostic facilities for referral
- Availability of audiologists (1:25 000 population) & speech language pathologists (1:10 000 population)

Local scenario - audiologists **1:370 000** population;
speech pathologists **1: 275 000** population
(60 audiologists; 80 speech language pathologists)

ORGANISATIONAL ASPECTS

2004

- Audiology services
 - in 26 public hospitals
- Speech pathology services
 - in 22 public hospitals

2003

- Equipment for diagnostic assessment
 - available in 13/16 hospitals that offer audiology services
- Equipment for fitting hearing aids
 - available in 19 public hospitals
- Rehabilitation services
 - Confined to urban areas
- Cochlear implants – University hospitals

RECOMMENDATIONS

- Targeted hospital based (high risk babies) screening program for hearing loss in hospitals where diagnostic & rehabilitative services available
- Universal screening in future