**ORIGINAL RESEARCH** 

# **Prevalence of hearing loss among neonates admitted in NICU in a tertiary care center**

<sup>1</sup>Dr.Guruprasad B, <sup>2</sup>Dr.Mahendra Kumar Banakar, <sup>3</sup>Dr. Bollakonda Maneesh Karthik

<sup>1</sup>Assistant Professor in Paediatrics, Department of Paediatrics, KVG Medical College, Sullia, Karnataka, India
<sup>2</sup>Assistant Professor in Paediatrics, Department of Paediatrics, KVG Medical College, Sullia, Karnataka, India
<sup>3</sup>Junior resident Department of Pediatrics, KVG Medical College, Sullia, Karnataka, India

## **Corresponding Author**

<sup>3</sup>Dr. Bollakonda Maneesh Karthik <sup>3</sup>Junior resident Department of Pediatrics, KVG Medical College, Sullia,Karnataka, India

Received: 29Dec, 2024

Accepted: 30Jan, 2025

## ABSTRACT

Indian studies performed using different hearing screening protocols have estimated the prevalence of neonatal hearing loss to vary between 1 and 8 per 1000 babies screened. The better prognosis of individual skills like language development, academic success, social integration and successful participation in the society can be attained by early identification and intervention for hearing loss i.e. by 6 months of age. After obtaining permission from the institutional ethical committee all neonates admitted in NICU fulfilling the inclusion and exclusion criteria were taken into the study after obtaining written informed consent from parents/guardian. Information regarding the condition of each neonate was collected in the form of a predesigned questionnaire which included: gestational age; family history of congenital hearing loss and consanguinity; presence of conditions including asphyxia(APGAR score<4), sepsis, respiratory distress syndrome, transient tachypnoea of newborn(TTN), congenital pneumonia, congenial heart disease(CHD) or hyperbilirubinemia( $\geq$ 18 mg/d1);and treatments used including phototherapy (>2 days), mechanical ventilation (>5 days), antibiotic therapy including aminoglycosides(>5 days), or oxygen therapy(>1 week and > 40% FIO2).

First OAE was done to all 320 neonates among whom 247 (77.8%) passed and 73 (22.8%) came as refer. Among those neonates who failed 1st OAE at first week of life repeat OAE was done around 45 days of age (6weeks).

Out of 320 neonates, 73 (22.8%) had failed in 1st OAE, among those 73 only 62 neonates came for second OAE (11 missed for follow up). Of these neonates 57 passed (91.93%) and 5 neonates failed 2nd OAE (8.064%). Out of 5 neonates who had failed in 2nd OAE, BERA was done and 4(80%) neonates failed while 1 passed BERA.

Key words: Hearing loss, neonates, BERA

This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-Non Commercial-Share Alike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

# INTRODUCTION

Hearing impairment of children in the world constitutes a serious obstacle to their ideal development, education and language acquisition. The prevalence of neonatal hearing loss is known to be more than twice that of other newborn disorders such congenital hypothyroidism as and phenylketonuria<sup>1,2</sup>.The Incidence of Bilateral congenital hearing impairment alone occurs in approximately 1 to 5 per 1000 live births, whereas when included with permanent unilateral hearing loss, incidence increases to 8 per 1000 live births<sup>3,4,5</sup>.Indian studies performed using different hearing screening protocols have estimated the prevalence of neonatal hearing loss to vary between 1 and 8 per 1000 babies screened<sup>6,7,8,9</sup>. The better prognosis of individual skills like language

development, academic success, social integration and successful participation in the society can be attained by early identification and intervention for hearing loss i.e. by 6 months of  $age^5$ .

The critical period of brain development occurs in the 1st year of life, especially the auditory pathway. An inadequate stimulus during this time lead to suboptimal development of auditory system, whereas optimal auditory experience has a good influence on the functional development<sup>10</sup>. Hence, early detection is extremely important in providing suitable care, provision of hearing aid and special training for deaf and hearing-impaired babies will help them enjoy equal opportunities in society alongside all other children.

American Academy of Pediatrics (AAP) in 1999 advocated Universal New Born Hearing Screening International Journal of Life Sciences, Biotechnology and Pharma Research Vol. 14, No. 2, February 2025

Online ISSN: 2250-3137 Print ISSN: 2977-0122

DOI: 10.69605/ijlbpr\_14.2.2025.180

Programme (UNHSP) and remedial intervention which is followed by most of the developed countries. In a developing country like India, the risk of infants to develop these difficulties is obviously more<sup>11,12</sup>. In India, newborn hearing screening are usually available only to newborns brought to tertiary referral hospitals<sup>13,14</sup>.

There was an introduction of two-stage screening protocol with otoacoustic emission (OAE) as the first screen, followed by auditory brainstem response (ABR) for those who fail the first screen<sup>15</sup>. The AAP Task Force on newborn and infant hearing recommends UNHS by 3 months of age with intervention by 6 months of age. The Joint Committee on Infant Hearing (JCIH) position statement provides guidelines that include Newborn Hearing Screening (NHS) soon after birth, before discharge from hospital, or before 1 month of age, diagnosis of hearing loss through audiological and medical evaluation before 3 months, and intervention through interdisciplinary programme for infants with confirmed hearing loss before 6 months of age<sup>16</sup>.

## **METHODOLOGY**

#### STUDY POPULATION

All neonates admitted in NICU for more than 48 hours.

# **INCLUSION CRITERIA**

a) All neonates admitted in NICU for more than 48 hours.

### **EXCLUSION CRITERIA**

- a) Neonates with congenital anomalies (craniofacial malformation).
- Neonates admitted for less than 48 hours. b)
- c) Neonates with family history of deafness.
- d) Neonates who died or who have not completed OAE.

#### SAMPLE SIZE AND SAMPLING

The sample size for the study was calculated based on the 50.5% prevalence of hearing impairment reported in the study by Nair VS et al., Based on this, the required sample size with 6% absolute precision and a confidence level of 95% was calculated using the formula,

Sample size,  $n = \frac{z^2 pq}{d^2}$ 

Where, z=1.96, Z score for 95% confidence level p=0.50, prevalence of hearing impairment <sup>[13]</sup> q=0.495 d=0.6, absolute precision of 6%. Thus

$$n=1.962 \times \frac{50.5 \times 49.5}{6^2} = 267$$

Taking into account a non-response rate of 20% i.e. 53, the total sample size will be 320.

# SYSTEMATIC SAMPLING

Technique was used where all neonates meeting the criteria was included in the study during the study duration until the sample size was reached.

## STUDY PROCEDURE

After obtaining permission from the institutional ethical committee all neonates admitted in NICU fulfilling the inclusion and exclusion criteria were taken into the study after obtaining written informed consent from parents/guardian. Information regarding the condition of each neonate was collected in the form of a predesigned questionnaire which included: gestational age; family history of congenital hearing loss and consanguinity; presence of conditions including asphyxia(APGAR score<4), sepsis. respiratory distress syndrome, transient tachypnoea of newborn(TTN), congenital pneumonia, congenial heart disease(CHD) or hyperbilirubinemia(≥18 mg/d1);and treatments used including phototherapy (>2)days), mechanical ventilation (>5 days), antibiotic therapy including aminoglycosides(>5 days),or oxygen therapy(>1 week and > 40% FIO2).

The screening procedure was done in a sound treated room in the department or in a quiet room adjacent to the respective wards of concerned departments. The presence of unilateral or bilateral hearing loss was considered as deafness in this study. The Instruments used is transitory evoked otoacoustic emission (TEOAE) set at a 1.5kHZ to 4 kHz screen with 3 of 4 frequency bands being required to be present for a pass. The intensity was calibrated at an 83 dB sound pressure level peak equivalent (3dB).

The firststepof thescreeningwasperformedatdischarge fromNICUwiththeOAE measurement. For babies who failed to pass the initial screening, another OAE test wasperformed within 1 month after discharge, and those failing to pass the test again were referred to a pediatric otologist for comprehensive audiological assessments at 3 months.

# DATA COLLECTION METHODS

Data was collected using a standard proforma on admission. After collecting basic information based on predesigned proforma screening for hearing loss was first performed at the time of discharge from NICU with the OAE measurement. For babies who failed to pass the initial screening, another OAE test was performed on the day of first immunization visit i.e.,6weeks(45days), and those failing to pass 2nd time were referred to a pediatric otologist for comprehensive audiological assessments at 3months.

# RESULTS

Assessment done	Number of neonates	Percentage (%)
1st OAE (n=320)		
Pass	247	77.8
Refer	73	22.8
2nd OAE (n=62)		
Pass	57	91.93
Refer	5	8.064
BERA (n=5)		
Pass	1	20
Refer	4	80

First OAE was done to all 320 neonates among whom 247 (77.8%) passed and 73 (22.8%) came as refer. Among those neonates who failed 1st OAE at first week of life repeat OAE was done around 45 days of age (6weeks).

Out of 320 neonates, 73 (22.8%) had failed in 1st OAE, among those 73 only 62 neonates came for

second OAE (11 missed for follow up). Of these neonates 57 passed (91.93%) and 5 neonates failed 2nd OAE (8.064%). Out of 5 neonates who had failed in 2nd OAE, BERA was done and 4(80%) neonates failed while 1 passed BERA.

Table 2: Comparison of Selected Va	ariables Based o	on Risk l	Factors of First OA	Е
			A 73	

D'-l. Fratan		Number of Frequency		Chi agrant		
Risk Factor		PASS	REFER	Chi-square	P-value	
Castational aga	Pre term	125(73.09%)	46(26.90%)	15.243	0.002	
Gestational age	Term	122(81.87%)	27(18.12%)	13.245		
Digth confermio	Present	19(86.36%)	3(13.63%)	1 1 2 0	0.29	
Birth asphyxia	Absent	228(76.51%)	70(23.48%)	1.138	0.28	
RDS	Present	136(75.55%)	44(24.44%)	0.622	0.22	
KDS	Absent	111(79.28%)	29(20.71%)	0.622	0.32	
MAS	Present	22(70.97%)	9(29.03%)	0.752	0.29	
MAS	Absent	225(77.85%)	64(22.14%)	0.753	0.38	
ΤΤΝΙ	Present	5(83.33%)	1(16.66%)	0.121	0.71	
TTN	Absent	242(76.82%)	73(23.17%)	0.131	0.71	
CLID	Present	27(77.14%)	8(22.85%)	0.000	0.995	
CHD	Absent	225(77.58%)	65(22.41%)	0.000		
<b>C</b> .:	Present	15(51.72%)	14(48.27%)	11.54	0.0001	
Seizures	Absent	232(79.72%)	59(20.27%)	11.74		
C	Present	71(67.61%)	34(32.38%)	0.1.(4	0.0004	
Sepsis	Absent	176(81.86%)	39(18.13%)	8.164		
TT 1'1' 1' '	Present	152(77.15%)	45(22.84%)	0.000	0.99	
Hyperbilirubinemia	Absent	95(77.23%)	28(22.76%)	0.000		
Distations	Present	152(77.15%)	45(22.84%)	0.000	0.862	
Phototherapy	Absent	95(77.23%)	28(22.76%)	0.298		
A	Present	89(70.78%)	38(29.92%)	6.04	0.014	
Antibiotics	Absent	158(81.86%)	35(18.13%)	6.04		
NEC	Present	5(50.0%)	5(50.0%)	4.22	0.031	
NEC	Absent	242(78.06%)	68(21.93%)	4.33		
	Present	155(74.16%)	54(25.83%)	2.121	0.07	
Oxygenrequirement	Absent	92(82.88%)	19(17.11%)	3.131		
	Present	3(0.93%)	0(0.0)	0.005	0.344	
Exchangetransfusion	Absent	244(76.25%)	73(22.81%)	0.895		
	Present	34(10.62%)	10(3.12%)	0.000	0.99	
CongenitalPneumonia	Absent	213(66.56%)	63(19.68%)	0.000		

Significance at the level (p<0.05) chi-square test

Among risk factors, out of 171 preterm,46 failed (26.90%);among 21 neonates with birth asphyxia, 3 (13.63%) failed; out of 180 neonates with RDS,44(24.44%) failed; out of 31 neonates with MAS, 9 (29.03%) failed; among 6 neonates with TTN,1(16.66%) failed; out of 35 neonates with CHD, 8 (22.85%) failed; among 29 neonates with seizures,14(48.27%) failed, out of 105 neonates with sepsis, 34(32.38%) failed; out of 197 neonates with hyperbilirubinemia,197 received phototherapy, of

which 45(22.84%) failed; out of 127 neonates who received aminoglycosides class of antibiotics, 38 (29.92%) failed; 10 neonates who had developed NEC, half of them (50%) failed; out of 44 neonates with congenital pneumonia, 10 (3.12%) failed; among 209 neonates who received oxygen 54 (25.83%) failed first OAE. Outof these factors gestationalage (preterm),seizures, sepsis,NEC and antibiotics values were statistically significant (*p*<0.05).

<b>Risk Factor</b>		Number of Frequency		Chi-square	P-value
KISK Factor		PASS	REFER	Cm-square	<b>r</b> -value
Gestational age	Pre term	35(92.10%)	3(7.89%)	14.321	0.111
Gestational age	Term	22(91.66%)	2(8.33%)	14.321	0.111
Birth asphyxia	Present	1(100.0%)	0(0.0)	3.34	0.341
	Absent	56(91.80%)	5(8.19%)	5.54	0.541
RDS	Present	32(94.11%)	2(5.88%)	1.32	0.751
KD3	Absent	25(89.28%)	3(10.71%)	1.52	0.751
MAS	Present	7(100.0%)	0(0.0)	1.018	0.79
MAS	Absent	50(90.90%)	5(9.09%)	1.018	0.79
TTN	Present	2(100.0%)	0(0.0)	0.106	0.991
IIN	Absent	55(91.66%)	5(8.33%)	0.100	0.991
CHD	Present	8(100.0%)	0(0.0)	1.24	0.745
CHD	Absent	49(90.74%)	5(9.25%)	1.24	
Seizures	Present	12(92.30%)	1(7.69%)	12.92	0.003
Seizures	Absent	45(91.83%)	4(8.16%)	- 13.83	
Sonsia	Present	26(89.66%)	3(10.33%)	9.272	0.026
Sepsis	Absent	31(93.99%)	2(6.06%)	9.272	
Hyperbilirubinemia	Present	41(95.34%)	2(4.65%)	3.02	0.389
Hyperollifuolitetilla	Absent	16(84.21%)	3(15.78%)	5.02	
Dhotothonony	Present	41(95.34%)	2(4.65%)	3.811	0.085
Phototherapy	Absent	16(88.88%)	2(11.11%)	5.811	
Antibiotics	Present	29(90.65%)	3(9.375%)	6.609	0.061
Antibiotics	Absent	28(93.33%)	2(6.66%)	0.009	
NEC	Present	5(100.0%)	0(0.0)	7.366	0.061
NEC	Absent	52(91.22%)	5(8.77%)	7.500	
Ourseen meguinement	Present	41(93.18%)	3(6.81%)	2.25	0.521
Oxygen requirement	Absent	16(88.88%)	2(11.11%)		
Exchange transfusion	Present	0(0.0)	0(0.0)	0.79	0.86
	Absent	58(93.54%)	4(6.45%)	0.78	
Conconital Draumeria	Present	9(14.51%)	0(0.0)	1.004	0.800
Congenital Pneumonia	Absent	49(79.03%)	4(6.45%)	1.004	

Significance at the level (p < 0.05) Chi-square test

Out of 73 neonates who failed in 1st OAE, 2nd OAE was done at the time of first vaccination i.e., 6weeks (45days). Out of 73, 2nd OAE was done only on 62 neonates as others could not be followed up due to increasing COVID cases and Lockdown. Among the risk factors, out of 38 pretern, 3 (7.89%) failed; out of 34 neonates with RDS, 2 (5.88%) failed; out of 13 patients with seizures,1(7.69%) failed; among 29

neonates with sepsis, 3(10.33%) failed; among 43 neonates who received phototherapy, 2(4.65%) failed; among 32 neonates who received antibiotics, 3 (9.37%) failed; out of 44 neonates who received oxygen, 3(6.81%) failed. Out of all these risk factors values of neonates with seizures (p=0.003) and sepsis (p=0.026) were statistically significant.

Dial- Easton		Number of Frequency		Chi agreene	D malma
Risk Factor		PASS	FAIL	Chi-square	P-value
Castational aga	Preterm	1(25.0%)	3(75.0%)	2.249	0.764
Gestational age	Term	0(0.0%)	1(100.0%)	3.348	
	Present	0(0.0%)	0(0.0%)	0.275	0.020
Birth asphyxia	Absent	1(20.0%)	4(80.0%)	0.375	0.829
RDS	Present	1(33.33%)	2(66.66)	0.942	0.656
KDS	Absent	0(0.0%)	2(100.0%)	0.842	0.656
MAC	Present	0(0.0%)	0(0.0%)	0.545	0.762
MAS	Absent	1(20.0%)	4(80.0%)	0.545	0.762
TTNI	Present	0(0.0)	0(0.0)	0.097	0.052
TTN	Absent	1(20.0%)	4(80.0%)	0.097	0.953
CHD	Present	0(100.0)	0(0.0)	- 5	0.2
CHD	Absent	1(0.0%)	4(100.0%)	5	0.2
S	Present	0(0.0)	1(100.0%)	0.212	1
Seizures	Absent	1(0.0%)	3(100.0%)	0.313	
Concio	Present	0(62.5%)	3(100.0%)	2.744	0.152
Sepsis	Absent	1(50.0%)	1(50.0%)	3.744	
Umarhilimhinamia	Present	1(33.33%)	2(66.66%)	0.850	0 (54
Hyperbilirubinemia	Absent	0(80.0%)	2(100.0%)	0.850	0.654
Dh at ath avarage	Present	1(33.33%)	2(66.66%)	0.956	0.931
Phototherapy	Absent	0(80.0%)	2(100.0%)	0.856	
Antibiotics	Present	0(0.0%)	3(100.0%)	2.756	0.252
Antibiotics	Absent	1(50.0%)	1(50.0%)	2.730	
NEC	Present	0(0.0)	0(0.0)	0.164	0.921
NEC	Absent	1(20.0%)	4(80.0%)	0.104	
Oww.gon Baguinomont	Present	0 (0.0%)	2(100.0%)	0.042	0.623
Oxygen Requirement	Absent	1(33.33%)	2(66.66%)	0.942	
Euchon as transfusion	Present	0(0.0)	0(0.0)	0.049	0.946
Exchange transfusion	Absent	1(20.0%)	4(80.0%)	0.048	
Concentral Draumania	Present	0(0.0)	0(0.0)	0.910	0.667
Congenital Pneumonia	Absent	1(20.0%)	4(80.0%)	0.810	0.007

## DISCUSSION

Hearing loss can be considered as the most common birth defect. Early detection of hearing impairment by screening at or shortly after birth helps in appropriate intervention that are critical for speech, language and cognitive development. Universal Neonatal Hearing Screening (UNHS) is done in all child irrespective of presence or absence of risk factor to identify hearing impairment as early as possible to provide interventions.

This study was conducted for a period of 10 months, included a total of 320 neonates after excluding newborns who died (10), early discharge before OAE screening (8) and having family history of hearing loss (3). First OAE was done on 320 neonates of which 73 (22.8%) failed. This high number of false positives (22.8%) may be due to premature outer hair cells or due to incomplete clearance of normal fetal

middle ear fluid as first OAE is done before discharge, which are the reasons to verify these results later with 2nd OAE after 6 weeks or with more specific methods such as BERA. Out of 73 who failed first OAE, 11 neonates were missed due to COVID and various other reasons so 2nd OAE was done only on 62 neonates, of which 5 (1.56%) failed and all these 5 neonates were followed up and among them 4 (1.25%) failed in BERA.

In our study 320 neonates were included and it showed that 4 out of 320 neonates failed BERA. Thus, the prevalence of hearing loss was 1.25%. A study by James M *et al.*, 48 found the prevalence rate among high-risk newborns as 0.63% which is almost similar to our study. Following table shows prevalence of hearing loss among high-risk newborns in various studies.

Study on hearing loss in high	Prevalence per 100	Prevalenceper100 with	Prevalenceper
risk newborns	with first OAE	Second OAE	100BERA
Our study (n=320)	73 (22.8%)	5(1.56%)	4 (1.25%)
JamesMetal. <sup>17</sup> (n=4628)	275 (6.4%)	44 (0.95%)	6 (0.63%)
Nair VS et al., <sup>18</sup> (n=200)	101 (50.5%)	-	1 (0.5%)
PaulAK <i>et al.</i> <sup>19</sup> (n=2031)	234 (11.52%)	78 (3.84%)	21 (1.03%)
Jose <i>et al.</i> , <sup>20</sup> (n=231)	38 (6.1%)	4 (1.73%)	2 (0.86%)

**Table 5: Comparison with other studies** 

Prevalence of individual risk factor as in JCIH (Joint committee on infant hearing) includes 171 neonates are preterm (53.4%), 22 neonates had perinatal asphyxia (6.7%),hyperbilirubinemia requiring phototherapy were 197 neonates (61.6%), 209 (65.3%) required oxygen and ototoxic medicine were given to 127 neonates (39.7%).

Present study found prematurity (p=0.002) to be significantly related to failure of initial screening. Similar observation was seen in study by Nair VS et al.,(p=0.01), James M et al., (p=0.000) and Pourarian S et al., (p=0.013) where prematurity was significantly associated with hearing loss. Prematurity should be considered as one of the risk factors for hearing loss in NICU population, as their respiratory system is not fully developed which necessitates the oxygen requirement and infections occurring due to their weekend immune system which requires antibiotic coverage. Among these variables' we found that antibiotics (aminoglycosides) given (p=0.014) was significantly associated with hearing loss which was similar to the finding in a study by James M et al. (p=0.000), oxygen requirement was also seen to be associated with hearing loss (p=0.07) but it was not statistically significant. This was similar to the finding in the study by Pourarian S et al.

Present study also showed that 10 (3.12%) neonates out of 320 had developed NEC, among which 5 (1.56%) of them failed the initial screening (1st OAE) which was statistically significant (p=0.031) but all these 5 neonates passed in 2nd OAE.

We also found 29 (9.1%) neonates out of 320 had developed seizures of which 14 (4.37%) failed the initial screening and out of 14 neonate's 2nd OAE was done on 13 neonates as 1 missed follow up, among these 13 neonates 1 (7.69%) failed the 2nd OAE screening, both of which are statistically significant (p=0.0001 and p=0.003 respectively). A study by Bergman I *et al.*, showed that 16.7% of neonates surviving seizures had developed hearing loss. However, there is lack of research on the correlation between seizures and NEC with hearing loss which needs further studies to find the association.

Sepsis is found to be associated with the failure of both 1st and 2nd OAE screening and is statistically significant (p=0.0004 & p=0.026) which is similar to the study by James M *et al.*, but is in contrast the study by Pourarian *et al.*, which may be explained by the smaller sample size.

Present study didn't show any statistically significant correlation between RDS,TTN, Congenital Pneumonia, CHD and Birth asphyxia with hearing loss, which is similar to the finding from the study by Pourarian *et al.*,

As described above many risk factors have been found to be associated with transient hearing loss among neonates (failure of initial screening). But the same association was not established for actual hearing impairment found in BERA. This could be attributed to premature outer hair cells in newborns or other physiological changes in newborn ear.

# CONCLUSION

Even though high risk newborns are at greater risk, hearing evaluation is important in all newborn irrespective of presence of risk factor for hearing impairment. Two step evaluation with OAE followed by BERA are tend to be useful. And also, while dealing with these newborns unnecessary oxygen therapy and antibiotics should be avoided. Hearing aids are to be advised to those babies with hearing impairment as early as possible to avoid significant morbidity and disabilities associated with hearing loss.

## REFERENCES

- Fisher DA, Dassault JH, Foley TP, Klein AH, Lanfranchi S, Larsen PR, *et al.*, Screening for congenital hypothyroidism: results of screening one million North. American infants.J Pediatr. 1979;94:700-5.
- Bickel H, Bachmann C, Beckers R, Brandt NJ, Clayton BE, Corrado G, *et al.*, Neonatal mass screening formetabolic disorders: summary of recent sessions of the committee of experts to study inborn metabolic diseases. Eur J Pediatr.1981;137:133–9.
- 3. Mehra S, Eavey RD, Keamy DG Jr. The epidemiology of hearing impairment in the United States: newborns, children, and adolescents. Otolaryngol Head Neck Surg. 2009;140:461-72.
- Stach BA, RamachandranVS. Hearing disorders in children. In: Madell JR, Flexer C eds. Pediatric Audiology: Diagnosis, Technology, and Management. New York: Thieme Medical Publishers Inc.; 2008. P. 3-12.
- 5. Judith A, Mason MS, Kenneth R, Herrmann MD.Universal infant hearing screening by automated auditory brainstem response measurement. Pediatrics. 1998;101:221-8.
- Jane E. Stewart, Aimee Knorr, Hearing Loss in Neonatal Intensive Care Unit Graduates: Manual of Neonatal Care. A Lippincott Manual. 7<sup>th</sup> edition. 2012: 65:846-850.
- 7. Nagapoornima P, Ramesh A, Srilakshmi, Rao S, Patricia PL, Gore M, *et al.*, Universal hearing screening. Indian J Pediatr. 2007;74:545-9.
- 8. Paul AK. Early identification of hearing loss and centralized newborn hearing screening facility-The Cochin experience.Indian Pediatr. 2011;48:355-9.
- Rai N, Thakur N. Universal screening of newborns to detect hearing impairment-Is it necessary? Int. J PediatrOtorhinolaryngol.2013;77:1036-41.

- Yoshinaga-Itano C, Sedey AL, Coulter DK, Mehl AL. Language of early and lateridentified children with hearing loss. Pediatrics. 1998;102:1161-1171
- 11. Report of the Collective Study on Prevalence and Etiology of Hearing Impairment. New Delhi: ICMR and Department of Science; 1983.
- 12. Kacker SK. The Scope of Pediatric Audiology in India. In: Deka RC, Kacker SK, Vijayalakshmi B, eds. Pediatric Audiology in India, 1st ed. New Delhi: Otorhinolaryngological Research Society of AIMS; 1997.p.20.
- Vaid N, Shanbag J, Nikam R, Biswas A. Neonatal hearing screening-The Indian experience. Cochlear Implants Int. 2009;10:111-4.
- Ramesh A, Nagapoornima M, Srilakshmi V, Dominic M. Swarnarekha. Guidelines to Establish a Hospital-based Neonatal Hearing Screening Programme in the Indian Setting. JAIISH.2008;27:105-9.
- 15. Watkin PM, Baldwin M, McEnery G. Neonatal at risk screening and the identification of deafness. Arch Dis Child. 1991;66:1130-1135.
- 16. Joint Committee on Infant Hearing; American Academy of Audiology; American Academy of Pediatrics; American Speech-Language-Hearing Association; Directors of Speech and Hearing Programmes in State Health and Welfare Agencies. Year 2000 Position Statement: Principles and guidelines for early hearing detection and intervention programmes. Pediatrics.2000;106;798-817.
- James M, Kumar P, Ninan PJ. A study on prevalence and risk factors of hearing Impairment among newborns. Int. J Contemp Pediatr. 2018;5:304-9
- Nair VS, Das P, Soundararajan P. Prevalence and risk factors of hearing impairment among neonates admitted in NICU in a tertiary care centre in South India. Int. J Contemp. Pediatr.2018;5:1342-7.
- Paul AK. Early identification of hearing loss and centralized newborn hearing screening facility: The Cochin experience. Indian Pediatr. 2011;48:355-9.
- 20. Jose DJ, Renjit RE, Manonmony S. Prevalence of hearing impairment among high risk neonates-a hospital-based screening study. Int. J Biomed Adv. Res. 2016;7(3):131-4.