

ORIGINAL PAPER

Sudden sensorineural hearing loss in children: Our experiences in tertiary care teaching hospital of eastern India

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ABSTRACT

Aim of the study: Sudden sensorineural hearing loss (SSNHL) is an uncommon clinical entity in children. The etiopathogenesis, clinical profile and management are not well understood and the incidence is not well known in medical literature. The goal of this study is to evaluate prevalence, etiology, treatment and outcome of SSNHL in the pediatric patients.

Material and methods: We had undertaken a retrospective study of the children below 18 years of age with SSNHL treated between July 2014 to August 2017. The data assessed were age, sex, clinical manifestations, onset of hearing loss, audiological profile, other diagnostic tools, treatment and outcome.

Results: The average age of the children presenting with SSNHL is 10.43 years (1-18 years). Twelve patients (6 right and 6 left) had unilateral hearing loss and 5 patients had bilateral hearing loss. Hearing loss ranges from profound (52.94%) to mild and unknown etiology in 70.58% and viral 29.41%. Hearing was completely improved in 5 cases (29.41%), partially improved in 8 cases (47.05%) and no improvement in 3 cases (17.64%). When comparing oral with oral plus intratympanic (IT) steroid, hearing output is statistically significant ($p = 0.04020$) with 95% confidential level is 0.69 to 0.02 and $t = 2.1381$.

Conclusions: The exact incidences of SSNHL in pediatric age groups are not established in medical literature. Pediatric patients are usually delayed presentations and associated with unknown etiology in majority of cases. SSNHL has serious impact on life and it should be followed up for preventing morbidities and for starting rehabilitation program as early as possible.

KEY WORDS:

sudden sensorineural hearing loss, children, tinnitus, steroid.

INTRODUCTION

Sudden sensorineural hearing loss (SSNHL) is a rare clinical entity in children. Sudden hearing loss was first

documented by de Kleyn in 1944 and defined as hearing loss of at least 30 decibel (dB) in three consecutive frequencies in the standard pure tone audiogram over three days or less [1]. SSNHL in children is considered as an

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otologic emergency and rarely reported in medical literature. The incidence of SSNHL ranges between 10.7 and 27 per 100,000 persons per year [2]. The incidence of SSNHL increases with age, with 8 per 100,000 under the age of 18 years and 70 per 100,000 over 65 years [2]. Early treatment is necessary for preventing permanent hearing loss. This is important in children as it affects speech and hearing if not improved which also affects academic and social performance. Often the children are unable to report their hearing loss, which may be not detected in acute stage when treatment is the most effective. The degree of hearing loss has an effect on the rate of recovery [3]. The causes of SSNHL in adults are usually idiopathic whereas the idiopathic cases are rare in children due to the small size of the available series. Presently the treatment protocol of SSNHL is available which includes systemic steroids as primary treatment and intratympanic steroid administration when there is no improvement after systemic steroid treatment or contraindications for systemic steroid [4]. This study discusses about prevalence, etiology, clinical profile and outcome of SSNHL among children in a tertiary care teaching hospital of eastern India.

MATERIAL AND METHODS

We performed a retrospective study of patients below 18 years of age between July 2014 to August 2017 with sudden sensorineural hearing loss. This study was approved by institutional ethical committee (IEC). The gap between sudden onsets of hearing loss and starting the treatment was within two weeks. All the children depending on the patient's age and developmental level, the testing methodology like visual reinforcement audiometry, conditioned play audiometry and conventional pure tone audiometry. The normal hearing is defined at ≤ 20 dB. In our study, the degree of hearing loss ranges from mild sensorineural hearing loss to profound loss across frequencies. The degree of hearing loss was determined as mild (20-40 dB), moderate (40-60 dB), moderate to severe (60-75 dB), severe (75-90 dB) and profound (> 90 dB). All patients with sudden hearing loss had done pure tone audiometry at frequencies between 250 Hz to 8000 Hz and had to be at least more than 20 dB in 3 consecutive frequencies in comparison to the unaffected ear. All children reported abrupt onset of hearing loss. All the children with hearing loss were treated with steroid. Eleven cases were treated with oral prednisolone given at an initial dose of 3 mg/kg body weight. The dose of the prednisolone was reduced every second day and for the maximum of 14 days or completed 2 days after the hearing had normalized in pure tone audiometry. In few cases, intravenous (IV) hydrocortisone 1 mg/kg/day given in divided doses. Whenever there was no improvement of hearing after systemic therapy or contraindication for steroid therapy, intratympanic injection of steroid like dexamethasone 1 mg every 12 hours for 7 days was given

through grommet inserted into the tympanic membrane. After intratympanic injection of steroid, children were positioned at the non-affected side for 3 minutes. All intratympanic injections were done under all aseptic precautions. Another pure tone audiometry was performed 2 to 3 weeks after therapy and combined with brainstem evoked response audiometry. The follow up for all children were done at 3 months, 6 months and 1 year. Complete hearing improvement was defined as hearing level is similar to the non-affected ear, partial hearing improvement is defined when more than 10 dB in at least one frequency and no improvement of hearing when there is no change in hearing in audiogram after treatment. In our study, data recorded are age, gender, symptoms associated with hearing loss like aural fullness, tinnitus, vertigo, and otalgia. Onset of hearing loss, audiological results, serological results, imaging, treatment and its outcome were documented.

RESULTS

We diagnosed 17 children with SSNHL where 11 were male and 6 were female child. All children were healthy and without any family history of hearing loss. All were presented with abrupt onset of hearing loss. Mean age was 9.64 years (1-18 years). The right ear was affected in 6 (35.29%) children, the left ear in 6 (35.29%) and both ears in 5 (29.41%) children. Fourteen children (82.35%) complained other symptoms in addition to hearing loss and most common one was tinnitus. Three patients complained otalgia (17.64%), two patients (11.76%) complained vertigo and one (5.88%) presented herpetic lesions but serology for herpes simplex was negative. There was higher incidence of profound hearing loss in 9 cases (52.94%), severe hearing loss in 3 cases (17.64%), moderate hearing loss in 3 cases (17.64%) and mild in 2 cases (11.76%). Initial treatment was oral steroids in 11 cases (64.70%), IV steroids in 6 cases (35.29%) and 6 cases (35.29%) undergone intratympanic steroids along with 6 oral steroids. The systemic steroids were continued for 1-2 weeks (median = 7 days). In three cases, parents of children decided to discontinue this treatment due to fear of side effects. Intratympanic steroids were given in 6 cases (35.29%) and lasted between 6 and 8 days (mean = 7). One patient stopped IT treatment after 4 days. There were no adverse effects following either systemic or IT steroid treatment. Non-contrast CT scan of temporal bone in 5 patients and MRI in 11 patients were within normal limits. None of the imaging revealed any pathology. Serological study revealed cytomegalovirus (CMV) in 3 cases, Epstein-Barr virus (EBV) IgM in one, EBV IgM in 1 case. Hearing was completely improved in 5 cases (29.41%), partially improved in 8 cases (47.05%) and there was no improvement in 3 cases (17.64%) (Table 1). There is no statistically significant between oral and IV steroid ($p = 0.729$) but when comparing oral with oral

TABLE 1. Clinical and audiological profile of the children

Serial No	Age (years)	Gender	Side of the ear	Day of the treatment started	Other symptoms	Steroids	Outcome
1	7	Male	Right	2	Tinnitus	Oral	PI
2	3	Male	Right	4	Tinnitus, fullness	Oral	PI
3	11	Female	Bilateral	0	Tinnitus	IV	CI
4	5	Male	Left	3	Otalgia	Oral, IT	PI
5	18	Male	Right	7	Tinnitus	Oral, IT	PI
6	9	Female	Left	1	Tinnitus	IV	PI
7	1	Female	Left	20	Tinnitus	Oral	NI
8	8	Male	Bilateral	5	Vertigo	Oral	NI
9	12	Male	Right	3	Tinnitus	IV	CI
10	14	Male	Bilateral	2	Tinnitus	Oral, IT	CI
11	4	Male	Left	32	Tinnitus, fullness	Oral	NI
12	7	Female	Right	4	Tinnitus	Oral	PI
13	9	Female	Bilateral	1	Vertigo	IV	PI
14	12	Male	Left	0	Tinnitus, otalgia	IV, IT	PI
15	13	Female	Left	1	Tinnitus	IV	PI
16	16	Male	Bilateral	2	Herpetic lesions, otalgia	Oral, IT	CI
17	15	Male	Right	0	Tinnitus	Oral, IT	CI

IV – intravenous; IT – intratympanic; PI – partial improvement; CI – complete improvement; NI – no improvement

plus IT steroid, it is statically significant ($p = 0.04020$) with 95% confidential level is 0.02 to 0.69 and $t = 2.1381$. The statistical analysis evaluated by SPSS 20 software. One patient was lost follow up.

DISCUSSION

Sudden sensorineural hearing loss (SSNHL) is an otological emergency in medicine. SSNHL in children is a rare clinical entity and less frequently discussed in the medical literature and the exact incidence is not documented. The causes of SSNHL in adults may be due to meningitis, labyrinthitis, temporal bone trauma, autoimmune disorders, retro-cochlear pathology, ototoxic medications, central nervous lesions and stroke [5]. However in majority of cases the cause is not clearly identified and is presumed to be vascular insults or infections [6]. In pediatric patients, the SSNHL is less likely by vascular insults and retro-cochlear pathology [7]. Majority of the cause for SSNHL is idiopathic in this study. One study reported Lyme disease and EBV are leading cause for SSNHL [7]. Congenital cytomegalovirus, radiographic findings of enlarged vestibular aqueduct and non-organic hearing loss are other causes and not common etiology in adult patients. In present study, no etiology was found in 70.58 % of the subjects. Adult patients with SSNHL usually complain hearing loss much earlier than children and sometimes present with associated symptoms like tinnitus, vertigo and otalgia. Our patients in this study also

presented with similar associated symptomatology. However, these presentations may be delayed in pediatric age groups. The delay in presentation will delay the treatment and can affect the outcome of the treatment. Hearing loss at very early age of life affects speech and language. It also affects academic and social performance in life. The diagnosis of SSNHL sometimes difficult particularly at a very young age, therefore a high index of suspicion is needed. The role imaging in SSNHL is not well defined and it does not have impact on the treatment. However to level the SSNHL to idiopathic, MRI is needed to rule out acoustic neuroma as it is seen in 2% cases of SSNHL and it is associated with partial recovery [8]. Currently American Academy of Otolaryngology guidelines recommended MRI for SSNHL because sudden hearing loss are associated with a retro-cochlear lesions in MRI [9]. The MRI is important diagnostic tool to rule out acoustic neuroma, which is not common in pediatric patients [9]. There is still a debate in pediatric literature whether to use CT scan or MRI for diagnosis of sensorineural hearing loss [10]. Serial audiometric testing are done along with the imaging for documenting progression of hearing loss. All patients had undergone pure tone audiometry at the time of first visit and next after completion of the treatment. Auditory brainstem response (ABR) in children is done to rule out non-organic hearing loss [11]. Laboratory testing were not done in our patients. Routine laboratory evaluation has low diagnostic yield and should only performed in case clinical suspicion [12]. It has seen that severe

hearing loss has poor prognostic factor [13]. Treatment of SSNHL is a subject of debate. Presently the available treatment options are systemic steroids, intratympanic steroids, both systemic and intratympanic steroids and hyperbaric oxygen therapy. The drugs like anti-viral, vasodilators, thrombolytic, vasoactive substance and anti-oxidants are not usually advocated in the treatment of SSNHL [6]. Oral steroids are usually given in the initial treatment for SSNHL. In one study, showed efficacy of oral steroids with 61% of patients improved with oral steroids whereas only 32% improved with placebo [14]. Intratympanic injections of steroid is another alternative to the oral steroid. Intratympanic steroid provides localized treatment with higher concentration at perilymph and avoid systemic side effects. Few complications may occur like otalgia during and after treatment, tympanic membrane perforation, dysgeusia, vertigo and transmission of infection to the middle ear. It has been seen that oral and intratympanic steroid treatments have equally efficacious in the treatment of SSNHL [15]. Gundogan *et al.*, showed better hearing outcome with both oral and intratympanic steroids than either one alone [16]. Hyperbaric oxygen therapy (HBOT) is another treatment option in medical literature. HBOT decreases the edema and hypoxia and may help with immune response to ischemia and infection [17]. It may be helpful in case of failed traditional treatment options [18]. The combined treatment of steroids and HBOT has better outcomes in comparison to medical treatment alone [19]. HBOT is costly and has potential adverse effects and power of the many studies is low. None of our patients under went HBOT in present study. The rate of recovery of hearing in case SSNHL among children varies between the different studies. In one study with 14 patients under the age of 18 years treated with IV prednisolone, the rate of complete recovery was 57% whereas the partial recovery rate was 36% [20]. Tarshish *et al.* showed complete recovery in 10% cases and some improvement in 10% cases after treatment with systemic steroids and without intratympanic steroids [21]. In our study, hearing completely improved in 5 cases (29.41%) and partial improved in 8 cases (47.05%) and no improvement in 3 cases (17.64%). Therefore, more such studies are required to elaborate on the recovery rate from SSNHL in children. When comparing recovery rate with adults, the complete recovery is higher and no improvement rate is significantly lower in children than adults [3]. In both pediatric and adult populations, hearing recovery is significantly higher in patients with mild SSNHL compared to profound SSNHL, but in children with moderate hearing loss, the rate of recovery is higher in comparison to adults, revealing that age is associated with a poorer prognosis [22]. The lower incidence of SSNHL in children is seen medical literature and it is still not clear work-up and treatment giving successful outcome for this population. The etiological role of viral infections is still not clear and it seems that pediatric CMV infection may play

role in SSHNL. Similar to adult patients, intratympanic steroids may be considered for the treatment along with systemic corticosteroids.

CONCLUSIONS

SSNHL in children is a rare clinical entity. This study showed that the most common etiology of SSNHL in pediatric age group is idiopathic. Detail history taking and serological evaluations for ruling out viral etiology are important for diagnosis and treatment. Most of the children in our study showed improvement in hearing. Intratympanic steroid has additional benefit to the patient along with systemic steroids. A larger study is needed to establish the standardize the diagnosis and treatment. The diagnosis of SSNHL should be prompt and time waste can be prevented by not doing unnecessary investigations. SSNHL has serious impact on life and it should be follow up for preventing morbidities and for starting rehabilitation program as early as possible.

This study is limited by its retrospective nature and the small number of patients. However, it provides further data on this subject.

DISCLOSURE

The authors declare no conflict of interest.

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