Therapeutic Effects of Carbogen Inhalation and Lipo-Prostaglandin E1 in Sudden Hearing Loss

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Purpose: Vascular disorders and viral infections are considered the main causes of sudden hearing loss (SHL), although its pathogenesis remain unclear. Treatments include carbogen inhalation and lipo-prostaglandin E1 (lipo-PGE1), both of which have circulation-enhancing effects. We investigated the effectiveness of carbogen inhalation and lipo-PGE1 in SHL. Materials and Methods: This retrospective review included 202 patients with idiopathic SHL who visited our clinic within 14 days of symptom onset between January 2006 and June 2010. All patients received oral prednisolone for 10 days. Of the 202 patients, 44 received no additional treatment, 106 received additional carbogen inhalation, and 52 received additional lipo-PGE1. Hearing improvement was measured using Siegel’s criteria. Results: Overall recovery rates were 67.9% in the carbogen group, 53.8% in the lipo-PGE1 group, and 52.3% in the steroid-only control group (p=0.097). Limited to type 1 and type 2 categories of Siegel’s criteria, the carbogen group had a significantly higher recovery rate (53.8%) than the lipo-PGE1 group (26.9%) and the steroid-only control group (38.6%) (p=0.005). Conclusion: Carbogen inhalation added to steroid was a more effective treatment than lipo-PGE1 added to steroid or steroid alone in patients with SHL.

Key Words: Sudden hearing loss, lipo-prostaglandin E1, carbogen

INTRODUCTION

Sudden hearing loss (SHL) is considered a syndrome and not a diagnosis and defined as a hearing loss of at least 30 dB in three sequential frequencies occurring over 3 days or less.1 SHL was first reported by De Kleyn2 in 1944, there have been many studies regarding the disease, but the pathogenesis remains unclear. SHL involves a variety of causative factors, and therefore should be considered a syndrome rather than a single disease.3 The most common causes include viral infection and vascular disorders; other causes include rupture of the inner ear membrane, immune disorders, and acoustic tumors.4
The most widely accepted treatment for SHL is systemic corticosteroids,\(^5\) due to their ability to reduce inflammation, inhibit immune mechanisms, and regulate electrolyte balance. However, studies have shown that corticosteroid treatment is not significantly more effective than placebo, and includes serious side effects.\(^6,7\) Thus, there have been continuous efforts to identify additional treatments.\(^8\) In the case of SHL caused by viral infection, studies on antiviral therapy have shown controversial therapeutic results.\(^9\)

Vasodilators such as carbogen inhalation and lipoprostaglandin E1 (lipo-PGE1) have been used to treat SHL based on the theory that vasodilatation of the inner ear may aid recovery if caused by a vascular disorder. Lipo-PGE1 is a prostanoid that acts as a vasodilator and improves vascular circulation. Carbogen inhalation increases arterial oxygen saturation and maximizes oxygen supply to the inner ear. Studies on the effects of carbogen and lipo-PGE1 in SHL have had controversial results. Shea and Kitabchi\(^10\) reported that carbogen inhalation had a therapeutic effect, while Cinamon, et al.\(^6\) found no therapeutic effect. Similarly, Zhuo, et al.\(^11\) reported that lipo-PGE1 had a therapeutic effect, while Ahn, et al.\(^12\) found no therapeutic effect. Despite controversial benefits, the theories remain that improving circulation may yield therapeutic results in cases where a vascular disorder underlies the SHL.

In this study, we compared 2 vasodilator treatments for SHL, carbogen inhalation and lipo-PGE1, to investigate their therapeutic effects.

**MATERIALS AND METHODS**

**Patients**

This retrospective review was performed in 236 patients with SHL who visited our clinic between January 2006 and June 2010. All patients underwent a thorough history and physical examination, pure tone audiometry, speech audiometry, and impedence audiometry. Auditory Brainstem Response latency and temporal magnetic resonance imaging were selectively performed to rule out retrocochlear pathology. To minimize the bias, we excluded patients who presented more than 14 days of symptom onset and patients with other causes of hearing loss such as Meniere’s disease, vestibular schwannoma, and ischemic brain lesion including AICA infarct.

All patients received identical initial treatment consisting of 1 mg/kg of prednisolone (Solondo\(^5\), 5 mg tablets, Yuhan Corp., Seoul, Korea) administered orally for 5 days and then tapered for 5 days (a half-dose on days 6 and 7, 20 mg on days 8 and 9, and 10 mg on day 10). Patients were divided into 3 groups: the steroid-only control group consisted of 44 patients who presented between January 2006 and May 2007 and received no additional treatment, the carbogen group consisted of 106 patients who presented between June 2007 and June 2009 and received additional carbogen inhalation treatment, and the lipo-PGE1 group consisted of 52 patients who presented between July 2009 and June 2010 and received additional lipo-PGE1 treatment.

Carbogen inhalation treatment involved inhaling mixed gas (5% CO\(_2\), 95% O\(_2\)) once a day for 1 hour for 5 days. Lipo-PGE1 treatment involved daily intravenous injection of 5 μg lipo-PGE1 mixed with 500 mL normal saline for 5 days.

**Therapeutic effect analysis**

The follow-up hearing testing schedule is given in Fig. 1. The Therapeutic effects were evaluated at least 2 months after end of treatment which known for hearing stabilazation,\(^13\) using the pure tone average (average of the 0.5, 1.0, 2.0, and 3.0 kHz hearing thresholds). We used Siegel’s criteria for hearing improvement analysis (Table 1). ‘Overall recovery’ was defined as showing any recovery after the conclusion of treatment (Siegel’s criteria I, II, and III). ‘Favorable recov-

![Fig. 1. Diagram for treatment and follow-up schedule. After a 10 day-course of corticosteroids and additional treatments, pure tone audiograms were measured. The final hearing test was performed at least 2 months after end of treatment. OPD, outpatient department.](image-url)
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en (45.5%). Their mean age was 49.6±16.9 years (range, 13 to 89). The average time between symptom onset and presentation to clinic was 3.5±3.1 days. There were no significant differences in age, gender, disease duration, follow-up period, or initial hearing threshold of the affected ear between groups (Table 2).

Hearing change
Overall average hearing thresholds before and after treatment were 73.2±27.6 dB and 47.1±31.7 dB, respectively. Overall recovery was achieved by 123 patients (60.9%), 88 (43.6%) of whom achieved favorable recovery, which corresponds to Siegel’s criteria I and II.

Comparing the recovery rate by the criteria of overall recovery, 72 of 106 (67.9%) in the carbogen group, 28 of 52 (53.8%) in the lipo-PGE1 group, and 23 of 44 (52.3%) in the control group attained hearing recovery. The higher rate of overall recovery in the carbogen group was not statistically significant ($p=0.097$). However, when comparing favorable recovery, the carbogen group (53.8%) experienced a significantly higher recovery rate than did the lipo-PGE1 group (26.9%) or the control group (38.6%) ($p=0.005$) (Fig. 2).

In regards to RHG, the carbogen group had higher RHG (29.58 dB) than did the lipo-PGE1 group (20.15 dB) or the control group (21.48 dB), but the difference was not statistically significant ($p=0.033$) (Table 3).

Word recognition score improved in correlation with av-

**RESULTS**

**Patient characteristics**
202 out of 236 patients with SHL were included in this study and were consisted of 110 men (54.5%) and 92 women (45.5%). Their mean age was 49.6±16.9 years (range, 13 to 89). The average time between symptom onset and presentation to clinic was 3.5±3.1 days. There were no significant differences in age, gender, disease duration, follow-up period, or initial hearing threshold of the affected ear between groups (Table 2).

**Statistical analysis**
Significant differences between groups were determined using the chi-square test, Fisher's exact test, and one-way ANOVA, with $p<0.05$ defined as the cut-off for statistical significance. All statistical analysis was performed using SPSS v16.0.

| Table 1. Siegel’s Criteria of Hearing Recovery |
|-----------------------------------------------|
| Type                                           |
| I. Complete recovery                           |
| Final hearing better than 25 dB                |
| II. Partial recovery                           |
| More than 15 dB gain, final hearing 25-45 dB   |
| III. Slight improvement                        |
| More than 15 dB gain, final hearing poorer than 45 dB |
| IV. No improvement                             |
| Less than 15 dB gain, final hearing poorer than 75 dB |

| Table 2. General and Hearing Characteristics of Patients |
|----------------------------------------------------------|
| Clinical characteristics                                  |
| Control (n=44)                                            |
| Lipo-PGE1 (n=52)                                         |
| Carbogen (n=106)                                         |
| $p$ value                                                 |
|-----------------------------------------------------------|
| Age (yrs)                                                 |
| 48.61±17.55                                              |
| 44.88±17.45                                              |
| 50.92±15.37                                              |
| 0.095*                                                    |
| Gender (male : female)                                   |
| 25 : 19                                                   |
| 28 : 24                                                   |
| 57 : 49                                                   |
| 0.947†                                                   |
| Ear (right : left)                                        |
| 21 : 23                                                   |
| 18 : 34                                                   |
| 55 : 51                                                   |
| 0.123†                                                   |
| Symptom duration (days)                                  |
| 3.57±3.51                                                |
| 3.23±2.85                                                |
| 3.60±2.99                                                |
| 0.762*                                                   |
| Mean follow-up (days)                                    |
| 54.35±29.10                                              |
| 50.74±16.44                                              |
| 51.67±40.31                                              |
| 0.925*                                                   |
| Rates of systemic disease (%)                            |
| 5/44 (11.4)                                              |
| 4/52 (7.7)                                               |
| 14/106 (13.2)                                            |
| 0.698†                                                   |
| Initial hearing threshold (dB)                           |
| 74.64±29.36                                              |
| 81.00±28.75                                              |
| 70.74±26.38                                              |
| 0.091*                                                   |
| Initial pattern of hearing (LFHL : HFHL : flat : others)  |
| 9 : 10 : 11 : 13                                         |
| 10 : 8 : 20 : 14                                         |
| 15 : 20 : 31 : 40                                        |
| 0.587†                                                   |

LFHL, low frequency hearing loss; HFHL, high frequency hearing loss; lipo-PGE1, lipo-prostaglandin E1.

*One way ANOVA test was used.
†Chi-square test was used.
‡Fisher’s exact test was used.
in overall and favorable recovery ($p=0.712$ and $p=0.756$, respectively) (data not shown).

Complications

There were no major complications during treatment. Although several patients experienced severe hyperglycemia, the patients’ blood glucose level were normalized after the end of treatment. The incidence of hyperglycemia was not significantly different between groups (data not shown) and it was likely due to the side effect of the systemic corticosteroids.

DISCUSSION

Although the underlying mechanism of SHL is unknown,
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