

## Vocal quality of patients treated for laryngeal tuberculosis, before and after speech therapy

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### Abstract

**Objectives:** To evaluate dysphonia in patients treated for laryngeal tuberculosis, and to assess the effect of speech therapy on patients' vocal quality.

**Materials and methods:** Seven of 23 patients with a confirmed diagnosis of laryngeal tuberculosis, treated at the Evandro Chagas Institute of Clinical Research, Oswaldo Cruz Foundation, underwent speech therapy for six months. These seven patients were evaluated by videolaryngoscopy and vocal acoustic analysis, before, during and after a course of speech therapy.

**Results:** The 23 patients with laryngeal tuberculosis comprised five women and 18 men, with ages ranging from 25 to 83 years (mean 41.3 years). Dysphonia was present in 91.3 per cent of these laryngeal tuberculosis patients, being present as the first symptom in 82.6 per cent. In laryngeal tuberculosis patients with dysphonia, laryngeal tuberculosis treatment resulted in dysphonia resolution in only 15.8 per cent. After speech therapy, dysphonia patients had better vocal quality, as demonstrated by statistical analysis of jitter, shimmer, fundamental frequency variability, maximum phonation time, and the ratio between maximum phonation time for voiceless and voiced fricative sounds.

**Conclusions:** Following treatment of laryngeal tuberculosis, the incidence of dysphonia was very high. Speech therapy improved patients' vocal quality.

**Key words:** Tuberculosis; Larynx; Dysphonia; Speech Therapy

### Introduction

Tuberculosis (TB) is a contagious, infectious disease of chronic evolution. Histologically, it is characterised by the presence of granulomas and central caseous necrosis, caused by *Mycobacterium tuberculosis*. Transmission occurs predominantly by air and particularly affects the lungs, but TB can occur in any organ.

Following the appearance of acquired immunodeficiency syndrome, TB has grown in importance.<sup>1,2</sup> Human immunodeficiency virus infection has changed not only the epidemiological trends of TB but also its clinical presentation, with a rising incidence of extra-pulmonary forms.<sup>3</sup>

Laryngeal TB is the most frequent granulomatous laryngeal disease.<sup>4,5</sup> Usually as secondary to pulmonary TB, but studies present up to 20 per cent of prevalence for primary laryngeal TB<sup>6</sup>, i.e. without pulmonary or any other form.

Two theories are accepted to explain the pathophysiology of laryngeal TB: bronchogenic and haematogenic. In the first, the laryngeal mucosa is contaminated by direct contact with bronchial secretions, explaining the preferential involvement of the vocal

folds and the posterior commissure.<sup>7</sup> In the second, the larynx is infected by circulatory transmission via blood or lymph, compromising preferably the false vocal folds, epiglottis and aryepiglottic folds.<sup>7,8</sup>

Tubercular lesions in the laryngeal mucosa may have various appearances, including granular, nodular, ulcerated, hyperaemic and/or oedematous.<sup>8</sup> The locations of the lesions, in decreasing order of incidence, are the vocal folds, epiglottis, false vocal folds, aryepiglottic folds and the interarytenoid region.<sup>9</sup>

Dysphonia is the main symptom of laryngeal TB, being present in 96.6 per cent of cases,<sup>8</sup> followed by odynophagia, dysphagia, odynophonia and cough.<sup>6,9,10</sup> After recovery from laryngeal TB, one would expect voice production to improve as the laryngeal inflammatory process resolved; however, we could find no reports addressing the evolution of dysphonia after laryngeal TB treatment.<sup>11</sup>

The vocal folds are formed in layers from the superficial layer of the lamina propria, flexible and rich in elastic fibres, to the vocal muscle, more rigid and rich in collagen fibres. Any change in one or more of these layers may alter their function.<sup>12–14</sup>

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The fibrosis associated with mucosal healing causes stiffness, which interferes with vocal production.<sup>9,15,16</sup>

This study aimed to evaluate the incidence, evolution and outcome of dysphonia in patients treated for laryngeal TB, as well as to assess the effect of speech therapy on patients' vocal quality.

### Materials and methods

A prospective, longitudinal, cohort study was performed. Initially, we reviewed the medical records of patients with laryngeal TB treated at the Evandro Chagas Institute of Clinical Research, Oswaldo Cruz Foundation, from 2000 to 2008, in order to identify those who had been treated for laryngeal TB and had vocal disorders, but who had not received previous speech therapy. These patients were invited to participate in the study. Those who agreed, and who gave written, informed consent, were included in the cohort.

Otorhinolaryngological evaluation consisted of videolaryngoscopy using a rigid optic Karl Storz 70° laryngoscope (Karl Storz, Tuttlingen, Germany) to assess the presence or absence of laryngeal lesions and their location.

Speech evaluation was performed at the first consultation and then every three months.

Acoustic speech analysis was undertaken using Vox Metria software (CTS Informática, Pato Branco, Brazil).<sup>17</sup> Patients' voices were recorded in a quiet room, in a digital computer file for best voice capture. A microphone was used (A-20 model; Plantronix, São Paulo, Brazil) with a mouth--microphone distance of 10 cm. The patient was asked to produce a sustained /e/ vowel in a normal fashion.

Acoustic analysis evaluated the following parameters: The indexes of disturbance related to the fundamental frequency are, jitter, and shimmer variability to semitones. The fundamental frequency is the number of cycles of vibration of the vocal folds per second, and its variability leads to oscillation of cycles; the fundamental frequency variability is considered abnormal when increased by more than two semitones. Jitter and shimmer supply information on the similarity of successive glottal cycles and the stability of the glottal source.<sup>11</sup> Jitter indicates the short-term variability of the fundamental frequency, and is considered abnormal when greater than 0.6 per cent. Shimmer indicates the short-term variability in the amplitude of the sound wave, and is considered abnormal when greater than 6.5 per cent. The glottal to noise excitation ratio calculates the acoustic noise in a series of pulses produced by the oscillation of the vocal folds; a value greater than 0.5 is considered to indicate an abnormal trend. The voice irregularity parameter represents the correlation between jitter and shimmer, and is regarded as abnormal if greater than 4.75.<sup>13,18</sup> The maximum phonation time was evaluated using a digital timer (model TI5G811; Timex, Manaus, Brazil), with the patient asked to sustain the vowel /e/; a time of less than 10 seconds was considered reduced. We also evaluated the ratio between the maximum phonation times of the voiceless fricative /S/ and the

voiced fricative /Z/; a value greater than or equal to 1.2 was considered to indicate lack of glottal closure.<sup>13,19</sup>

Data were processed as follows: arrangement of study protocol variables, data entry, data quality control, and data analysis using the Statistical Package for the Social Sciences version 11.0 software package.

During exploratory data analysis, the simple frequencies of categorical variables and summary measures (i.e. mean and standard deviation (SD)) of quantitative variables were described.

We applied the Student's *t*-test to compare the means of paired data (evaluation of voice quality before and after speech therapy), considering a significance level of 5 per cent.

This project was approved by the Research Ethics Committee of the Evandro Chagas Institute of Clinical Research, Oswaldo Cruz Foundation (protocol number 0043.0.009.000-07).

### Results and analysis

We retrieved the clinical records of 23 patients with a confirmed diagnosis of TB and laryngeal lesions. These patients comprised five women and 18 men, with a mean age  $\pm$  SD of 41.3  $\pm$  13.9 years at the time of illness.

Dysphonia was present in 21 (91.3 per cent) of the 23 patients with laryngeal TB, and constituted the first symptom in 19 (82.6 per cent) of these 23. After treatment, dysphonia was completely resolved in 15.8 per cent of patients, reduced in 68.4 per cent and unchanged in 15.8 per cent.

Nine patients accepted the invitation to participate in the study. One had a normal voice at the time of first evaluation, and another left the study before commencement of speech therapy. Thus, seven patients (five men and two women) underwent longitudinal study involving voice assessment and speech therapy. All these patients participated in speech therapy, and were evaluated three times over a six month period.

The results of videolaryngoscopic evaluation before and after speech therapy are presented in Table I. In patients one and two, biopsy was used to investigate possible associated disease. Videolaryngoscopic views for patient five, before and after speech therapy, are shown in Figure 1.

The parameters selected for evaluation of voice quality, and the results of statistical analysis, are shown in Table II.

Figure 2 shows changes in patients' vocal quality, indicated by the voice irregularity parameter, over the treatment period. All patients experienced a reduction in voice irregularity, and five attained a degree of irregularity within the normal range ( $p = 0.002$ ).

### Discussion

The residual dysphonia in patients treated for laryngeal TB has not attracted much interest in the literature. However, its high incidence (84.2 per cent) and its impact on quality of life prompted us to investigate

TABLE I  
PATIENTS\* VIDEOLARYNGOSCOPIC FINDINGS BEFORE AND AFTER SPEECH THERAPY

Pt no	Before speech therapy			After speech therapy		
	Cicatricial lesion location	Cicatricial lesion characteristics	Glottal chink?	Cicatricial lesion location	Cicatricial lesion characteristics	Glottal chink?
1	Epiglottis L FVF	Nodular destruction	Invisible	Epiglottis L FVF	Nodular Destruction	No
2	Arytenoids L & R FVF L Ary F L TVF	Infiltration Granulomatous Granulomatous Granulomatous	Irregular	Arytenoids L & R FVF L Ary F L TVF	Infiltration Granulomatous Granulomatous Granulomatous	Irregular
3	Arytenoids L TVF	Hyperaemia Thickened	Posterior, medium, triangular	No lesion	No lesion	No
4	Epiglottis Arytenoids L & R TVF	Oedema Hyperaemia	Ampoule	Epiglottis	Destruction	No
5	L & R Ary F	Oedema	Fusiform	No lesion	No lesion	No
6	Epiglottis Arytenoids	Oedema Oedema	No	No lesion	No lesion	No
7	L & R FVF L & R TVF	Oedema hyperaemia	No	No lesion	No lesion	No

\*Initial *n* = 9. Pt no = patient number; L = left; R = right; FVF = false vocal fold; TVF = true vocal fold; Ary F = aryepiglottic fold

the influence of speech therapy on its evolution. In this study, speech therapy significantly improved voice quality in all patients.

Consistent with other studies, we found a higher incidence of laryngeal TB in males,<sup>1,6,8,11</sup> with an average patient age of approximately 40 years.<sup>2,6,8</sup>

As in other studies,<sup>1,2,6,8,10</sup> laryngeal TB lesions were found most frequently on the vocal folds. This finding explains the fact that dysphonia is the primary symptom of laryngeal TB, caused by impairment of vibratory movement and glottal closure.<sup>5,9</sup>

The laryngeal pathology we observed after laryngeal TB treatment (and before speech therapy) can be explained by the inflammatory process of laryngeal TB and the compensatory mechanisms developed

after treatment.<sup>12</sup> The cicatricial tissue observed is made up of dense collagen fibres which are more rigid than the normal vocal fold mucosa. The location of the scar may vary, affecting one or both vocal folds, and may create asymmetry. The scar is localised, so the associated stiffness is restricted to the portion affected, and compensatory mechanisms may facilitate the development of secondary disorders.<sup>5,9</sup> We also observed glottal chinks, which may develop due to loss of vocal fold vibration (caused by laryngeal TB) preventing complete glottal closure. In these cases, an improvement in the amplitude of vibratory movement may occur after laryngeal TB treatment, although insufficient to allow complete glottal closure.<sup>9</sup>

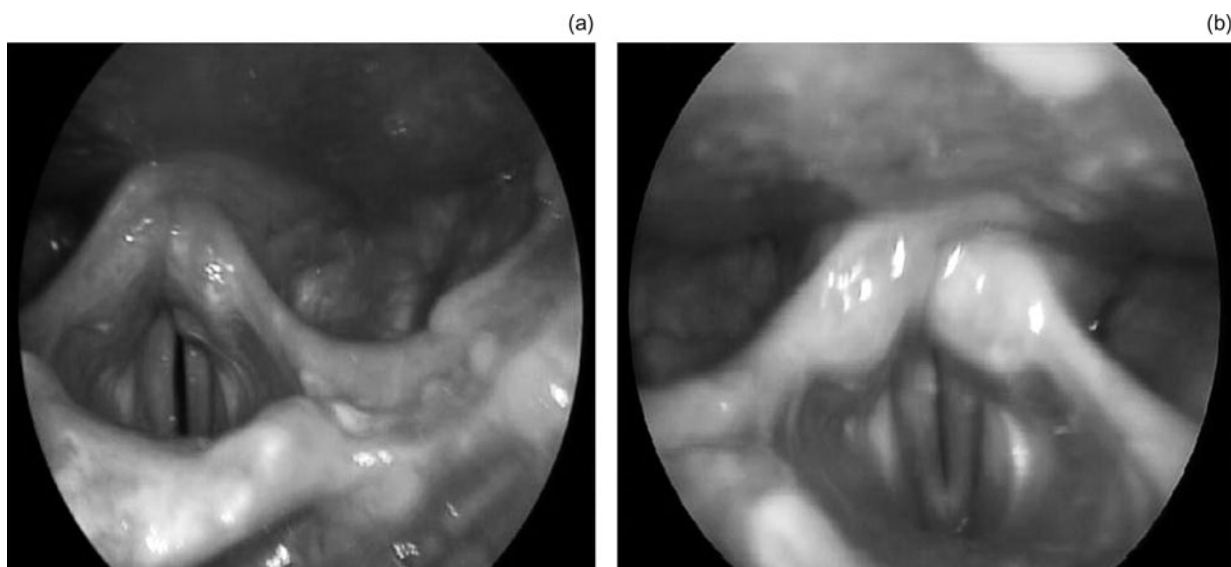


FIG. 1

Videolaryngoscopic views for patient five: (a) fusiform chink seen before speech therapy, and (b) full glottal closure after three months of speech therapy.

TABLE II

PATIENTS' VOICE PARAMETERS BEFORE AND AFTER SPEECH THERAPY

Parameter	Pre-therapy	Post-therapy	<i>p</i> *
Jitter (%)	2.01 ± 1.3	0.48 ± 0.42	0.017
Shimmer (%)	12.21 ± 6.25	6.03 ± 3.13	0.025
GNE ratio	0.64 ± 0.27	0.67 ± 0.19	0.632
VF <sub>0</sub> (semitones)	9.71 ± 8.04	2.30 ± 1.97	0.032
MPT (seconds)	8.00 ± 0.58	17.86 ± 2.19	0.000
S/Z ratio	1.74 ± 0.55	1.10 ± 0.22	0.008

Data represent means ± standard deviations unless otherwise specified. \*Student's *t*-test for paired data. GNE = glottal to noise excitation; VF<sub>0</sub> = fundamental frequency variability; S/Z = ratio between maximum phonation time for /S/ vs /Z/

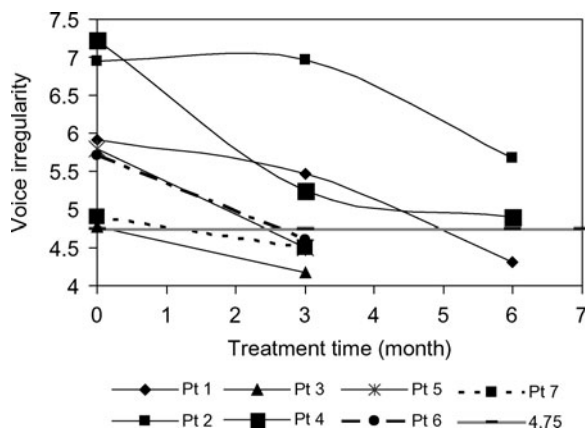


FIG. 2

Change in patients' voice irregularity over the speech therapy treatment period. (A voice irregularity value of more than 4.75 was considered abnormal.)

According to Yelken *et al.*,<sup>11</sup> the action of anti-TB drugs improves voice quality. Unlike the present study, these authors reported the absence of dysphonia after laryngeal TB treatment. However, other studies have noted the permanence of dysphonia after laryngeal TB treatment.<sup>5,9</sup> This difference may be related to the location of lesions, and the time elapsed between infection and its diagnosis and treatment.

- **Tuberculosis (TB) is an infectious disease of chronic evolution, histologically characterised by the presence of granulomas and central caseous necrosis, and caused by *Mycobacterium tuberculosis***
- **Laryngeal TB is the most frequent granulomatous laryngeal disease**
- **This study evaluated dysphonia after laryngeal TB treatment, and the effect of subsequent speech therapy on vocal quality**
- **There is a high incidence of dysphonia after laryngeal TB treatment; such dysphonia can be improved by speech therapy**

We could find no previous reports of speech therapy in patients with dysphonia after laryngeal TB treatment. The results of the current study indicated statistically significant improvements in jitter, shimmer, fundamental frequency, maximum phonation time and S/Z ratio. After speech therapy, all patients had complete glottal closure, except for one who had an associated vocal fold injury under ongoing investigation. The lack of statistically significant improvement of the glottal to noise excitation ratio, following speech therapy, was probably due to noise remaining in the voice because of an irreversible process associated with cicatrization.<sup>5,9</sup> Even when some degree of dysphonia remained, patients reported less vocal fatigue and more comfortable speech.

### Conclusion

The permanence of dysphonia after the treatment of laryngeal TB leads us to conclude that affected patients need assistance in rehabilitating their damaged speech functions. In this study, speech therapy had a positive effect on vocal rehabilitation in such patients, and resulted in improved voice quality.

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### References

- 1 Agarwal P, Bais AS. A clinical and videostroboscopic evaluation of laryngeal tuberculosis. *J Laryngol Otol* 1998;**112**:45–8
- 2 Kandiloros DC, Nikolopoulos TP, Ferekidis EA, Tsangaroulakis A, Yiotakis JE, Davilis D *et al.* Laryngeal tuberculosis at the end of the 20th century. *J Laryngol Otol* 1997;**111**:619–21
- 3 Nasti G, Tavio M, Rizzardini G, De Paoli P, Morassut S, Barzan L *et al.* Primary tuberculosis of the larynx in a patient infected with human immunodeficiency virus. *Clin Infect Dis* 1996;**23**:183–4
- 4 Yencha MW, Linfesty R, Blackmon A. Laryngeal tuberculosis. *Am J Otolaryngol* 2000;**21**:122–6
- 5 Pease BC, Hoasjoe DK, Stucker FJ. Videostroboscopic findings in laryngeal tuberculosis. *Otolaryngol Head Neck Surg* 1997;**117**:230–4
- 6 Porras AE, Martin MA, Perez RJ, Avalos SE. Laryngeal tuberculosis. *Rev Laryngol Otol Rhinol* 2002;**123**:47–8
- 7 Rodriguez BR, Rodriguez BA, Vidal JL, Noguera AA. Dysphonia and laryngeal tuberculosis: presentation of two cases and review of the literature [in Spanish]. *Aten Primaria* 2002;**30**:530–2
- 8 Lim JY, Kim KM, Choi EC, Kim YH, Kim HS, Choi HS. Current clinical propensity of laryngeal tuberculosis: review of 60 cases. *Eur Arch Otorhinolaryngol* 2006;**263**: 838–42
- 9 Ozudogru E, Cakli H, Altuntas EE, Gurbuz MK. Effects of laryngeal tuberculosis on vocal fold functions: case report. *Acta Otorhinolaryngol Ital* 2005;**25**:374–7
- 10 Lindell MM, Jing BS, Wallace S. Laryngeal tuberculosis. *Am J Roentgenol* 1977;**129**:677–80
- 11 Yelken K, Guven M, Topak M, Gultekin E, Turan F. Effects of antituberculosis treatment on self assessment, perceptual analysis and acoustic analysis of voice quality in laryngeal tuberculosis patients. *J Laryngol Otol* 2008;**122**:378–82

- 12 Hirano MB. *Stroboscopic Examination of the Larynx* [in Portuguese]. Artes médicas, Porto Alegre, Brazil, 1997; 35–7
- 13 Behlau M. *Laryngeal Anatomy and Physiology of Vocal Production* [in Portuguese]. Revinter, Rio de Janeiro, Brazil, 2001
- 14 Hirano S, Bless DM, Massey RJ, Hartig GK, Ford CN. Morphological and functional changes of human vocal fold fibroblasts with hepatocyte growth factor. *Ann Otol Rhinol Laryngol* 2003;**112**:1026–33
- 15 Woo P, Casper J, Colton R, Brewer D. Diagnosis and treatment of persistent dysphonia after laryngeal surgery: a retrospective analysis of 62 patients. *Laryngoscope* 1994;**104**: 1084–91
- 16 Hansen JK, Thibeault SL. Current understanding and review of the literature: vocal fold scarring. *J Voice* 2006; **20**:110–20
- 17 Caporrino Neto J, Cervantes O, Jotz GP, Abrahão M. *Granulomatous Diseases of Larynx* [in Portuguese]. Acta AWHO, São Paulo, Brazil, 1998;6–10
- 18 Carrillo L, Ortiz KZ. Vocal analysis (auditory - perceptual and acoustic) in dysarthrias. *Pro Fono* 2007;**19**:381–6
- 19 Garcia RT, Garcia RA, Diaz RT, Canizo RA. The outcome of hydration in functional dysphonia [in Spanish]. *An Otorrinolaringol Ibero Am* 2002;**29**:377–91

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