

Multimodal Therapy for Chronic Tinnitus

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Abstract: From 2001 to 2006, we performed a retrospective study of patients suffering from chronic unilateral or bilateral tinnitus that was previously ineffectively treated by oral drugs [betahistine (Betaserc), extract of *Ginkgo biloba* (EGb 761), tanakan (Tebokan), and cinnarizine-dimenhydrinate (Arlevert), singly or in combination]. We divided 150 tinnitus patients (80 men, 70 women) into seven treatment groups. Treatments consisted of application of intravenous pentoxifylline, lidocaine, or vinpocetine (Cavinton) and combination of these agents with physiotherapy and soft laser. Mean duration (\pm standard deviation) of tinnitus in these patients was 7.4 ± 6.0 years; their mean age was 55.6 ± 12.5 years. The aim of our study was to compare treatment modalities and define their effectiveness for tinnitus relief. The most effective treatment was defined as a combination of Cavinton and physiotherapy. We evaluated pure lidocaine infusion therapy as ineffective. None of the treatment modalities had an objective correlate of improvement, though improvement was reported by a visual analog scale.

Key Words: chronic tinnitus; infusion therapy; physiotherapy; soft laser

Tinnitus is the phantom perception of sound in the absence of overt acoustic stimulation [1]. It is well known that chronic tinnitus is difficult to treat, though lots of modalities are used for treatment. Guidelines for inclusion of each patient into an appropriate treatment group are still missing. In our study, we evaluated some of the current treatment modalities to define the most effective for a certain group of patients.

Current possibilities for chronic tinnitus treatment vary from neuroprotective substances [2], calcium-channel blockers, corticosteroids, glutamate agonists, and thrombolytic drugs [3–5] to intravenous or intratympanic lidocaine application [6–10]; vinpocetine (Cavinton), pentoxifylline (Agapurin, Trental), or piracetam infusions [4]; low-level laser therapy (so-called soft laser) [11–13]; physiotherapy, acupuncture, tinnitus retraining therapy, or other approaches. Because the studies available on MEDLINE report different outcomes [8,10–12,14], we tried to evaluate the most widely used therapeutic approaches in our hospital: drop infusion of lidocaine, vin-

pocetine, or pentoxifylline and combination of these drugs with low-level laser therapy and physiotherapy.

SUBJECTS AND METHODS

Patient Population

We divided 150 chronic tinnitus patients (80 male, 70 female; mean age, 55.6 ± 12.5 years) into seven study groups (Table 1). The mean duration of treatment was 9.2 ± 3.2 days, and the mean duration of tinnitus was 7.4 ± 6.0 years. All patients had previously been treated ineffectively by various oral drugs [betahistine (Betaserc), extract of *Ginkgo biloba* [EGb-761], tanakan [Tebokan], cinnarizine-dimenhydrinate (Arlevert)], or their combination] for at least 6 months. Patients with decompensated systemic or psychiatric disease were excluded from the study.

Study Design

Permanent tinnitus remaining for more than 6 months was considered to be chronic. Before and after treatment, each patient underwent basic ear, nose, and throat examination, including otomicroscopy, basic vestibular examination, pure-tone audiometry, and tinnitometry.

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Table 1. Demographic and Anamnestic Data of the Patient Population

Regimen	Male/		Age ± SD (yr)	Duration of Tinnitus ± SD (yr)
	No.	Female		
PTX	22	8/14	55.6 ± 11.1	9 ± 6.5
Lidocaine	17	9/8	54.7 ± 12.6	4.0 ± 2.8
Lidocaine-RHB	28	17/11	54.4 ± 9.5	5.3 ± 5.2
Lidocaine-RHB-soft laser	29	13/16	51.8 ± 16.4	6.8 ± 6.6
Vinpocetine	17	9/8	58.7 ± 14.2	9.2 ± 7.5
Vinpocetine-RHB	20	14/6	59.1 ± 12.0	8.8 ± 7.1
Vinpocetine-RHB-soft laser	17	10/7	55.2 ± 11.5	7 ± 6.2
Total	150	80/70	55.6 ± 12.5	7.4 ± 6.0

PTX = pentoxifylline; RHB = physiotherapy; SD = standard deviation.

We randomly divided the patients into seven distinct treatment groups and administered treatment as specified.

- The *pentoxifylline (PTX) group* received daily for 10 days an intravenous infusion of 100 mg pentoxifylline and 250 ml physiological saline.
- Those in the *Cavinton group* received daily for 10 days an intravenous infusion of 20 mg Cavinton plus 250 ml physiological saline, and oral Cavinton 2 × 10 mg (0–2–2).
- To those in the *lidocaine group*, we gave a daily intravenous infusion of 40 mg lidocaine hydrochloride plus 250 ml physiological saline for 10 days.
- The *Cavinton-plus-physiotherapy (Cavinton-RHB) group* received a daily intravenous infusion of 20 mg Cavinton plus 250 ml physiological saline, and oral Cavinton 2 × 10 mg, combined with physiotherapy (soft techniques, including temporomandibular joint massage), the entire procedure lasting 20 minutes once daily for 10 days.
- For the *Cavinton-RHB-soft laser group*, we administered a daily intravenous infusion of 20 mg Cavinton plus 250 ml physiological saline, and oral Cavinton 2 × 10 mg, combined with soft laser

(5,000 BTL; 400 mW/830 nm; pulse beam, 30 J/cm³) once daily for 10 minutes and physiotherapy, all for 10 days.

- Those in the *lidocaine-RHB group* were given a daily intravenous infusion of 40 mg lidocaine plus 250 ml physiological saline, combined with physiotherapy, for 10 days.
- To those patients in the *lidocaine-RHB-soft laser group*, a daily intravenous infusion of 40 mg lidocaine plus 250 ml physiological saline, combined with physiotherapy and soft laser (for soft-laser specifications, see earlier in this paragraph), were given for 10 days.

In addition, there were groups treated by *pentoxifylline plus physiotherapy (PTX-RHB)* and *pentoxifylline plus physiotherapy and soft laser (PTX-RHB-soft laser)*, which consisted of a total of 9 persons. As the outcomes were considered to be statistically insignificant, the findings from these groups are not evaluated further.

The visual analog scale from 0 (no tinnitus) to 10 (unbearable tinnitus) was used for daily evaluation. Patients' symptoms were evaluated as improved if their tinnitus was perceived to be better by two or more points on this scale.

STATISTICAL EVALUATION

We used an enhanced method for construction of a binary classification tree—a regression method computed by condition of maximized homogeneity of treatment methods used [by Agapurin (Trental), Cavinton, lidocaine] in terminal leaves. We had four metrics—age, duration of tinnitus, gender, and audiometry—each of which could be used to divide cases into branches. According to audiometry, we had three subgroups: bilateral normal hearing, unilateral normal hearing, and bilateral hearing loss. We crossed all boundaries that are created as a midpoint between all recorded numerical data on all metrics

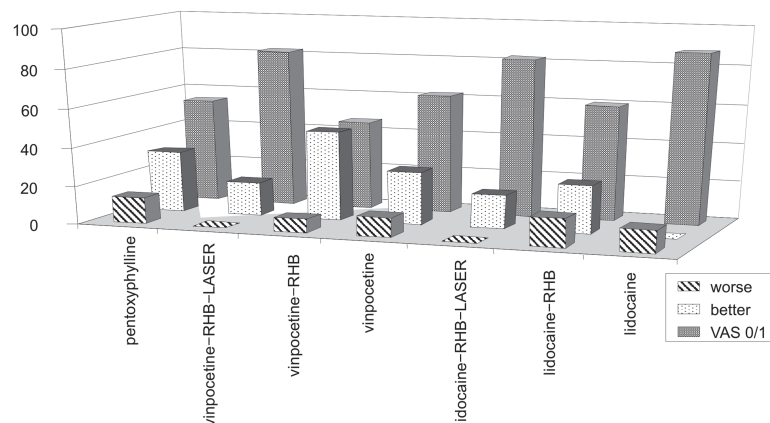


Figure 1. Effectiveness of therapy in each of seven treatment groups, determined by using a visual analog scale. Change is noted in percentages.

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