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Early hearing detection and intervention (EHDI) programmes for infants and young children in low-income and middle-income countries in Asia: a systematic review

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ABSTRACT

Background Early hearing detection and intervention (EHDI) measures initiated in high-income countries (HICs) were attempted in low-income and middle-income countries (L&MICs). However, information regarding the models of EHDI, context-specific adaptations made to strategies and outcomes are not known.

Aims The aims of this systematic review were to identify the various models of EHDI used in Asian L&MICs in the published scientific literature and to describe their efficacy and validity.

Methods The studies were eligible if the programme was from Asian L&MICs, implemented for children below 6 years of age and published between 2010 and 2021. Google Scholar, PubMed, Web of Science, Scopus, EBSCOHost and EBSCO–CINAHL were used to find articles. Data were extracted from each selected article, and the risk of bias was assessed. The search results were summarised using the Preferred Reporting Items for Systematic Reviews and Meta-Analyses flow diagram. For primary outcomes, narrative synthesis was used, and forest plots were generated for secondary outcomes.

Results In all, 82 studies were included, and these studies were divided into two categories: newborn and infant screening programmes and screening programmes for older children. Predominantly, a two-stage objective otoacoustic emission (Distortion Product/Transient Evoked) or automated auditory brainstem response screening, followed by a detailed auditory brainstem response to confirm the hearing loss, was used in newborn and infant screening programmes. Audiologists were the most frequent screening personnel. Screening of older children was mostly done by otolaryngologists, school instructors and nurses. They performed a singlestage pure tone audiometry screening followed by a detailed examination.

Conclusion The screening tools and protocols used were similar to those used in HICs. However, no uniform protocols were followed within each country. Long-term viability of EHDI programmes was not known as there was limited information on impact outcomes such as cost–benefit.

PROSPERO registration number CRD42021240341.

WHAT IS ALREADY KNOWN ON THIS TOPIC

⇒ Early hearing detection and intervention (EHDI) programmes are mandated in several high-income countries (HICs) for over two decades. These screening programmes are based on guidelines and standards provided by the Joint Committee on Infant Hearing, the American Audiology Association, the Newborn Hearing Screening Programme England, WHO, the European Consensus Statement on Neonatal Hearing Screening, etc. Systematic reviews have documented screening protocols and programme outcomes predominantly in the context of HICs.

WHAT THIS STUDY ADDS

⇒ Unlike several HICs, EHDI programmes are not mandated in many low-income and middle-income countries (L&MICs). In this context, we conducted a systematic review and gathered information on hearing screening programmes mainly to identify different models of EHDI that were implemented in the context of Asian L&MICs. This review provides information on various screening protocols, tools, personnel, diagnostic tools, use of information and communication technology, barriers and facilitators in different EHDI programmes of L&MICs.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

⇒ We found that the screening tools and protocols used were similar to those used in HICs, yet no uniform protocols were followed within each country. Long-term viability of EHDI programmes is not known in this context due to limited impact outcome-based studies(eg, costbenefit, rate of intervention, etc); hence, future research should focus on these aspects. Further, policy makers and programme planners in these countries should build consensus to implement uniform countrywise protocols suited to the context.

INTRODUCTION

Currently, 34 million children below 15 years are estimated to have hearing loss, with a higher prevalence in low-income and middleincome countries (L&MICs) (2.4%) than in

| | | Validity of | the screen | ing program | Efficacy of | f the screenii | ng program |
|--|------------------------|---|---|---|-----------------------------------|-----------------------------|------------------------------|
| Citation | Country | Use of validated screening tools | Valid diagnostic testing tools within the scope of the program | Implementatio n phase Outcome of HL identification/i ntervention or Impact evaluation/> 2 years duration | Program's economic analysis | Frequency of identification | Frequency of intervention |
| Biswas et al., 2012 | India | | | | | | |
| Paul et al.,2011 Sharma et al., 2013 | India India | | | | | | |
| Ramesh et al., 2013 | India | | | | | | |
| Rai & Takhur et al.,2013 | India | | | | | | |
| Kumar et al.,2015 | India | | | | | | |
| Gupta et al., 2015 | India | | | | | | |
| Vignesh et al.,2015 Vishwakarma et al.,2015 | India India | | | | | | |
| Paul et al.,2016 | India | | | | | | |
| Kumar et al., 2016 | India | | | | | | |
| Bhat et al.,2018 | India | | | | | | |
| Sachdeva & Sao et al.,2017 | India | | | | | | |
| Kumar et al.,2017 | India | | | | | | |
| Swain et al.,2017 | India | | | | | | |
| Bhat et al.,2018 Bishnoi et al.,2018 | India India | | | | | | |
| Bishnoi et al.,2018 Parab et al.,2018 | India | | | | | | |
| Jacob et al.,2020 | India | | | | | | |
| Nishad et al.,2020 | India | | | | | | |
| Sija et al., 2022 | India | | | | | | |
| Zhang et al., 2012 | China | | | | | | |
| Huang et al.,2012 Chen et al.,2012 | China China | | | | | | |
| Shang et al.,2012 | China | | | | | | |
| Wenjin et al., 2018 | China | | | | | | |
| Wang et al., 2019 | China | | | | | | |
| Dai et al.,2019 | China | | | | | | |
| Zeng et al., 2020 | China | | | | | | |
| Wen et al., 2020 | China China | | | | | | |
| Guo et al., 2020 Guomei et al.,2022 | China China | | | | | | |
| Ahmad et al.,2011 | Malaysia | | | | | | |
| Wong et al.,2020 | Malaysia | | | | | | |
| Mazlan et al.,2022 | Malaysia | | | | | | |
| Tasci et al.,2010 | Turkey | | | | | | |
| Sennaroglu & Akmese, 2011 | Turkey | | | | | | |
| Ulusoy et al.,2014 Kemaloğlu et al., 2016 | Turkey Turkey | | | | | | |
| Kemaloğlu et al., 2016 Yorulmaz et al., 2017 | Turkey | | | | | | |
| Celik et al.,2016 | Turkey | | | | | | |
| Ozturk et al.,2017 | Turkey | | | | | | |
| Hamdi, 2018 | Turkey | | | | | | |
| Yücel et al., 2019 | Turkey | | | | | | |
| Arslan et al., 2013 | Turkey | | | | | | |
| Çıkrıkçı et al., 2020 Arjmandi et al., 2012 | Turkey Iran | | | | | | |
| Islami et al.,2013 | Iran | | | | | | |
| Firoozbakht et al.,2014 | Iran | | | | | | |
| Zahed et al., 2014 | Iran | | | | | | |
| Farhat et al., 2014 | Iran | | | | | | |
| Haghshenas et al., 2014 | Iran | | | | | | |
| Baradaranfar et al., 2014 | Iran | | | | | | |
| Azizi et al., 2016 Tajik & Ahmadpour-Kacho, 201 | Iran Iran | | | | | | |
| Saki et al.,2017 | Iran | | | | | | |
| Rahimi et al.,2018 | Iran | | | | | | |
| Thungvachirakul et al.,2011 | Thailand | | | | | | |
| Poonual et al.,2016 | Thailand | | | | | | |
| Poonual et al., 2017b | Thailand | | | | | | |
| Poonual et al.,2017 | Thailand | | | | | | |
| Pitathawatchai et al.,2019 | Thailand | | | | | | |
| Ray et al.,2021 | Nepal | | | | | | |
| Shameem et al.,2022 Khaimook et al.,2022 | Bangladesh Thailand | | | | | | |
| ritainiook et al.,2022 | maliand | | | | | | |

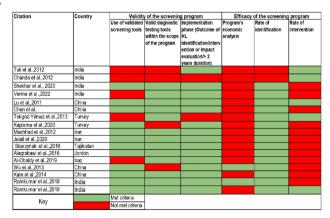


Figure 1 Validity and efficacy of screening programmes (A) for newborns and infants and (B) for older children. HL, hearing loss.

high-income countries (HICs) (0.5%).¹ Early hearing detection and intervention (EHDI) for children with hearing loss is critical to maximise linguistic competence and literacy development. EHDI is a concept that emanated in the USA in the 1990s and is intended as an at-birth hearing screening of newborns prior to hospital discharge. Infants who do not pass the screening are recommended for diagnostic evaluation and, when confirmed to have hearing loss, are enrolled in early intervention programmes. Subsequently, the Joint Committee on Infant Hearing (2007) in the USA recommended that all infants be screened for hearing by 1 month of age and diagnosed by 3 months and receive intervention by 6 months of age.² It is practised as a mandatory universal screening in the entire country.

The concept was subsequently adopted in the UK and practised as universal screening since 2006. Subsequently, several other HICs (Australia and Canada, to name a few) adopted this strategy. Alternative strategies for EHDI have been implemented in L&MICs due to financial, human resource and infrastructural challenges.³ These include high risk-based screening,⁴ screening during immunisation,⁵ community-based hearing screening by health workers^{6 7} and school entry-level screening.^{8 9} Several of these programmes have also integrated telepractice to either improve coverage of screening or provide better diagnostic follow-up.^{10 11} However, there remains a lack of clarity on the range of strategies implemented in L&MICs and which should be promoted.

The aims of this systematic review were to identify different models of EHDI that have been implemented in the context of Asian L&MICs in the published scientific literature and to describe evidence of their efficacy and validity.

METHOD

The protocol for this systematic review was registered in the International Prospective Register of Systematic Reviews (registration number CRD42021240341).

Patient and public involvement statement

This systematic review did not involve any subject/patient and public directly.

Inclusion criteria

All types of study designs were eligible for this review, including (1) cross-sectional, (2) cohort, (3) casecontrol, (4) randomised controlled trials, (5) quasiexperimental and (6) field trials. Both qualitative and quantitative types of studies were included.

The EHDI model is operationally defined for the purpose of this systematic review as programmes for identification and referral of young children with hearing loss. Studies that described EHDI programmes related to triaging children suspected with hearing loss using methods such as objective or subjective screening, parental questionnaire-based screening, implemented

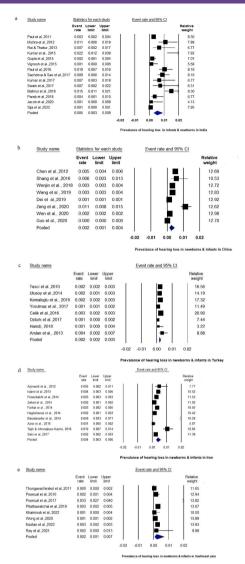


Figure 2 Forest plot of prevalence of hearing loss in (A) newborns and infants in India; (B) newborns and infants in China; (C) newborns and infants in Turkey; (D) newborns and infants in Iran; and (E) newborns and infants in other Asian countries (Thailand, Malaysia and Nepal).

in the context of low-income countries (LICs), lower middle-income countries (LMICs) and upper middleincome countries (UMICs) including hospital, community, school based or any other alternative approach were included.

Studies were eligible regardless of screening strategies (eg, at birthing hospital/community/school), protocol used (eg, single stage/two-stage), provider stakeholder (eg, private/public) involved, tools for screening (eg, checklist, otoacoustic emission (OAE), automated auditory brainstem response (AABR) etc), or personnel involved in screening, diagnosis and intervention (eg, nurse, audiometrists, audiologists and ENT). We also included studies that explored evidence of validity (eg, sensitivity/specificity) and reported implementation barriers and facilitators to EHDI.

According to World Bank classification (2021), LICs, LMICs and UMICs (L&MICs) in the Asian continent (South East Asia, Central Asia and Western Asia/Middle East) were considered as eligible for the review. In the L&MICs, 6 years and below was predominantly considered as the age band for 'early' detection and intervention. Therefore, this review included studies describing EHDI among neonates, infants and children below 6 years of age. Studies were eligible if they had been published from 2010 to 2022.

Exclusion criteria

We excluded studies that described hearing screening programmes for individuals older than 6 years of age or for other disabilities not including hearing. In addition, studies from HICs, studies published in languages other than English and studies published before the year 2010 were excluded.

Search strategy

Since EHDI is an interdisciplinary programme often implemented by ENT/paediatrics/neonatology/audiology/nursing, databases that captured articles from multiple disciplines was preferred. The primary databases used for the search include PubMed, Scopus, Web of science, EBSCOHost, EBSCO–CINAHL (humanities and social sciences) and Google scholar. Hand searching was conducted for the *International Journal of Audiology* (2015–2022) and bibliographies of the selected papers based on the eligibility criteria. Grey literature search included ProQuest Dissertations & Theses Global (nterdisciplinary) and first 500 searches for articles/reports in Google Search. We excluded social media articles, newspaper articles, editorials and website information.

A search strategy for each of the aforementioned databases was designed using 2Dsearch online tool.¹² The search strategy included Medical Subject Headings terms and Boolean operators . A pilot search was conducted in each database to identify the keywords. Synonyms of the keywords were then identified and included in the search strategy.

Screening for eligibility and quality

Title screening was conducted as per the inclusion and exclusion criteria using database search. The Rayyan software¹³ was used to screen the abstract and full texts. Screening was conducted by two reviewers (DJ and VR), and any discrepancies were discussed between the reviewers and decisions were made. Joanna Briggs Quality assessment tools specific to the research design were used to assess the quality of the articles.

Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) flowchart¹⁴ was used to represent the search results.

Data extraction and synthesis

A Google Sheet was used for data extraction, which was undertaken by two authors (DJ and LSN) and verified by another author (VR).

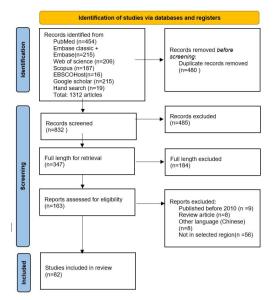


Figure 3 Preferred Reporting Items for Systematic Reviews and Meta-Analyses flowchart representing the selection of article at each stage.

Narrative synthesis of available data was conducted using textual approach to describe strategies adopted for EHDI including screening methods, service delivery points, use of information and communications technology (ICT), the target age groups of such programmes, personnel involved in delivery of the programme, and reported barriers and facilitators of the programme. The Joanna Briggs Institute (JBI) tool for critical appraisal¹⁵ was used for quality assessment. The Synthesis Without Meta-analysis (SWiM) guideline was used for analysis of secondary outcomes.¹⁶ If a country had at least three studies that reported data on children with confirmed hearing loss, then that country was included for estimation of prevalence per 1000 using forest plots.

The primary outcomes of interest were the validity and efficacy of the screening programmes. We developed a checklist (figure 1A,B) to assess the validity and efficacy using three criteria each. The items in the validity checklist included (1) the use of a *validated screening tool*, (2) the use of a *validated diagnostic tool*, whether the screening programme reported was in the (3) *design phase* (eg, pilot/feasibility/ validity/only reported coverage rate or referral rate or follow-up rate) or *implementation phase* (eg, scale programme). The efficacy was assessed if the study reported (1) evidence of early identification, (2) evidence of early intervention and (2) inclusion of an economic analysis.

The secondary outcome of interest was to estimate the incidence and prevalence outcomes of EHDI programmes in the Asian L&MICs. For secondary outcomes analysis, in screening programmes for newborns and infants, the prevalence of hearing loss in infants reported in each country was analysed using the SWiM guidelines. Using a random effect model, Forest plots (figure 2A–E) were

constructed for each country based on two criteria: if more than five studies in a country reported prevalence outcomes and if the number of children screened was more than 1000.

RESULTS

Our electronic search yielded 1312 citations. Based on the inclusion/exclusion criteria and multiple levels of screening by the two reviewers independently, a total of 82 studies qualified for the current review. The article selection process is presented in the PRISMA flowchart (figure 3). Sixty-five studies (79%) reported on newborn hearing screening (NHS), and only 17 studies (21%) reported hearing screening among older children. Predominantly, studies were conducted in India (n=27), followed by Turkey (n=13), Iran (n=13), China (n=15), Thailand (n=6), Malaysia (n=3), Nepal (n=1), Bangladesh (n=1), Iraq (n=1), Jordan (n=1) and Tajikistan (n=1).

These studies included 75 cross-sectional studies and 7 cohort studies. Results of quality appraisal using appropriate JBI tool are provided in online supplemental file 1.

The screening programmes identified in this review were grouped based on the age group of the children: (1) screening programmes for newborns and infants (0–3 years of age) and (2) screening programmes for older children even beyond 6 years of age.

Hearing screening programmes for newborns and infants (below 2 years) included 65 studies. Most studies (49) reported single-hospital programmes, whereas others (16 studies) reported multiple-centre programmes. Of these studies, 55 were undertaken in the private sector and 10 in the public sector. There were 17 studies of hearing screening programmes for older children aged 3-17. Fifteen of these studies were schoolbased hearing screenings, while two were communitybased. Of these studies, nine were undertaken in the private sector and eight in the public sector. Table 1A-E represents the summary of included studies describing hearing screening programmes for newborns and infants in each country. Table 2 represents the summary of included studies hearing screening programmes for older children.

Screening protocol and tests

Newborn and infant hearing screening

Two-stage hearing screening protocols were employed most frequently for newborn and infant hearing screening (n=47), followed by three-stage protocols (n=13) and onestage protocols (n=4). One study reported employing a five-step hearing screening protocol.

Sixteen studies that reported a two-stage hearing screening protocol, employed OAE (TE/DP-OAE) or AABR as screening tests (individually or combined in either stage).¹⁷⁻³¹ The other 25 studies used only OAEs (DP/TE)³²⁻⁴⁹ or AABR screening^{50 51} for testing in both stages. Those studies that reported the use of AABR in

| (A) | | | | | | | | |
|----------------------------------|----------|--------------------------|------------------------------|--|---|--|--|---------------------------------|
| Author and year | Citation | Duration of programme | Population screened | Screened (n) | Screening protocol | Screening test used | Screening personnel | Diagnostic test |
| Biswas <i>et al</i> 2012 | 104 | 2 years | Newborns | 490 | 1 stage | DPOAE | Not mentioned | Not mentioned |
| Paul 2011 | 18 | 7 years | Newborns | 10165 | 2 stage | OAE+OAE (non-mentioned DP/TE) | Person with basic knowledge in computer with training on NHS | Diagnostic ABR |
| Mishra et al 2013 | 32 | 3 years | 0–2 years | 1101 | <6 months of age, 5 stage; 6 months-1 year, 4 stage; 1–2 years, 3 stage | DPOAE | Not mentioned | Diagnostic ABR |
| Ramesh <i>et al</i> 2012 | 85 | 2 years | Newborns | 425 | 1 stage | Calibrated noise maker-based Behavioral Observation Audiometry (BOA) | Trained health workers (30 hours of training) | Diagnostic ABR, OAE and BOA |
| Rai <i>et al</i> 2013 | 83 | 1 year | Newborns | 500 | 3 stage | TEOAE+TEOAE+TEOAE | ENT Specialist (Ear Nose Throat) | Diagnostic ABR |
| Kumar <i>et al</i> 2015 | 17 | 1 year 8 months | High risk <2 years of age | 500 | 2 stage | TEOAE+AABR | Audiologist | Not mentioned |
| Gupta <i>et al</i> 2015 | 50 | 1 year | Newborns | 2265 | 2 stage | AABR+AABR | Single specialist staff | Not mentioned |
| Vignesh <i>et al</i> 2015 | 18 | 1.5 years | Newborns | 1405 | 2 stage | TEOAE+AABR | Not mentioned | Diagnostic ABR |
| Vishwakarma <i>et al</i> 2015 | 54 | 1 year 8 months | Newborns | Well babies: 2000 High risk:1020 | 3 stage | TEOAE+TEOAE+AABR | Nurse, resident doctor/ certified audiologist | Diagnostic ABR |
| Paul <i>et al</i> 2016 | ŝ | 11 years | Newborns | Well babies: 84 <i>7</i> 74 High risk: 16914 | 2 stage | OAE+OAE (non-mentioned DP/TE) | Person with basic training in hearing screening | Diagnostic ABR |
| Sharma <i>et al</i> 2018 | 86 | 3 years | Newborns | 2534 | 2 stage | DPOAE | Not mentioned | Diagnostic ABR |
| Kumar <i>et al</i> 2016 | 105 | 2 years | Newborns | 1537 | 2 stage | TEOAE+TEOAE+AABR | Not mentioned | Not mentioned |
| Sachdeva <i>et al</i> 2017 | . 87 | 10 months | Newborns | 2254 | 2 stage | (HRR+BOA+DPOAE)+DPOAE | Not mentioned | Confirmatory, diagnostic ABR |
| Kumar <i>et al</i> 2017 | 82 | No info | Newborns | 600 | 2 stage | TEOAE+DPOAE | Not mentioned | Not mentioned |
| Swain <i>et al</i> 2017 | 88 | 1.5 years | Newborns | 410 | 2 stage | DPOAE+DPOAE | Not mentioned | Diagnostic ABR |
| Bhat <i>et al</i> 2018 | 19 | 1 year | High-risk newborns | 195 | 1 stage | TEOAE | Not mentioned | Diagnostic ABR |
| Bishnoi <i>et al</i> 2018 | 88 | No info | Newborns | 2000 | 2 stage | (OAE and Tymp)+OAE (non- mentioned DP/TE) | Not mentioned | Diagnostic ABR |
| Parab <i>et al</i> 2018 | 34 | 3 years | Newborns | 8192 | 2 stage | TEOAE+TEOAE | Audiologist | Diagnostic ABR |
| Jacob <i>et al</i> 2020 | 35 | 2 years | Newborns | 773 | 2 stage | TEOAE+TEOAE | Not mentioned | Diagnostic ABR |
| Nishad <i>et al</i> 2020 | 36 | 1 year | Newborns | 1000 | 2 stage | OAE+OAE (non-mentioned DP/TE) | Not mentioned | Diagnostic ABR |

| Table 1 Continued (A) (A) | pel | | | | | | | |
|---|----------|--------------------------|------------------------|---------------|------------------------------|--|-------------------------------|---|
| Author and year | Citation | Duration of programme | Population screened | Screened (n) | Screening protocol | Screening test used | Screening personnel | Diagnostic test |
| Sija et al 2022 | 37 | 4 years | Newborns | 16265 | 2 stage | DPOAE+DPOAE | Trained nurse | Diagnostic ABR |
| (B) | | | | | | | | |
| | : | | Population | | | | | |
| Author and year | Citation | programme | screened | Screened (n) | Screening protocol | Screening test used | Screening personnel | Diagnostic test |
| Zhang <i>et al</i> 2012 | 106 | 1.5 years | Newborns | 10 043 | 2 stage+genetic screening | TEOAE+(TEOAE and AABR) | Nurse | Not mentioned |
| Tobe <i>et al</i> 2013 | 83 | 2 years | Newborns | Not mentioned | 2 stage | OAE+AABR (non-mentioned DP/TE) | Trained personnel, no info | Not mentioned |
| Chen <i>et al</i> 2012 | 8 | 2 years | Newborns | 11568 | 2 stage | TEOAE | Audiologist | Diagnostic ABR, TFT, impedance, ASSR at hospital |
| Shang <i>et al</i> 2016 | 52 | 6 months | Newborns | 1064 | 2 stage | First protocol: TEOAE+TEOAE Second protocol: (TEOAE and ABR screen)+TEOAE | Not mentioned | Diagnostic ABR |
| Wenjin <i>et al</i> 2018 | 50 | 2 years | Newborns | 19 098 | 2 stage | Well babies: DPOAE+ABR screening High-risk babies: (DPOAE and ABR screening) (DPOAE and ABR screening) | Nurse | Otoscopy, diagnostic ABR at 30 dBHL, Tymp; DPOAEs |
| Wang et <i>al</i> 2019 | 21 | 5 years | Newborns | 55977 | 2 stage | OAE+AABR (non-mentioned DP/TE) | Nurse | Comprehensive diagnostic audiometry around 3 months of age |
| Dai <i>et al 2</i> 019 | 53 | 1 year | Newborns | 180469 | 2 stage+genetic screening | TEOAE+(TEOAE and AABR) | Not mentioned | Diagnostic ABR, ASSR, DPOAE, immitance |
| Zeng <i>et al</i> 2020 | 53 | 1 year | Newborns | 4205 | 2 stage+genetic screening | OAE+AABR screening (non- mentioned DP/TE) | Not mentioned | No |
| Wen <i>et al</i> 2020 | 24 | 2 years | Newborns | 467 980 | 2 stage | OAE+(OAE and AABR) (non- mentioned DP/TE) | Not mentioned | Not mentioned |
| Guo <i>et al 2</i> 020 | 25 | 2 years 4 months | Infants >3 months | 287 430 | 2 stage+genetic | OAE+AABR (non-mentioned DP/TE) | Not mentioned | Diagnostic ABR |
| Guomei <i>et al</i> 2022 | 53 | 9 months | Newborns | 2174 | 2 stage+genetic | OAE+OAE (non-mentioned DP/TE) | Not mentioned | Diagnostic ABR |
| | | | | | | | | Continued |

| (c)(| Table 1 Continued | ued | | | | | | | | |
|---|-------------------------------------|----------|--------------------------|--------------------------|---------------------------|-----------------------|-----------------------|---------------------------------------|---|---|
| Image: constraint of the second se | (C) | | | | | | | | | |
| II ⁴⁴ Malaysia (MIC) Syears Newborns 1600 3 stage DPOAE-DPOAE Table in a data data data data data data data d | Author and year | Citation | Country | Duration of programme | Population screened | Screened (n) | Screening protocol | Screening test used | Screening personnel | Diagnostic test |
| 1 0 Malaysia 2 years Newborns 2 stage 2 stage 2 stage 2 stage 2 stage 2 stage 2 strage 2 strand 2 stran | Ahmad <i>et al</i> 2011 | 64 | Malaysia (MIC) | 5 years | Newborns | 16 000 | 3 stage | DPOAE+DPOAE+DPOAE | Technician, staff nurse, ward attendants | Diagnostic ABR |
| of Thailand Tytain Interflored OAE-OAE (not mentioned) Not mentioned 16 "U,MIC) montrix Newborns 3120 2 stage Automated Not mentioned Not mentioned 17 "U,MIC) montrix Newborns 3120 2 stage Automated Not mentioned Not mentioned 17 "U,MIC) montrix Newborns 3120 2 stage COBAF-ROAE Not mentioned Not mentioned 17 "U,MIC) montrix Newborns 3120 2 stage COBAF-ROAE Not mentioned Not mentioned 17 "U,MIC) tyear Newborns 5120 2 stage COAF-ABR Not mentioned Not mentioned 16 "U,MIC) tyear Newborns 5120 2 stage COAF+TEOAE Not mentioned Not mentioned 16 "U,MIC) tyear Newborns 1596 2 stage TEOAF+TEOAE Not mentioned Not mentioned 16 MU TEOAF Stage | Wong <i>et al</i> 2021 | 26 | Malaysia (UMIC) | 2 years | Newborns | 28 432 | 1 and 2 stage | 1 stage: AABR 2 stage: DPOAE+AABR | Nurses | Diagnostic ABR |
| 16 11 Trailand 1 werborns 12 1 stage 1 functional TEOAE Notmentioned 17 1 Thailand Iventioned 1 Newborns 1 1 mentioned Notmentioned 17 1 UMIC) Newborns 1 1 Newborns 1 1 Notmentioned Notmentioned Notmentioned 17 1 UMIC) Iver Newborns 1 2 | Tungvachirakul <i>et</i> al 2011 | 30 | Thailand (UMIC) | 1 year 11 months | Newborns | 4043 | 2 stage | OAE+OAE (not mentioned DP/TE) | Not mentioned | ASSR |
| 17 ¹² Thelland Net Net Net COBRAHR Net mentioned 11 ¹⁶ (MMC) mentioned 19er COBRAHR Not mentioned 11 ¹⁶ (MMC) 19er Newborns 3120 2 stage COBRAHR Not mentioned 11 ¹⁰ (MMC) 19er Newborns 3120 2 stage TEOAE+ABR Not mentioned 12 ¹⁰ (MMC) 19er Newborns 540 2 stage DAE+OAE (not mentioned Not mentioned 12 ¹⁰ Newborns 540 2 stage TEOAE+ABR Not mentioned 12 ¹⁰ Newborns 540 2 stage TEOAE+ABR Not mentioned 13 ¹⁰ Newborns 540 2 stage TEOAE+ABR Not mentioned 14 Newborns 109 2 stage TEOAE+ABR Not mentioned Not mentioned 14 Newborns 109 2 stage TEOAE+ABR Not mentioned </td <td>Poonual <i>et al</i> 2016</td> <td></td> <td>Thailand (UMIC)</td> <td>1 year 7 months</td> <td>Newborns</td> <td>3120</td> <td>2 stage</td> <td>Automated TEOAE+conventional TEOAE</td> <td>Not mentioned</td> <td>Diagnostic ABR</td> | Poonual <i>et al</i> 2016 | | Thailand (UMIC) | 1 year 7 months | Newborns | 3120 | 2 stage | Automated TEOAE+conventional TEOAE | Not mentioned | Diagnostic ABR |
| 17 % Thailand 1year Newborns 3120 2 stage TEOAE+ABR Not mentioned ef % Uuldic) 1year Newborns 6140 2 stage TEOAE+TEOAE Nurses 2 % Uuldic) 1year Newborns 6140 2 stage DOE+OAE (not mentioned Nurses 2 % Malaysia 10 years Newborns 50633 2 stage DOE+ABR Nurses 2 % Malaysia 10 years Newborns 50633 2 stage DOE+TEOAE Nurses 2 % Malaysia 10 years Newborns 50633 2 stage TEOAE+TEOAE Nurmetioned Nurmetioned 2 Bangladesh 2 years Newborns 50633 2 stage TEOAE+TEOAE Nurmetioned | Poonual <i>et al</i> 2017 | | Thailand (UMIC) | Not mentioned | Newborns | 3120 | 3 stage | COBRA HRR tool+TEOAE+AABR | Not mentioned | Not mentioned |
| of 140 19ar 7 Newborns 6140 2 stage TEOAE+TEOAE Nurses 1 Nead (MIC) 2 vars Newborns 540 2 stage DAF-ICS Not mentioned Not mentioned 2 2 Malaysia 10 vars Newborns 540 2 stage DAF-ICS Not mentioned Not mentioned 2 2 Malaysia 10 vars Newborns 50 633 2 stage TEOAE+TEOAE Not mentioned 2 2 Malaysia 10 vars Newborns 50 633 2 stage TEOAE+TEOAE Not mentioned 4 Liveic Newborns 1696 2 stage TEOAE+TEOAE Not mentioned Not mentioned 4 UUNC) Screened Screening test used Trained nurse and Motiongist Motiongist 4 14 motins Newborns 1896 Screening test used Motiongist centrice Motiongist centrice Motiongist centrice 5 10 vars Newborns 1840 Screening | Poonual <i>et al</i> 2017 | | Thailand (UMIC) | 1 year | Newborns | 3120 | 2 stage | TEOAE+AABR | Not mentioned | ABR at 3 and 8 months |
| 4 Nepal (LMIC) 2 years Newborns 540 2 stage OAE+OAE (not mentioned Not mentioned 2 2 Malysia 10 years Newborns 5053 2 stage TEOAE+TABR Trained nurses and istributioning ists 4 8 Malysia 10 years Newborns 5053 2 stage TEOAE+TEOAE Net mentioned 4 8 Malysia 2 years High-risk 426 2 stage TEOAE+TEOAE Not mentioned 4 8 Networns 1696 2 stage TEOAE+TEOAE Not mentioned Not mentioned 4 9 14mich Newborns 1696 2 stage TEOAE+TEOAE Not mentioned Not mentioned 4 14mich Newborns 1697 Screening test used Not mentioned No | Pitathawatchai <i>et</i> a/ 2019 | 40 | Thailand (UMIC) | 1 year 7 months | Newborns | 6140 | 2 stage | TEOAE+TEOAE | Nurses | Not mentioned |
| 2 ¹⁰ Malaysia 10 years Newborns 5633 2 stage TEOAE+ABR Trained nurses and medical technologits 1 (UMIC) (UMIC) Bangladesh 2 years High-risk house 456 2 stage TEOAE+TEOAE Not mentioned 1 1 (UMIC) 6 months 80% 1696 2 stage TEOAE+TEOAE Not mentioned 1 1 Notice 1696 Screened (M) 70% 2 stage TEOAE+TEOAE Not mentioned 1 Partition Revolution Screened (M) Protocol 2 stage Screening test used Screening test used Screening test used Screening test used 1 1 1 Nowborns 16975 3 stage TEOAE+TEOAE+ ABR Redionoptist or function 1 1 1 1 1 Screening test used Screening test used Screening test used Noticologist or function 1 1 1 1 1 Screening test used TEOAE+TEOAE+ ABR Noticologist or function 1 1 1 1 1 Screening t | Ray <i>et al</i> 2021 | 41 | Nepal (LMIC) | 2 years | Newborns | 540 | 2 stage | OAE+OAE (not mentioned DP/TE) | Not mentioned | Diagnostic OAE and diagnostic ABR |
| ⁴ Bangladesh (LMIC) 2 years ewborns High-risk ewborns 26 Stage (LMIC) TEOAE+TEOAE Net mentioned ⁴ Thaliand 6 months Newborns 1696 2 stage TEOAE+TEOAE Net mentioned r Thaliand 6 months Newborns 1696 2 stage TEOAE+TEOAE Net mentioned r tration of Borours 1697 Screening test used Net diologist of autoins <i>i</i> ⁴ 14 months Newborns 16975 3 stage TEOAE+TEOAE+ ABR Audiologist of autoins <i>i</i> ⁴ 14 state Newborns 1757 3 stage TEOAE+TEOAE+ ABR Audiologist of autoins <i>i</i> ¹ 10 state 1607 3 stage TEOAE+TEOAE+ ABR Audiologist of autoins <i>i</i> ¹ 10 state 1607 3 stage TEOAE+ABR Audiologist of autoins <i>i</i> 10 state 10 state 1607 3 | Mazlan <i>et al</i> 2022 | 28 | Malaysia (UMIC) | 10 years | Newborns | 50 633 | 2 stage | TEOAE+AABR | Trained nurses and medical technologists | Diagnostic ABR |
| 43Thailand (UMIC)6 months16962 stageTEOAE+TEOAETrained nurse and audiologist111111111111111111111112121414169753 stepsTEOAE+TEOAEScreening personnel10131414169753 stepsTEOAE+TEOAE+ABRAudiologist or14141418402 stageTEOAE+ABRAudiologist or15141415753 stageTEOAE+ABRAudiologist or1453 vears115753 stageTEOAE+ABRAudiologist or1510 vearsNewborns115753 stageTEOAE+ABRAudiologist or16510 vearsNewborns19436 (NP)3 stageTEOAE+ABRAudiologist or1510 vearsNewborns19436 (NP)3 stageTEOAE+ABRAudiologist or1655 vearsNewborns136333 stageTEOAE+TEOAE+ABRAudiology students1655 vearsNewborns136333 stageTEOAE+TEOAE+ABRAudiology students1655 vearsNewborns136333 stageTEOAE+TEOAE+ABRAudiology students1655 vearsNewborns136333 stageTEOAE+TEOAE+ABRAudiology students1655 vearsNewborns136332 stageTEOAE+TEOAE+ABRAudiology students1655 vear | Shameem <i>et al</i> 2022 | 42 | Bangladesh (LMIC) | 2 years | High-risk newborns | 426 | 2 stage | TEOAE+TEOAE | Not mentioned | Diagnostic ABR |
| Image: section constraints Duration constraints Constraints Screening sections Screening personnel Image: section constraints Buration constraints Reveloped Screening personnel Screening personnel Screening personnel Screening personnel Image: section constraints Screening personnel Screening personnel Screening personnel Screening personnel Screening personnel Image: section constraints Image: section constraints Image: section constraints Screening personnel Screening personnel Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Image: section constraints Im | Khaimook <i>et al</i> 2022 | 43 | Thailand (UMIC) | 6 months | Newborns | 1696 | 2 stage | TEOAE+TEOAE | Trained nurse and audiologist | Diagnostic ABR+Tymp |
| Image: bit is the state is the sta | (D) | | | | | | | | | |
| 5614 monthsNewborns169753 stepsTEOAE+TEOAE+ABRAudiology technician141 yearNewborns18402 stageTEOAEAudiologist or14563 yearsNewborns115753 stageTEOAE+AABR2 audiometrists and 11553 yearsNewborns115753 stageTEOAE+AABR2 audiometrists and 1510 yearsNewborns19436 (//P)3 stageTEOAE+TEOAE+ABR2 audiometrists and 155 yearsNewborns19436 (//P)3 stageTEOAE+TEOAE+ABRAudiology students55 yearsNewborns136933 stageTEOAE+TEOAE+ABRAudiology students66 yearsNewborns136933 stageTEOAE+TEOAE+ABRAudiology students | Author and year | Citation | Duration of programme | Population screened | Screened (n) | Screening protocol | Screening | test used | Screening personnel | Diagnostic test |
| 1441 yearNewborns18402 stageTEOAEAudiologist or audiometrist4563 yearsNewborns115753 stageTEOAE+AABR2 audiometrist5710 yearsNewborns19436 (/P)3 stageTEOAE+TABR2 audiometrist and 1585 yearsNewborns136933 stageTEOAE+TEOAE+(TEOAE and AABR)Audiology students585 yearsNewborns136933 stageTEOAE+TEOAE+ABRAudiology students586 yearsNewborns136933 stageTEOAE+TEOAE+ABRAudiology students | Tasci <i>et al</i> 2010 | 55 | 14 months | Newborns | 16975 | 3 steps | TEOAE+TE | EOAE+ ABR | Audiology technician | Diagnostic ABR |
| 4 ⁵⁰ 3 vaars Newborns 11575 3 stage TEOAE+ABR 2 audiometrists and 1 nurse ⁵⁷ 10 vears Newborns 19436 (/P) 3 stage TEOAE+TEOAE+(TEOAE and ABR) 2 audiometrists and 1 nurse ⁵⁸ 10 vears Newborns 19436 (/P) 3 stage TEOAE+TEOAE+(TEOAE and ABR) and audiology students ⁵⁸ 5 years Newborns 13693 3 stage TEOAE+TEOAE+ABR Audiology students ⁴⁶ 6 years Newborns 142 128 2 stage TEOAE+TEOAE+ABR Not mentioned | Sennaroglu <i>et al</i> 2011 | 4 | 1 year | Newborns | 1840 | 2 stage | TEOAE | | Audiologist or audiometrist | Diagnostic ABR; |
| 5710 yearsNewborns19436 (I/P)3 stageTEOAE+TEOAE+(TEOAE and ABB)Audiology technicians585 yearsNewborns136933 stageTEOAE+TEOAE+ABBAudiology students585 yearsNewborns136933 stageTEOAE+TEOAE+ABBAudionetrist466 yearsNewborns142 1282 stageTEOAE(twice same day)+TEOAENot mentioned | Ulusoy <i>et al</i> 2014 | 56 | 3 years | Newborns | 11575 | 3 stage | TEOAE+AA | ABR | ometrists and | Diagnostic ABR, the level 3 centre |
| ⁵⁸ 5 years Newborns 13693 3 stage TEOAE+TEOAE+AABR Audiometrist ⁴⁵ 6 years Newborns 142 128 2 stage TEOAE (twice same day)+TEOAE Not mentioned | loğlu et | 57 | 10 years | Newborns | 19436 (I/P) 2083 (O/P) | 3 stage | TEOAE+TE | COAE+(TEOAE and AABR) | Audiology technicians and audiology students | Diagnostic ABR |
| ⁴⁵ 6 years Newborns 142 128 2 stage TEOAE (twice same day)+TEOAE Not mentioned | Yorulmaz <i>et al</i> 2017 | 28 | 5 years | Newborns | 13693 | 3 stage | TEOAE+TE | :OAE+AABR | Audiometrist | Diagnostic ABR, Tymp, acoustic reflexes, ASSR |
| | Çelik <i>et al</i> 2016 | 45 | 6 years | Newborns | 142 128 | 2 stage | TEOAE (tw | ice same day)+TEOAE | Not mentioned | Diagnostic ABR |

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| (D)(D)Author and yearCitationDAuthor and yearCitation \mathbf{P} Ozturk et al 2017 5^{3} 2 Yücel et al 2018 6^{0} 2 Yücel et al 2013 4^{6} 8 Arslan et al 2013 4^{6} 8 Çıkrıkçı et al 2020 5^{1} 1.1 Citation \mathbf{p} \mathbf{P} Author and yearCitation \mathbf{p} Arjmandi et al 2013 4^{8} 1 Islami et al 2013 4^{8} 1 Firoozbakht et al 2013 4^{8} 1 Zahed et al 2013 2^{8} 8 2014 2^{12} 2^{8} 8 | | | | | | | |
|--|--|--|--|--|--|---|---|
| or and year Citation ik et al 2017 59 ii 2018 60 et al 2019 65 et al 2013 46 r) et al 2013 46 cpi et al 2020 51 cpi et al 2013 46 or and year Citation or and year Citation or and year 2012 i et al 2013 48 bi et al 2013 48 conditiet al 2013 48 conditiet al 2013 48 conditiet al 2013 48 | | | | | | | |
| k et al 2017 ⁵⁹ di 2018 ⁶⁰ et al 2019 ⁶⁵ cpi et al 2013 ⁴⁶ cpi et al 2020 ⁵¹ and year Citation andi et al 2012 ⁴⁷ ti et al 2013 ⁴⁸ bi et al 2013 ⁴⁸ cbakht et al ²⁹ | Duration of programme | Population screened | Screened (n) | Screening protocol | Screening test used | Screening personnel | Diagnostic test |
| li 2018 ⁶⁰ 1: et al 2019 ⁶⁵ 1: n et al 2013 ⁴⁶ 1: c; et al 2020 ⁵¹ c; et al 2020 ⁵¹ ntridi et al 2012 ⁴⁷ i et al 2013 ⁴⁸ zbakht et al ²⁹ | 2 years | Newborns | 7502 | 3 stage | Wellbabies: DPOAE+DPOAE+ABR screening Highrisk babies: Direct ABR | R Audiologist | Diagnostic ABR |
| et al 2019 ⁶⁵ n et al 2013 ⁴⁶ cpi et al 2020 ⁵¹ cpi et al 2020 ⁵¹ and year Citation and et al 2012 ⁴⁷ i et al 2013 ⁴⁸ 2bakht et al ²⁹ ch et al 2014 ³⁰ | 2 years | Newborns | 1808 | 3 stage | TEOAE+TEOAE+ABR screening | Nurses (trained) | Diagnostic ABR |
| r et al 2013 ⁴⁶ cp et al 2020 ⁵¹ 51 51 51 51 51 51 51 51 | 2 years | Newborns | 7230 turkish | 3 stage | (TEOAE and Tymp)+TEOAE+ABR | Not mentioned | Detailed testing |
| cçi et al 2020 ⁵¹ or and year Citation andi et al 2012 ⁴⁷ i et al 2013 ⁴⁸ zbakht et al ²⁹ | 8 months | Newborns | 2229 | 2 stage | TEOAE+TEOAE | Nurse | Diagnostic ABR |
| or and year Citation and et al 2012 47 i et al 2013 48 zbakht et al 29 chet al 2014 30 | 1.5 years | Newborns | 702 turkish 172 syrian | 2 stage | AABR+AABR | Not mentioned | Diagnostic ABR |
| or and year Citation andi <i>et al</i> 2012 ⁴⁷ i <i>et al</i> 2013 ⁴⁸ zbakht <i>et al</i> ²⁹ | | | | | | | |
| andi <i>et a</i> / 2012 ⁴⁷ i <i>et a</i> / 2013 ⁴⁸ zbakht <i>et a</i> / ²⁹ ch <i>et a</i> / 2014 ³⁰ | Duration of programme | Population screened | Screened (n) | Screening protocol | Screening test used | Screening personnel | Diagnostic test |
| i <i>et a</i> l 2013 ⁴⁸ zbakht <i>et a</i> l ²⁹ d <i>et a</i> l 2014 ³⁰ | 1 year | Newborns | 1232 | 2 stage | TEOAE+TEOAE | Not mentioned | Diagnostic ABR |
| zbakht et <i>al</i> ²⁹ d et al 2014 ³⁰ | 1.5 years | Newborns | 7250 | 2 stage | TEOAE+TEOAE | Audiologists | Diagnostic ABR |
| 30 | 8 years | Newborns | 3 350 995 | 2 stage | TEOAE+AABR | Audiologists, nurses, midwives and trained health technicians. | Comprehensive test |
| | 8 years | Newborns | 40 930 | 2 stage | TEOAE+ABR | Audiologists | ABR/ASSR and immittance audiometry, |
| Farhat <i>et al</i> 2014 ⁹⁰ 2 | 2 years | Newborns | 8987 | 2 stage | TEOAE+TEOAE | Not mentioned | ASSR |
| Haghshenas <i>et al</i> ⁶¹ 2014 | 2 years | Newborns | 15165 | 3 stage | OAE+OAE+(OAE and AABR) (not mentioned DP/TE) | Audiologist | ABR screening |
| Baradaranfar <i>et al</i> ¹⁰⁸ 1) 2014 | 1 year | Newborns | 514 | 2 stage | TEOAE+TEOAE | Not mentioned | Diagnostic ABR |
| Azizi et al 2016 ⁴⁹ 1. | 1.5 years | Newborns | 3818 | 2 stage | TEOAE+TEOAE | not mentioned | ABR, |
| Tajik <i>et al</i> 2016 ³¹ 4 | 4 years | Newborns | 3362 | 2 stage | TEOAE+(TEOAE and ABR) | Not mentioned | Not mentioned |
| Saki et a/ 2017 ⁸⁴ 3 | 3 years | Newborns | 92 52 1 | 2 stage | First and second:TEOAE+AABR | Audiologists | Diagnostic OAE and ABR |
| Rahimi <i>et al</i> 2018 ⁶² 5 | 5 years | Newborns | 4729 | 3 stage | TEOAE+TEOAE+AABR | Audiologist | Diagnostic ABR |
| AABR, automated auditory brainstem response; ABR, auditory brainstem response; ASSR otoacoustic emission; LMIC, lower middle-income country; MIC, middle-income country; levoked otoacoustic emission; Tymp, tympanometry; UMIC, upper middle-income country, | em response; AF · middle-income p, tympanometr | BR, auditory brair country; MIC, mi y; UMIC, upper n | astem response; A ddle-income cour niddle-income cou | SSR, auditory stea htry; NHS, newborr intry. | AABR, automated auditory brainstem response; ABR, auditory brainstem response; ASSR, auditory steady-state response; BOA, behavioral observation audiometry; DPOAE, distortion product otoacoustic emission; LMIC, lower middle-income country; MIC, middle-income country; NHS, newborn hearing screening; OAE, otoacoustic emission; PTA, pure tone audiometry; TEOAE, transient evoked otoacoustic emission; Tymp, tympanometry; UMIC, upper middle-income country. | servation audiometry; DPOAE, dist imission; PTA, pure tone audiometr | tortion product y; TEOAE, transient |

| **1012 ματ5 ματ1111042 ματ2 ματ <th>Author and year</th> <th>Citation</th> <th>Country</th> <th>Duration of programme</th> <th>Age of screening (years)</th> <th>Screened (n)</th> <th>Screening protocol</th> <th>Screening test used</th> <th>Pass/fail criteria</th> <th>Screening personnel</th> <th>Diagnostic test</th> <th>Diagnostic person</th> | Author and year | Citation | Country | Duration of programme | Age of screening (years) | Screened (n) | Screening protocol | Screening test used | Pass/fail criteria | Screening personnel | Diagnostic test | Diagnostic person |
|--|-------------------------------------|----------|----------------------|--------------------------|-----------------------------|-----------------|-----------------------|--|--|---|---|------------------------------|
| 0 1da 3 yara 6 - 7 yara 1 data 0 denotions Denotion <thdenotions< th=""></thdenotions<> | Tuli e <i>t al 2</i> 012 | 99 | India | 2 years | 5-16 years | 111 | 1 stage | Case history, audiological and ENT evaluation, awareness and SIFTER | Not mentioned | Not mentioned | ENT and PTA and diagnostic ABR | Audiologist |
| 1 1da 2 yaas Bith-5 yaas 2 stages D Oct-D Oct SN3.36 Toped vilge hash Biellopentic ABI- stages 1 1 usis 2 wars Bith-5 yaas 2 stages D Oct-D Oct SN3.46 Toped vilge hash D option diation 1 1 usis 6 words Bith-5 yaas 2 stages D volce | Chadha <i>et al</i> 2013 | 67 | India | 3 years | 5-12 years | 15718 | 1 stage | Otoscopy, 10-Question Screening Index for Disabilities in English and Hindi | Positive history of hearing or speech defects, a positive finding on examination | Proforma: parents, otoscopy: otolaryngologists | Not mentioned | Not mentioned |
| 0 1dia 2 years 2 Harbo | Ramkumar <i>et al</i> 2018 | | India | 2 years | Birth–5 years | 1335 | 2 stages | DPOAE+DPOAE | >SNR 3dB | Trained village health worker | Telediagnostic ABR | Audiologist |
| 0 10dia 6 monts 617 years 537 1 stage I vanito de test Normetioned P vanitonid P vanitonid 0 10dia Nici 6 -14 years 674 1 stage P vanitonid EV secondation EV seco | Ramkumar <i>et al</i> 2019 | | India | 2 years | Birth-5 years | 2815 | 2 stages | DPOAE+DPOAE | >SNR 3dB | Trained village health Worker | Diagnostic ABR— in person and telediagnostic ABR | Audiologist |
| 0UndaKur benitomedEvityeeVariationEvityee <td>Jerma et al 2025</td> <td></td> <td>India</td> <td>6 months</td> <td>6-17 years</td> <td>597</td> <td>1 stage</td> <td>Tuning fork test</td> <td>Not mentioned</td> <td>Not mentioned</td> <td>PTA and Tymp</td> <td>Audiologist</td> | Jerma et al 2025 | | India | 6 months | 6-17 years | 597 | 1 stage | Tuning fork test | Not mentioned | Not mentioned | PTA and Tymp | Audiologist |
| °° China 1 service 3-6 years 1 stage FA 1 stage FA Service service service PA (5-6 years) PA | Shekhar <i>et al</i> 2022 | 68 | India | Not mentioned | 5-14 years | 474 | 1 stage | PTA | Not mentioned | ENT specialist | ENT examination | ENT specialist |
| ⁶⁰ China Tyters 364 years 2654 bits 1 stage TeXes Sthod nurses and solutions Contonic straining Contonic strainin Contonic strainin Contonic s | Lü <i>et al</i> 2011 | 69 | China | 1 year | 3–6 years | 21427 | 1 stage | PTA | 1, 2 and 4 kHz >20dB | Screening person with training (training programme with certificate) | PTA (5–6 years) VRA or play PTA (3–4 years) | Not mentioned |
| "" Unitary Lations Software-based rew and Matz Software-based rew and Matz Software-based rew and Matz Internitions Internitoned Internitions Inten | Chen <i>et al</i> 2013 | | China | 1 year 5 months | 3–6 years | 28546 | 1 stage | TEOAE | >SNR 3dB | School nurses and doctors 2 hours of training | Comprehensive test | Not mentioned Audiologist |
| 1 ¹ China Not SodBHLat 1, 2 Atomatic test: nurses Tymp. PPOAE and with 25-8.0.kHz 2 Turkey 3-5 years 3-5 years 239 1 stage PTA Atomatic test: nurses Tymp. PPOAE and with 25-8.0.kHz 2 Turkey 3 years 3-5 years 239 1 stage PTA Not mentioned Automatic test: nurses Tymp. PPOAE and with 25-8.0.kHz 3 Turkey 1 year 3-5 years 239 1 stage PTA Not mentioned Automatic test: nurses For examination 3 Turkey 1 year 69-84 months 23664 2 stage PTA Not mentioned Automatic test: nurses For examination 3 Turkey 1 year 69-84 months 23664 2 stage PTA Not mentioned PTA complexity PTA complexity PTA complexity PTA | Wu et al 2014 | 02 | China | Not mentioned | 3-6 years | 6288 | 1 stage | Software-based new PTA | >30dBHL at 1, 2 and 4 kHz | Preschool teachers – minimally trained | Not mentioned | Not mentioned |
| 7 Turkey 3-5 years 3-5 years 1 stage PTA Not mentioned Motionolst and SLP EN rexamination 7 Turkey 1 year 69-34 months 2364 2 stage PTA, 500, 1000, 2000 Certified nurses, EN rexamination 7 Turkey 1 year 69-34 months 2364 2 stage PTA, 500, 1000, 2000 Certified nurses, EN rexamination 7 Turkey 1 year 69-34 months 2364 2 stage PTA, 500, 1000, 2000 Certified nurses, EN rexamination 7 Turkey 1 year 6-7 years 237 Not PTA Not | Kam <i>et al</i> 2014 | | China | Not mentioned | 3–7 years | 6231 | 1 stage | Automated PTA | >30 dBHL at 1, 2 and 4 kHz | Automatic test: nurses with 2 hours' training as facilitator | | Not mentioned |
| ¹⁰ Turkey I year 69–84 months 23664 P and a conditionation of a conditionationation of a conditionationationationationationationation | Tokgöz-Yılmaz ə <i>t al</i> 2013 | 22 | Turkey | 3 years | 3–5 years | 239 | 1 stage | PTA | Not mentioned | Audiologist and SLP | ENT examination | ENT specialist |
| ¹² Iran I year 6-7 years 237 Not mentioned PTA, Not mentioned PTA, Weber, Rinnetes mentioned PTA, PTA | Caplama <i>et al</i> 2020 | 62 | Turkey | 1 year | 69-84 months | 23664 | 2 stage | PTA, 10 questionnaire | 500, 1000, 2000 and 4000 Hz >20 dB 10 questions – refer in 1 question | | ENT examination | ENT specialist |
| ⁷² Iran 4 months 6-13 years 2019 1 stage PTA 0.5-4.0 kHz >15 Not mentioned BNL and comprehensive audiological and comprehensive audiological examination tube for the formation mentioned mentioned for the formation mentioned mentioned for the formation mentioned | larvijEslami <i>et a</i> 2017 | | Iran | 1 year | 6-7 years | 2237 | Not mentioned | PTA | Not mentioned | Not mentioned | PTA, Weber, Rinne test | Not mentioned |
| ⁶ Tajikistan Not 6–8 years 143 1 stage Questionnaire, PTA module (500- Medical doctors Detailed PTA (LMIC) mentioned PTA using SZOK 8 kHz) >25 dB at Other specialists telemed model one frequency | Jalali <i>et al</i> 2020 | | Iran | 4 months | 6-13 years | 2019 | 1 stage | PTA | 0.5–4.0kHz >15 dBHL | Not mentioned | ENT examination and comprehensive audiological examination | Not mentioned |
| | Skarzyński <i>et al</i> 2016 | | Tajikistan (LMIC) | Not mentioned | 6-8 years | 143 | 1 stage | Questionnaire, PTA using SZOK telemed model | PTA module (500– 8kHz) >25dB at one frequency | Medical doctors Other specialists | Detailed PTA | Audiologists |

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| yearCitationCountryprogramme(years)(n)Alaqrabawi et al74Jordan4 years5-15 years16492016(UMIC)(UMIC)4 years5-15 years1649Al-Obeidy et al75Iraq (UMIC)1 year6 years4252019201975Iraq (UMIC)1 year1 year1 year | Duration of Age of screening Screened | Screened Screening | | | | | Diagnostic |
|---|---------------------------------------|--------------------|---------------------|---|--|--|---------------|
| Jordan 4 years 5–15 years (UMIC) I year 6 years Iraq (UMIC) 1 year | | protocol | Screening test used | | Pass/fail criteria Screening personnel Diagnostic test | Diagnostic test | person |
| 75 Iraq (UMIC) 1 year 6 years | | 1 stage | PTA | 500 Hz, 1, 2 and 4 Not mentioned kHz >25 dB | Not mentioned | Audiometry, otoscopy and Tymp | Audiologists |
| | | 1 stage | HR questionnaire | Not mentioned | Not mentioned | ENT examination, TFT (Weber, Rinne and absolute bone conductio). HRR children: PTA | Not mentioned |

the initial stage of screening either employed AABR solely for both stages⁵⁰ or a combination of AABR and OAE to screen only high-risk newborns.^{20 52} Four studies from China used two-stage screening coupled with genetic hearing screening.^{21–23 25 53}

When a three-stage protocol was used, generally the first two stages included OAE (DP/TE) screening followed by AABR/auditory brainstem response (ABR) screening⁵⁴⁻⁶² or included OAE (DP/TE) for all three stages.^{63 64} Only one study reported combining tympanometry and TEOAE in the initial stage of its three-stage screening protocol.⁶⁵ Studies from Turkey (n=7) reported a three-stage screening protocol.

Screening for older children

Fourteen studies for older children employed a singlestage screening protocol⁸ ^{66–75} with three employing a two-stage protocol.³ ⁷⁶ Ten studies reported using subjective hearing screening tests, two studies used questionnaire or otoscopy for screening⁶⁷ ⁷⁵ and another three studies used TEOAE.⁷⁶ ⁷⁷ Pure tone audiometry (PTA) was the most commonly used subjective test for screening older children.⁶⁸ ⁶⁹ ^{72–74} ⁷⁸ Two studies reported the use of automated software-based PTA.⁷⁰ ⁷¹ PTA was combined with questionnaires⁸ ⁷⁹ or otoscopy.⁶⁷ ⁷⁵ Only one study reported the use of TEOAE screening.⁸⁰

Pass/refer criteria

In several programmes for newborn and infant screening, screening results were based on data generated from the screening instrument automatically. The pass criteria for DP/TEOAE was between 3 dB and 6 dB signal-to-noise ratio, ¹⁹ 20 ²⁵ 37 ³⁸ 40 ⁴³ 45 ⁴⁹ 54 57 63 64 ⁸¹ ⁸² and for AABR, it varied between 30, 35 and 40 dB neural hearing loss (NHL).^{20 52 56 58 61} Predominantly, refer results in one ear was considered for follow-up screening.

For screening older children, the pass criteria for PTA ranged from 15 dB HL to 30 dB HL. All studies used the four frequencies from 0.5 kHz to 4.0 kHz for pure tone testing. In questionnaire-based studies, failing one item or a family history of hearing loss was the referral criterion.^{67 68}

Screening personnel

Audiologistswere the primary screening personnel in many newborn and infant programmes, ^{17 30 34 38 44 48 54 59 61 62 83 84 followed by nurses.^{20 21 26 28 29 37 40 43 46 54 56 60 64} In five studies, the training provided for nurses to perform hearing screening was also briefly mentioned, ^{28 29 40 46 60} including some certifications.⁵⁶ Other than nurses, some studies reported audiometrists^{44 56 58} and audiologist technicians⁵⁵ as personnel involved in screening. Other nonspecialists that were engaged in hearing screening were technicians,⁶⁴ ward attendants,⁶⁴ trained health workers^{29 85} social workers⁸³ and midwives.^{29 33 81} In a few programmes, otolaryngologists⁶³ performed the hearing screening. Out of 59 studies, 29 did not provide any information regarding the screening individual.} Screening for older children was conducted by otorhinolaryngologists^{8 67 68} audiologists⁷² and audiometrists.⁷⁹ Other non-specialists involved in the hearing screening included trained nurses/midwives,^{38 71 79} trained village health workers or volunteers,^{76 77} and school teachers with training.⁷⁰

Studies have reported a variety of training programmes. They included hearing screening certification,^{67,79} 2 hours of TEOAE training,³⁸ TEOAE training and telediagnostic testing facilitation,⁷⁶ and minimal training/2 hours of training for facilitating automated PTA.^{70,71}

Confirmation of hearing loss

Diagnostic ABR was the only testing carried out to confirm the hearing loss in studies in newborns and infants.²⁵ ²⁸ ³² ^{35–37} ⁴² ⁵⁴ ⁶³ ^{86–89} Comprehensive test battery including the diagnostic BERA, OAE, and tympanometry was mentioned only in 11 studies.²⁰ ²⁹ ⁵⁸ Four studies also reported the inclusion of the auditory steady-state response (ASSR) in the test battery.³⁰ ⁵⁸

Two programmes used solely ASSR,^{39 90} and studies also used ABR screening at 30²⁰ or 35 dB NHL⁶¹ for hearing loss diagnosis.

However, 11 of the 65 programmes made no mention of the diagnostic confirmatory test used for confirmation of hearing loss. More than half of the studies (n=37), reported that the diagnostic confirmatory test was performed at the same hospital where screening was conducted. In another 18 studies, children were referred to more specialist or tertiary care facilities for diagnostic confirmatory tests. The diagnostic site was not mentioned or could not be inferred in 10 studies.

In studies reporting screening for older children, a test battery approach was used in three studies where they included PTA with tympanometry and DPOAE⁷¹ or PTA with otoscopy and tympanometry⁷⁴ or PTA and detailed ABR.⁶⁶ Two studies reported the use of comprehensive test battery but did not mention the tests included.³⁸

PTA was frequently included in the diagnostic test battery,^{71 74 91} but in three studies, PTA was the only diagnostic test used.^{8 73 78} Of the studies that reported the use of PTA for diagnosis, only four studies^{72–74 78} mentioned information related to bone conduction testing. Apart from these studies, ENT examination was included in five studies.^{68 72 73 75 79} The diagnostic testing sites included a hospital,⁷³ a school,⁶⁸ a speech and hearing centre,⁷¹ and a telemedicine platform.^{8 76}

Use of ICT

In studies related to newborn and infant hearing screening, three programmes reported the use of ICT for storing and forwarding results,³⁴ database management^{28 83} and sending reminders for follow-up screening.

In studies reporting screening of older children, five studies reported using telepractice for screening, diagnosis or both. Telediagnostic ABR^{76 77} was reported in India. Use of m-health-based automated hearing screening was reported in China.^{70 71} .A telesensory

screening platform including hearing screening was reported (SZOK - (Sense Examination Platform) paradigm) in Tajikistan, where both screening and diagnosis were carried out via telemedicine.⁸

Validity and efficacy of the screening programmes

Validity of screening programmes as reported in the studies was evaluated based on three criteria: use of a validated screening tool, use of a validated diagnostic tool, and whether the programme was in the design phase or in the implementation phase.

Among the studies that reported newborn and infant hearing screening, 48 studies fulfilled all three criteria of the validity tool; 11 studies fulfilled two out of three criteria; and 6 studies fulfilled one out of three criteria (figure 1A). The validated screening tool was used by 63 studies and 54 studies used a validated diagnostic tool. As per the criteria we used, 55 studies could be classified to be in the implementation phase and 10 studies were in the design phase.

Economic analysis, frequency of identification and intervention were the three criteria included to assess efficacy. Only 2 studies fulfilled all the three efficacy criteria; 17 studies fulfilled two out of the three criteria; and 37 studies fulfilled only one of the three criteria, whereas the remaining 9 studies did not fulfil any of the criteria. Fifty-one studies reported only the frequency of identification, whereas 14 reported both the frequency of identification and intervention. Twelve per cent of the studies did not mention either of these outcomes. Economic analysis was very limited (n=3) and was reported majorly in public programmes.

Among the studies that reported screening programmes for older children, 10 studies fulfilled all the three criteria; 3 studies fulfilled two out of three criteria; and 3 studies fulfilled one out of three criteria. Only one study did not meet any of the criteria⁶⁷ since only a questionnaire and an otoscopic examination were used to estimate the incidence of conductive hearing loss in older children.

With respect to efficacy, it was observed that none of the studies among older children fulfilled all the three criteria. Only five studies fulfilled two out of three criteria, whereas the remaining 12 studies fulfilled only one criterion.

Fourteen studies have reported frequency of identification, but only five studies have reported the frequency of intervention (eg, medical intervention for conductive pathology). The intervention-related screening programmes were reported from India, China and Turkey. The economic analysis was reported in only two studies.^{71 77} Except for the economic analysis, only 2 of the 17 studies fulfilled all validity and efficacy criteria.^{69 76}

Prevalence of hearing loss

Across 48 studies, the mean prevalence of hearing loss among newborns and infants was 5/1000 in India, 2/1000 in China, 2/1000 in other Southeast Asian nations (Thailand, Malaysia and Nepal), 2/1000 in Turkey, and 4/1000 in Iran. Figure 2A–E shows the forest plots for prevalence of each country.

In screening programmes for older children, 11 studies reported number of cases with hearing loss including conductive and sensori neural hearing losses. However, in four studies,^{67 68 79 80} the specific audiological tests conducted to diagnose were not mentioned, and in seven studies,^{69 72–75 78} details of diagnostic audiometry were provided. In this age group, the percentage of conductive hearing loss reported was higher compared with sensori neural hearing loss across all the studies. In two studies, the type of loss was not differentiated.^{8 67} The percentage of children identified with a certain type of hearing loss was calculated based on the information on the number of children diagnosed that was provided in each of the studies. The study outcomes are reported in table 3.

Barriers and facilitators

Barriers

Loss to follow-up for second screening and diagnostics^{20 29 35–37 40 43 48 54 56 59 81 87} was reported as a major challenge. Loss to follow-up was linked to parental rejection for diagnosis,^{33 43 50} poor tracking system,^{20 29} financial burden of parents, low socioeconomic status⁵¹ and travel distance to testing distance. Other major challenges highlighted in relation to outcomes included limited coverage^{35 82} and a high referral rate,^{18 37 54} poor longterm outcomes with respect to coverage and referral rate.²⁴

Other factors that had an indirect impact on programme outcomes included the lack of dedicated screening personnel,⁵⁰ lack of professional resources/audiologists,^{29 84} high ambient noise in the testing environment⁸² and the absence of diagnostic facilities.⁵⁶ A few studies mentioned challenges affecting programme implementation, such as the use of a three-step protocol only with OAE,⁵⁵ the difficulties of centralised programme implementation in remote locations²⁹ and delay in diagnosis in remote locations due to referral to regional facilities.⁸⁴

In screening for older children, children's attention was regarded as a major challenge resulting in poor accuracy.⁷¹ Other key factors influencing programme outcomes included inadequate internet connectivity^{8 76} and poor follow-up due to social stigma.

Facilitators

Use of appropriate tracking or data management systems, were reported to be helpful in minimising loss to follow-up.²⁰ ²⁸ ³³ ³⁵ ⁸¹ Combining hearing screening with other screenings improved follow-up rates.²⁵ ⁶² Several studies highlighted strategies to minimise false referral rates, including (1) employing a conducive environment and trained individuals,⁵⁴ (2) adding AABR in the initial stage of screening protocol,⁵² (3) screening between 3 days and 5 days of age⁶² and (4) incorporating tympanometry into the screening protocol.⁸⁹ Financial assistance in the form of funding²⁸ ³⁷ ⁸³ and centralised hearing screening facilities or grouping more centres³³ ⁸¹ were

| Table 3 | Table 3 Secondary outcomes: studies reporting number of cases identified with CDHL/SNHL in older children in each country | idies reporting r | number of cases iden | tified with CDI | HL/SNHL in older c | children in ea | ch country | | |
|------------|---|-----------------------|----------------------------|--------------------|--------------------------|------------------|---------------------------|------------------------------|---|
| Country | Author and year | Screened (n) | CDHL identified (n) | % of CDHL | NHL identified (n) | % of SNHL | Overall HL identified (n) | % of HL | 95% CI (LB to UP) |
| India | Chadha <i>et al</i> . ⁶⁷ 2013 | 15718 | NA | NA | NA | NA | 1578 | 10.30 | 9.57 to 10.52 |
| | Shekhar <i>et al</i> . ⁶⁸ 2020 | 474 | 146 | 30.80 | - | 0.21 | 147 | 31.01 | 26.87 to 35.39 |
| Turkey | Tokgöz-Yılmaz <i>et al.⁷² 2</i> 013 | 239 | 25 | 10.46 | - | 0.42 | 26 | 10.88 | 7.23 to 15.53 |
| | Kaplama <i>et al.</i> ⁷⁹ 2020 | 23664 | 186 | 0.79 | 89 | 0.37 | 275 | 1.16 | 1.03 to 1.31 |
| Iran | TarvijEslami <i>et al.</i> ⁷⁸ 2017 | 2284 | 28 | 1.23 | 8 | 0.35 | 36 | 1.58 | 1.11 to 2.18 |
| | Jalali <i>et al.</i> ⁷³ 2020 | 2019 | 19 | 0.94 | 8 | 0.39 | 27 | 1.33 | 0.88 to 1.94 |
| Tajikistan | Skarzyński et al. 2016 | 143 | NA | NA | NA | NA | 34 | 23.70 | 17.06 to 31.61 |
| Jordan | Alaqrabawi <i>et al.</i> ⁷⁴ 2016 | 1649 | 54 | 3.27 | 36 | 2.18 | 06 | 5.45 | 4.41 to 6.61 |
| Iraq | Al-Obeidy <i>et al.</i> ⁷⁵ 2019 | 425 | 28 | 6.59 | 2 | 0.47 | 30 | 7.06 | 4.81 to 9.92 |
| China | Lu <i>et al.</i> ⁴⁴ 2011 | 21547 | 285 | 1.32 | 16 | 0.07 | 301 | 1.39 | 1.24 to 1.56 |
| | Chen <i>et al.</i> ³⁸ 2012 | 28546 | 344 | 1.21 | 22 | 0.08 | 366 | 1.29 | 1.15 to 1.42 |
| China | Al-Obeidy <i>et al.</i> ^{1,9} 2019 Lu <i>et al.</i> ⁴⁴ 2011 Chen <i>et al.</i> ³⁸ 2012 | 425 21547 28546 | 28 285 344 | | 6.59 1.32 1.21 | | 2 16 22 | 2 0.47 16 0.07 22 0.08 | 2 0.47 30 16 0.07 301 22 0.08 366 |
| CDHL, con | CDHL, conductive hearing loss; HL, hearing loss; LB, lower bound; NA, not mentioned; NHL, neural hearing loss; SNHL, sensorineural hearing loss; UB, upper bound. | oss; LB, lower bound | d; NA, not mentioned; NHL, | neural hearing los | s; SNHL, sensorineural h | earing loss; UB, | upper bound. | | |

strategies reported in studies to improve coverage rates. Multicentre-based or a centralised hearing screening programme was reported to be resource efficient with respect to cost, infrastructure and professionals.⁸¹

DISCUSSION

The primary purpose of this review was to describe the models of hearing screening programmes implemented in young children in various Asian L&MICs in the published scientific literature. The inclusion of countries was based on the World bank classification rather than culturally defined regions; this led to a heterogenous inclusion with central Asian and middle eastern countries as well. Out of 61 L&MICs in Asia, only 14 countries reported hearing screening programmes that fit our inclusion criteria. In a recent systematic review, highquality literature with hearing screening programmes was reported to be primarily in HICs⁹²; yet, it is also likely that resources for research and publication are low and hence are also low on priority in the L&MICs context. Though studies from both L&MICs were included, our results show that most of the studies reporting on hearing screening were from the middle-income countries and more specifically from UMICs. This suggests greater adoption of EHDI measures in UMICs, possibly due to greater availability of resources in comparison to LMICs and LICs.

Our review gathered evidence on hearing screening programmes in general, including screening protocols, screening tests, pass/fail criteria, screening personnel, diagnostic tests, use of ICT, and programme validity and efficacy. The hearing screening tools and protocols used for newborns, infants and older children were similar to those used in HICs.⁹³ Despite the fact that the majority of programmes used a two-stage OAE (DP/TE) and ABR screening as preferred screening tools across countries, there was no consistency in protocol stages or screening tests undertaken. This was consistent with Kanji *et al* 's assessment of NHS protocols, which revealed non-uniformity in the protocols followed.

It was also noted that objective hearing screening was most commonly reported over subjective hearing screening for newborns and infants. Only one study⁸⁵ found good sensitivity and specificity for behavioural hearing assessment for neonates and infants using calibrated noise makers. The use of objective screening in L&MICs implies a preference for international best practices based on Western contexts and guidelines.² However, it is important to assess the sustainability and long-term outcomes of these efforts. Subjective singlestage PTA screening, on the other hand, was extensively used in various screening programmes for older children above the age of 3. This is comparable to HICs where PTA screening is mandatory for children over the age of 3.94 95 In contrast, the current review found a few public initiatives^{75 87 96} that used questionnaire methods, and this

implies that mass screening was being done by low-cost tools like questionnaires where resources were limited.

Audiologists were the most common screening personnel in newborn screening programmes across Asian L&MICs. This is in contrast to HICs, where nurses mostly performed hearing screening.⁹⁷ While the majority of NHS programmes in Asian L&MICs were started by audiologists or otolaryngologists in private hospitals, in most HICs, the screening programmes were generally universal and followed as a part of other normal newborns screening before discharge. Screening of older children was mostly done by otolaryngologists, school instructors and nurses. This could be because many of the screening programmes for older children were conducted in schools or community settings in the absence of audiologists on site. In contrast, hearing screenings are carried out at child health clinics by a dedicated school nurse/ audiologist in HICs.⁹⁷

Use of the test battery was limited in diagnostic confirmation of hearing loss. Detailed ABR testing was considered as the standard diagnostic tool in many countries as it examines the entire peripheral auditory pathway responsible for hearing. Apart from this, studies from China employed a test battery containing a variety of tests altogether (eg, ASSR, ABR and tympanometry) to confirm hearing loss. In WHO guidelines for hearing screening, diagnostic test battery including ABR/ASSR, tympanometry, acoustic reflex, otoscopic examination and medical evaluation was suggested.⁹⁸ Therefore, in HICs, the diagnostic test battery approach is mostly preferred.⁹⁷ In screening programmes for older children, medical (ENT) examination in cases of conductive pathology and routine PTA with or without tympanometry were prioritised as tests to confirm hearing loss. This is inconsistent with the WHO guidelines⁹⁸ and with the programmes from HICs⁹⁷ It is important to note that PTA is a crucial test to differentiate CDHL and sensorineural hearing loss. However information on bone conduction testing was was limited.

Few studies reported the use of ICT to screen, manage data or perform diagnostic tests.⁸⁷⁶ Lack of use of ICT could be due to lack of adequate infrastructure, skills to support use of such tools. Yet, this is not unique to L&MICs as evidence on use of ICT is limited even among HICs.^{92 93 97 99}

We assessed the validity and efficacy of the screening programme for infants and older children using a purposively developed tool. None of the programmes reported met all of the criteria. The majority of programmes made use of validated screening and diagnostic tools and reported the rate of hearing loss identification. However, information on economic analysis was scarce, even though cost effectiveness is a key variable for determining programme success.¹⁰⁰ Furthermore, studies predominantly reported only identification but not intervention. The importance of EHDI programmes is to intervene children so that the pervasive impact of childhood hearing loss can be mitigated^{101 102}; therefore, it is

pertinent to know whether such programmes resulted in early intervention.

Mean prevalence of hearing loss in newborns and infants was identified to be high in India (5/1000), followed by Iran (3/1000) and China (2/1000). This is similar to the findings of Bussé and colleagues (2021) where the highest prevalence was found in India and Nigeria, followed by Iran. In another review, prevalence was found to be highest in Asian countries compared with other regions.⁹⁹ A world report on hearing also stated that prevalence of congenital hearing loss in L&MICs is high compared with HICs.

Barriers identified from our review were similar to those previously identified and discussed in various studies including L&MICs.^{97 101-103} However, a recent study in HICs found that when hearing screening programmes were integrated as part of national screening with a dedicated screening person, database management system and appropriate guidelines, they were more successful. Therefore, EHDI in L&MICs is also likely to be more successful when implemented through the government.

There were some limitations to the review which must be considered. No article was excluded based on quality assessment owing to the limited literature available from L&MICs, yet the risk of bias in many included studies was moderate to high. Furthermore, due to heterogeneity in the information obtained across studies, no meta-analysis was performed. The generalisability of the findings was limited to Asian L&MICs. Further, there were potential for publication bias as not all programmes would have published their results. The coverage of EHDI in these countries was not assessed.

From this study, it is evident that strategies for EHDI in Asian L&MICs were similar to those recommended in HICs. However, there is inadequate evidence related to the intended outcome of early intervention in this context. Therefore, programme planners and researchers must focus on impact evaluations that demonstrate the longterm viability of EHDI programmes in the L&MI context.

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