Laryngeal electromyography in dysphonic patients with incomplete glottic closure

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Abstract

The lack of specificity in laryngoscopical examination requires that the diagnosis of superior laryngeal and recurrent laryngeal nerve involvement be carried out with the aid of electromyography.

Objective: This study aims to assess the electrophysiological function of the superior and inferior laryngeal nerves by measuring the electrical activity of the muscles they innervate in dysphonic patients with incomplete closure of the vocal folds during phonation.

Method: Thirty-nine patients with incomplete glottic closure were enrolled in a prospective study and had their cricothyroid, thyroarytenoid, and lateral cricoarytenoid muscles examined bilaterally through electromyography. Insertion activity, electrical activity at rest (fibrillation, positive wave and fasciculation) and during muscle voluntary contraction (recruitment, amplitude, potential length and latency between electrical activity and phonation) were measured.

Results: No altered test results were observed for parameters insertion activity and electrical activity at rest. None of the patients had recruitment dysfunction. The mean electrical potential amplitude values were within normal range for the tested muscles, as were potential durations and latency times between the onset of electrical activity and phonation.

Conclusion: No signs of denervation were seen in the thyroarytenoid, cricothyroid, and lateral cricoarytenoid muscles of the studied patients.
INTRODUCTION

The clinical relevance of electromyography (EMG) has been recognized in recent years\(^1\). EMG has become an important method in assessing laryngeal diseases, more specifically in patients with involved vocal fold mobility. The technical difficulty that characterizes this test, however, has limited its use. It is particularly useful in diagnosing inferior motor neuron disorders, differentiating fixation paralyses, and in locating lesions and establishing prognosis\(^2\), as it is the most objective method to assess muscle activity\(^3\).

Inferior laryngeal nerve unilateral involvement can be diagnosed by analyzing the patient’s phonation and the observation of varying degrees of breathiness. When the superior laryngeal nerve is involved, subjects have difficulty adjusting voice loudness and singing.

Recurrent laryngeal nerve dysfunction is also easily verified through laryngoscopy, as vocal fold mobility is impaired in these cases. The alterations consequent to the loss of function of the cricothyroid muscle - innervated by the superior laryngeal nerve - are not evident, and thus hamper diagnostic efforts even with the aid of videolaryngoscopy\(^4\). The most common alterations are amplitude and mucosal wave phase asymmetries, delayed abduction and adduction of the involved vocal fold, and glottic clefts\(^5\), all of which are non-specific findings. Therefore, isolated cases of superior laryngeal nerve paralysis or paresis with preserved vocal fold mobility - an occurrence of remarkable relevance for voice professionals - are hard to diagnose. While disorders involving the recurrent laryngeal nerve and the lateral cricoarytenoid (LCA) muscle are more easily diagnosed in clinical examination by the absence of rotation of the arytenoid cartilage and fixed position of the vocal folds, the isolated involvement of branches of the thyroarytenoid muscle introduces only a few disturbances. Examination with a laryngoscope reveals little evidence of involvement, such as non-specific spindle-shaped clefts\(^6\)-\(^7\).

Superior laryngeal nerve injuries can be diagnosed through EMG of the cricothyroid muscle\(^8\). The examination of the muscles innervated by the recurrent laryngeal nerve requires accurate knowledge on the anatomy of the larynx and experience with EMG. EMG of the thyroarytenoid and LCA muscles assesses the activity of the branches of the recurrent laryngeal nerve in the main larynx adductors, and rules out the need to examine the bilaterally innervated interarytenoid muscles.

This study aimed to assess the electrophysiological function of the recurrent laryngeal and superior laryngeal nerves by looking into the electrical activity of the muscles they innervate in patients with phonation disorders and incomplete coaptation of the vocal folds during phonation as seen in video laryngoscopy.

METHOD

This project was approved by the Research Ethics Committee at the Federal University of São Paulo – Paulista Medical School (permit n° 0122/04).

This cross-sectional prospective study enrolled individuals seen between October 2005 and March 2007 with complaints of dysphonia and incomplete closure of the vocal folds observed in video laryngoscopy. The subjects underwent laryngeal examination by EMG. A control group was not required, as the parameters have been clearly established.

Enrollment criteria: subjects had to be 18 or older, present complaints of dysphonia and incomplete glottic closure as observed in video laryngoscopy, except for presence of a posterior triangular cleft in females.

Exclusion criteria: lack of vocal fold mobility as seen in laryngoscopy, presence of lesions or degenerative disease in the vocal folds.

Sample characterization: 39 subjects, 16 males and 23 females, aged between 18 and 80 years, with a mean age of 51.

Procedure

All patients underwent ENT examination including video laryngostroboscopy or fibrolaryngoscopy with stroboscopy. The subjects were examined through EMG after ENT examination.

Video laryngostroboscopy was carried out with the patients under topical anesthesia with 10% lidocaine; subjects were asked to utter vowel ‘é’ as closely as possible to the way they would in spontaneous phonation. EMG was performed by the same examiner for every patient. Patients were asked to sit and place their heads against a slightly tilted head rest, at an angle of approximately 30 degrees. Before the test was done the patients had their skin cleaned with 70% ethanol. EMG examination was done using a Nilon Kohden® Neuropack device with a 10-kHz low pass filter and a 10-kHz high pass band filter with two channels. The first channel was used for the needle electrode recording muscle electrical activity. The second channel was used to capture the sounds uttered by the patient into a microphone positioned 10 cm away from their mouths. The ground wire was placed close to the patients’ sternum. Time was measured in units of 10 ms and with sensitivity ranging from 100 to 500 µV. All exams were carried out using a concentric bipolar needle electrode.

Examination was conducted as follows, starting on the left side, then moving on to the right side, in accordance with the technique:

1. Palpation of the neck and identification of the anatomic landmarks to position the needle: thyroid and cricothyroid cartilages, cricothyroid membrane.
2. Introduction of the bipolar concentric electrode by percutaneously puncturing the skin over the cricothyroid membrane approximately 0.5 cm to 1.0 cm away from the median sagittal line, between the upper border of the cricoid cartilage and the lower border of the thyroid cartilage, all the way to the cricothyroid muscle at a depth of 0.5 cm to 1.0 cm. The muscle is identified by the presence of audible potentials verified on a computer screen as the patients are asked to produce high frequency sounds.

3. Percutaneous introduction of the needle through the cricothyroid membrane, driving the needle at an angle of 30 to 45 degrees in the lateral and superior directions depending on the subject's neck anatomy, until the thyroarytenoid muscle is reached, at a depth of approximately two centimeters.

4. Introduction of the needle by percutaneously puncturing the skin over the cricothyroid membrane in the lateral portion of the larynx, in the area close to the end of the thyroid cartilage, until the LCA muscle is reached.

EMG was performed to assess spontaneous and voluntary muscle activity; the data sets gathered from the examinations were recorded onto the device's hard drive for further analysis. The following parameters were recorded for each studied muscle: electrical activity at rest, i.e., during normal breathing; electrical activity as patients uttered vowel 'e' at their usual intensity and frequency levels as the thyroarytenoid and LCA muscles were tested; and electrical activity as patients produced high frequency sounds to assess the cricothyroid muscle. The analysis of EMG records included assessment of insertion activity levels, spontaneous activity levels at rest, and activity levels during phonation.

Insertion activity is the electrical response obtained as the needle is driven into the muscle, probably generated by the mechanical process of the needle penetrating the muscle tissue. This parameter was categorized as normal, reduced, or increased.

Spontaneous activity is the electrical response generated spontaneously while the muscle is at rest and the needle is kept still. This parameter was categorized as absent - usually observed in normal muscle tissues - or present. Presence of spontaneous activity was characterized by the identification of fibrillations, positive waves, fasciculations, and high frequency complexes. Fibrillations are low amplitude potentials (10 to 200µV) of short duration (1 to 2 ms). Positive waves present a positive component in amplitudes ranging between 20 and 200 µV and a negative minimum component. Fasciculations are discharges with varied morphology arising from one or part of a motor unit. High frequency discharges are bizarre discharges of variable duration identified mainly by sound.

The analysis of voluntary activity included the assessment of the following parameters:

1. Motor unit recruitment patterns were categorized as normal or with rarefaction, based on the presence of peaks leading to the disappearance of the baseline, at an acquisition rate of 100 ms/cm.

2. Wave duration is the time between the point at which the wave starts from the baseline and its return. This parameter was measured using the software on the EMG device. A window was manually configured to identify the beginning and the end of the wave, and the software calculated its duration. The durations of two clearly defined waves were measured to allow the calculation of the mean duration value. Mean values were categorized as normal (3-7 ms)14-26, increased, or reduced.

3. Motor unit action potential (MUAP) amplitude is the measurement obtained between the most negative and the most positive peaks found for the wave. It was measured using a software that used the data stored on a text file to find the mean values of the acquired potentials by calculating the square roots of the potentials to the square of each wave component, thus covering negative and positive values. In the measurement of the mean amplitude values, the minimum and maximum times of analysis considered were 1 and 2 seconds respectively. Mean amplitudes were categorized as normal (188-1485 μV)29, increased, or reduced.

4. During the examination of the thyroarytenoid muscle, one of the voluntary activity values was obtained by pushing the device's button that quantifies the intensity of the sound generated by electrical activity at the minimum volume level, i.e., without sound. This file was used to measure the latency between the onset of electrical activity acquired on channel 1 and the appearance of the sound uttered by the subject acquired on the microphone connected to channel 2 on the device. Latency times were measured from the onset of increased electrical activity from the baseline on channel 1 to the onset of activity on the voice channel (channel 2). Latency times were deemed normal when equal to or under 408 ms29 or increased when above the threshold.

Based on the findings related to electrical activity at rest and during voluntary activation, EMG examination of each laryngeal muscle may indicate the following:

1. Absence of electrophysiological alterations: normal insertion activity, absence of spontaneous activity, normal amplitude and duration, normal latency time.
2. Neurogenic lesion: normal, increased, or reduced insertion activity; normal latency time. Acute involvement was characterized when fibrillation-type spontaneous activity or positive wave and rarefaction were found. The presence of rarefaction and absence of spontaneous activity were interpreted as neuropathic. Chronic involvement was identified when fibrillation-type spontaneous activity or positive waves were not found, fasciculation or high frequency discharge was present or absent, and rarefaction and waves with increased duration and amplitude were found. The presence of rarefaction and absence of spontaneous activity, along with normal wave duration and amplitude, was characterized as chronic neurogenic injury with central involvement.

3. Myopathic lesion: normal insertion activity, reduced MUAP amplitude and duration.

4. Inconclusive: findings did not fit any of the categories described above.

The electrical activity of the thyroarytenoid and LCA muscles was analyzed to assess recurrent laryngeal nerve function, whereas the cricothyroid was used to look into the superior laryngeal nerve bilaterally. The thyroarytenoid and the LCA are innervated by the recurrent laryngeal nerve, but both must be analyzed as patients may have selective cases of paresis - e.g.: thyroarytenoid paresis - which are difficult to characterize through laryngoscopy and require EMG examination to confirm the diagnosis.

The subjects were provided with clarification on the research and on the tests they signed up to undergo and signed a free informed consent form.

RESULTS

Nine of the 39 patients submitted to EMG of the larynx did not have their LCA muscles examined due to technical hazards in locating their muscles or because they asked for a shorter course of tests. The thyroarytenoid muscle was normal in all such patients. In three patients the cricothyroid muscle was not found - two on the right side and one on the left side. The difficulty identifying the three cricothyroid muscles may be explained by the approached picked to perform the examination: the same puncture was used to look at the cricothyroid and thyroarytenoid muscles, and the puncture may have been performed medially to the cricothyroid. In one patient, only the cricothyroid muscles were examined and in another patient only the muscles on the left side were observed. All other patients (25) had all the described muscles examined.

The mean latencies between electrophysiological activity and start of phonation in the right and left thyroarytenoid muscles was 230.9 ms and 194.4 ms respectively; MUAP durations in the left (LCT) and right (RCT) cricothyroid muscles, left (LTA) and right (RTA) thyroarytenoid muscles, left (LLCA) and right (RLCA) lateral cricoarytenoid muscles were 4.1 ms, 3.5 ms, 4.7 ms, 4.6 ms, 4.8 ms, and 5.0 ms respectively.

The mean MUAP amplitude on muscles LCT, RCT, LTA, RTA, LLCA, and RLCA were 626.7 μV, 703.9 μV, 688.8 μV, 725.7 μV, 662.0 μV, and 679.9 μV respectively.

The analysis of EMG results shows that wave morphology was normal and none had pathologic spontaneous activity or rarefaction. MUAP duration was found to be within normal range, as were latencies between electrophysiological activation and phonation (Table 1). Mean MUAP amplitude was within normal range for all tested muscles (Table 2).

Therefore, no functional electrophysiological alterations were seen in the superior and recurrent laryngeal nerves of the patients with incomplete glottic closure and dysphonia.

DISCUSSION

EMG is an extremely useful tool in diagnosing neurogenic diseases, diseases of the neuromuscular junction, and myopathies, and has been broadly applied in large skeletal muscles. In patients suspected for neuromuscular junction involvement and myopathies, EMG can be done in accessible muscles such as the levator palpebrae superioris, the orbicularis oculi, the biceps, and the quadriceps.

In laryngeal mobility disorders, EMG allows the examiner to locate the lesion, define which nerves were involved, and draw out prognostic parameters.

EMG of the larynx has its particularities, and requires knowledge on the anatomy of the structures of the larynx, more specifically the location of the intrinsic muscles and the pathophysiology of the diseases that may hamper the function of the larynx as the protective barrier for the lower airways and the organ responsible for phonation.

The intrinsic laryngeal muscles are small and located close to each other and other muscles in the neck area, as the extrinsic laryngeal muscles. The larynx has antagonist muscles which, by means of the motion of the arytenoids - mainly its rotation on the cricoid cartilage, lead to the abduction of the vocal folds and, consequently, the free inflow and outflow of air, or adduction, during swallowing or phonation.

The innervation of the antagonist muscles is done by the recurrent laryngeal nerve, a branch of the vagus nerve, whose motor neurons are located in the nucleus ambiguous, bilaterally in the brainstem. The recurrent laryngeal nerve stretches throughout the neck, and is thus susceptible to trauma and compressive injuries, and lesions subsequent to thyroidectomy.
### Table 1. Latency between electrophysiological activity and onset of phonation (ms) and MUAP duration in the examined muscles (ms).

| Patient | Latency | Duration |
|---------|---------|----------|
|         | LTA     | RTA      | LCT     | RCT     | LTA  | RTA  | LLCA | RLCA  |
| 1       | 404     | 350      | 4.7     | 3.8     | 4.2  | 5.3  | -    | -     |
| 2       | 274     | Ni       | 4.2     | 3.7     | 4.3  | 5.1  | 5.7  | 4.6   |
| 3       | 145     | 195      | 3.7     | 3.2     | 4.7  | 4.5  | -    | -     |
| 4       | Ni      | Ni       | 2.9     | 5.3     | 4.2  | 4.6  | -    | -     |
| 5       | 204     | 250      | 2.9     | 3.5     | 3.7  | 4.2  | 3.3  | 3.6   |
| 6       | 178     | 78       | 4.9     | 5.7     | 4.1  | 3.3  | 5.3  | 6.5   |
| 7       | 190     | 178      | 2.6     | 4.7     | 5.3  | 3.4  | 6.8  | 6.1   |
| 8       | 125     | 115      | 2.5     | 4.2     | 3.4  | 3.3  | 4.6  | 2.8   |
| 9       | 274     | 129      | 4.0     | 4.2     | 3.9  | 4.3  | 5.0  | 4.6   |
| 10      | 321     | 418      | 3.5     | 3.8     | 5.2  | 4.5  | -    | -     |
| 11      | Ni      | Ni       | 4.8     | 4.2     | 3.7  | 4.3  | 5.2  | 5.7   |
| 12      | 123     | 42       | 3.5     | 3.3     | 4.3  | 2.7  | 3.2  | 3.5   |
| 13      | 236     | Ni       | 2.7     | 3.2     | 3.3  | 3.4  | -    | -     |
| 14      | 197     | Ni       | 2.7     | -       | 5.6  | -    | 5.7  | -     |
| 15      | 379     | 261      | 4.4     | 4.9     | 5.7  | 4.8  | 2.8  | 4.7   |
| 16      | 123     | 144      | 3.9     | 3.7     | 3.3  | 5.2  | -    | -     |
| 17      | 311     | 253      | 2.4     | 5.2     | 4.2  | 3.1  | 4.7  | 4.8   |
| 18      | 231     | 140      | 4.7     | 4.4     | 2.7  | 5.0  | -    | -     |
| 19      | 220     | 240      | 5.9     | 6.5     | 5.9  | 5.8  | 3.1  | 3.7   |
| 20      | 350     | 325      | 3.3     | 3.8     | 3.4  | 5.4  | 5.9  | 6.7   |
| 21      | 275     | 120      | 3.6     | 3.8     | 3.2  | 4.8  | 4.6  | 4.8   |
| 22      | 196     | 239      | 5.6     | 6.6     | 5.9  | 5.7  | 5.4  | 5.5   |
| 23      | 63      | 91       | 3.9     | 3.6     | 5.4  | 4.8  | 5.8  | 5.9   |
| 24      | Ni      | Ni       | 5.2     | 3.5     | 5.8  | 4.6  | 3.3  | 5.3   |
| 25      | Ni      | Ni       | 3.7     | 4.2     | -    | -    | -    | -     |
| 26      | 197     | 78       | 5.5     | 3.8     | 6.5  | 5.2  | -    | -     |
| 27      | 312     | 335      | 4.9     | 3.4     | 4.7  | 3.8  | 4.7  | 5.3   |
| 28      | 95      | 233      | 5.9     | 4.8     | 4.9  | 4.2  | 5.2  | 4.2   |
| 29      | 179     | 124      | 4.9     | 3.5     | 4.6  | 3.9  | 4.6  | 5.3   |
| 30      | 215     | Ni       | 6.5     | 4.2     | 4.0  | 6.4  | 5.9  | 4.7   |
| 31      | 153     | 115      | 3.0     | -       | 4.7  | 4.0  | 4.1  | 6.1   |
| 32      | 417     | 278      | 4.3     | 5.4     | 5.7  | 4.7  | 6.8  | 5.3   |
| 33      | 279     | 82       | 3.9     | 3.3     | 4.4  | 5.8  | 5.7  | 4.5   |
| 34      | 346     | 124      | 4.2     | 3.2     | 5.0  | 4.3  | 4.6  | 5.7   |
| 35      | 225     | Ni       | 3.0     | 3.0     | 4.7  | 4.0  | 5.3  | 5.2   |
| 36      | 283     | 248      | 5.9     | 6.4     | 5.6  | 5.7  | 3.4  | 4.8   |
| 37      | 154     | 226      | 2.7     | 3.2     | 5.1  | 2.9  | 4.5  | 4.8   |
| 38      | Ni      | 158      | 4.6     | 5.1     | 6.9  | 5.7  | 2.8  | 5.3   |
| 39      | 176     | 264      | 5.1     | 3.8     | 6.1  | 6.8  | 5.7  | 4.4   |
| mean    | 230.9   | 194.4    | 4.1     | 3.5     | 4.7  | 4.6  | 4.8  | 5.0   |
| median  | 217.5   | 186.5    | 4.0     | 3.8     | 4.7  | 4.6  | 4.9  | 4.8   |
| minimum | 63      | 42       | 2.4     | 3.0     | 2.7  | 2.7  | 2.8  | 2.8   |
| maximum | 417     | 418      | 6.5     | 6.6     | 6.9  | 6.8  | 6.8  | 6.7   |

Muscles: left thyroarytenoid (LTA); right thyroarytenoid (RTA); left cricothyroid (LCT); right cricothyroid (RCT); left lateral cricoarytenoid (LLCA); right lateral cricoarytenoid (RLCA); NI: not included.
Table 2. Mean MUAP amplitude (µV).

| Patient | LCT  | RCT  | LTA  | RTA  | LLCA | RLCA |
|---------|------|------|------|------|------|------|
| 1       | 491.2| 948.7| 1152.1| 1059.8|      |      |
| 2       | 416.3| 531.1| 996.5| 1104.8| 798.3| 854.7|
| 3       | 168.0| 425.6| 1399.2| 547.9|      |      |
| 4       | 1146.1| 402.4| 599.7| 363.5|      |      |
| 5       | 474.2| 410.0| 795.4| 331.7| 543.7| 438.6|
| 6       | 795.8| 632.7| 674.8| 685.4| 411.3| 1144.4|
| 7       | 355.4| 739.6| 626.7| 475.4| 845.0| 600.1|
| 8       | 765.4| 586.1| 1319.9| 937.1| 677.5| 730.0|
| 9       | 291.4| 896.9| 548.3| 799.2| 509.5| 691.7|
| 10      | 391.3| 620.7| 612.8| 505.3|      |      |
| 11      | 573.8| 699.3| 575.7| 546.6| 361.3| 377.5|
| 12      | 668.2| 367.5| 628.2| 400.5| 354.2| 367.2|
| 13      | 766.5| 888.9| 753.7| 547.9|      |      |
| 14      | 1128.8| NI   | 419.3| NI   | 1309.9| NI   |
| 15      | 225.2| 419.2| 326.5| 600.8| 409.1| 548.9|
| 16      | NI   | 1276.8| 864.3| 1294.9|      |      |
| 17      | 789.1| 872.9| 602.5| 827.6| 454.4| 841.7|
| 18      | 628.9| 605.3| 603.5| 595.1|      |      |
| 19      | 713.6| 459.1| 744.3| 467.3| 1203.5| 589.0|
| 20      | 376.6| 566.4| 373.1| 809.9| 487.9| 568.9|
| 21      | 667.7| 413.3| 415.3| 500.5| 787.5| 865.8|
| 22      | 385.7| 907.5| 513.4| 831.6| 493.9| 376.5|
| 23      | 385.1| 767.3| 410.5| 577.5| 596.0| 394.1|
| 24      | 958.5| 453.4| 347.6| 491.1| 453.3| 476.9|
| 25      | 626.1| 636.7| NI   | NI   |      |      |
| 26      | 500.7| 1142.5| 682.9| 638.6|      |      |
| 27      | 547.5| 424.0| 356.5| 580.0| 1484.7| 1013.5|
| 28      | 762.0| 1370.0| 534.7| 609.1| 657.3| 754.4|
| 29      | 515.4| 1272.7| 795.1| 1430.6| 456.7| 643.2|
| 30      | 1143.1| 759.1| 1399.2| 1016.6| 857.6| 753.0|
| 31      | 933.9| NI   | 298.1| 421.4| 487.4| 435.9|
| 32      | 607.9| 537.1| 851.4| 788.5| 787.1| 653.3|
| 33      | 197.7| 498.2| 789.3| 1281.0| 710.9| 1432.6|
| 34      | 758.3| 678.9| 546.7| 678.9| 489.5| 578.9|
| 35      | 978.4| 1105.5| 678.1| 864.8| 534.9| 650.2|
| 36      | 537.8| 789.2| 567.7| 675.3| 748.2| 890.2|
| 37      | 764.8| 598.3| 1120.5| 978.9| 734.7| 756.8|
| 38      | 890.4| 796.7| 540.0| 749.0| 557.8| 638.9|
| 39      | 489.5| 546.3| 758.3| 837.5| 655.8| 534.8|

Mean 626.7 703.9 688.8 725.7 662.0 679.9
Median 617.0 632.7 619.8 675.3 577.9 641.1
Minimum 168.0 367.5 298.1 331.7 354.2 534.8

**muscles:** left thyroarytenoid (LTA); right thyroarytenoid (RTA); left cricothyroid (LCT); right cricothyroid (RCT); left lateral cricoarytenoid (LLCA); right lateral cricoarytenoid (RLCA). NI: not included.
Laryngeal mobility disorders bear significant clinical consequences and repercussions upon phonation when occurring unilaterally, and upon breathing when involvement is bilateral.

The vagus nerve, through its superior laryngeal branch, innervates the cricothyroid muscle which, by the forward rocking of the thyroid cartilage and backward rocking of the cricoid cartilage, elongates and tenses the vocal folds, making it possible for people to utter high frequency sounds which are critical in singing. Injuries to the superior laryngeal nerve - paralysis or paresis - do not lead to laryngeal immobility, but only to indirect signs suggestive of involvement. The diagnosis of muscle paralysis is done through EMG. Thus, in order to assess the recurrent and superior laryngeal nerves, the electrophysiological activity of at least two laryngeal intrinsic muscles should be considered: the cricothyroid, evaluated in high frequency utterances, and the thyroarytenoid through phonation at low and high frequencies.

The recurrent laryngeal nerve branches out after it penetrates the larynx, and usually its first branch moves toward the posterior cricoarytenoid abductor muscle. The second branch reaches the interarytenoid muscle’s both sides, and follows on to the LCA to finally reach the terminal branch of the thyroarytenoid. Albeit rate, selective paresis of its branches may occur and diagnosis can only be performed using EMG. In this case, knowledge of clinical alterations may require the assessment of other intrinsic muscles which are harder to access and, thus, are not routinely examined.

Incomplete closure of the vocal folds leads to dysphonia, varying degrees of breathiness, reduced intensity, compromised voice efficiency, and exhaustion when speaking. Some alterations in the configuration of the larynx during phonation may be consequent, among other factors, to neurological involvement. Thus, spindle-shaped clefts have been related to paresis and paralysis of the recurrent and superior laryngeal nerves of 16% of their patients. However, other findings were observed through laryngoscopy, such as arytenoid rotation and hypermobility, which resulted in indication to perform EMG. Our study was different in this regard to that done by Koufman et al., as in our sample we had no patients with altered laryngeal mobility. Therefore, more studies are required to shed light on the factors connected to glottic insufficiency.

CONCLUSION

Electrophysiological assessment of the superior and recurrent laryngeal nerves failed to detect disorders in patients with dysphonia and incomplete glottic closure seen through video laryngostroboscopy.

REFERENCES

1. Benninger MS, Crumley RL, Ford CN, Gould WJ, Hanson DG, Ossof RH, et al. Evaluation and treatment of the unilateral paralyzed vocal fold. Otolaryngol Head Neck Surg. 1994;111(4):497-508.
2. Bevan K, Griffiths MV, Morgan MH. Cricoarytenoid muscle paralysis: its recognition and diagnosis. J Laryngol Otol. 1980;103(2):191-5.
3. Dedo HH. The paralyzed larynx: an electromyographic study in dogs and humans. Laryngoscope. 1970;89(10):1455-517.
4. Dedo HH, Hall WN. Electrodes in laryngeal electromyography. Re-liability comparison. Ann Otol Rhinol Laryngol. 1969;78(1):172-80.
5. Dursun G, Sataloff RT, Spiegel JR, Mandel S, Heuer RJ, Rosen DC. Superior nerve paresis and paralysis. J Voice. 1996;10(2):206-11.
6. Eckley CA, Sataloff RT, Hawkshaw M, Spiegel JR, Mandel S. Voice range in superior laryngeal nerve paresis and paralysis. J Voice. 1998;12(3):340-8.
7. Iroto I, Hirano M, Tomita H. Electromyographic investigation of human vocal cord paralysis. Ann Otol Rhinol Laryngol. 1968;77(2):296-304.
8. Koufman JA, Walker FO, Joharii GM. The cricothyroid muscle does not influence vocal fold position in laryngeal paralysis. Laryngoscope. 1995;105(4 Pt 1):D136-72.
9. Min YB, Finnegan EM, Hoffman HT, Luschei ES, McCulloch TM. A preliminary study of the prognostic role of electromyography in laryngeal paralysis. Otolaryngol Head Neck Surg. 1994;111(7):770-5.
10. Parans SM, Satya-Murti S. Predictive value of laryngeal electromyography in patients with vocal cord paralysis of neurogenic origin. Laryngoscope. 1989;99(11):1523-6.
11. Rodriguez AA, Myers BR, Ford CN. Laryngeal electromyography in the diagnosis of laryngeal nerve injuries. Arch Phys Med Rehabil. 1990;71(8):587-90.
12. Rontal E, Rontal M, Silverman B, Kileny PR. The clinical differentiation between vocal cord paralysis and vocal cord fixation using electromyography. Laryngoscope. 1993;103(2):133-7.
13. Simpson DM, Sternman D, Graves-Wright J, Sanders I. Vocal cord paralysis: clinical and electrophysiologic features. Muscle Nerve. 1993;16(9):952-7.
14. Thunfart WF. Electrodiagnosis of laryngeal nerve disorders. Ear Nose Throat J. 1988;67(5):380-4.
15. Woodson GE, Murry MP, Schweizer V, Hengestag A, Chen N, Yeung D. Unilateral cricothyroid contraction and glottic configuration. J Voice. 1998;12(3):355-9.
16. Yin SS, Qiu WW, Stucker FJ. Value of electromyography in differential diagnosis of laryngeal joint injuries after intubation. Ann Otol Rhinol Laryngol. 1996;105(6):446-51.
17. Lindestad PA. Electromyographic and laryngoscopic studies of normal and disturbed voice function. Stockholm: Huddinge University Hospital; 1994. 48p.
18. Dejonckere P, Hamoir M. Etiology of neurogenic laryngeal lesions diagnosed by electromyography. Acta Otorhinolaryngol Belg. 1980;34(3):285-99.
19. Crumley RL. Unilateral recurrent laryngeal nerve paralysis. J Voice. 1994;8(1):79-83.
20. Kotby MN, Haugen LK. Clinical application of electromyography in vocal fold mobility disorders. Acta Otolaryngol. 1970;70(5):428-37.
21. Eckel HE, Wittenkindt C, Klaussmann JP, Schroeder U, Sittel C. Management of bilateral arytenoid cartilage fixation versus recurrent laryngeal nerve paralysis. Ann Otol Rhinol Laryngol. 2003;112(2):103-8.
22. Munin MC, Rosen CA, Zullo T. Utility of laryngeal electromyography in predicting recovery after vocal fold paralysis. Arch Phys Med Rehabil. 2003;84(6):1150-5.
23. Koufman JA, Postma GN, Whang CS, Rees CJ, Amin MR, Belafsky PC, et al. Diagnostic laryngeal electromyography: The Wake Forest experience 1995-1999. Otolaryngol Head Neck Surg. 2001;124(6):603-6.
24. De Biase N, Pontes P, Nobrega JAM, De Biase S. Paralisia do músculo cricaritenóideo lateral: relato de caso como contribuição ao estudo da anatomia funcional da laringe. Rev Bras Otorrinolaringol. 2003;69(1):121-6.
25. Watson BC, Schaefer SD, Freeman FJ, Dembowski J, Kondraske G, Roark R. Laryngeal electromyographic activity in adductor and abductor spasmodic dysphonia. J Speech Hear Res. 1991;34(5):473-82.
26. Knutsson E, Martinsson A, Martinsson B. The normal electromyogram in human vocal muscles. Acta Otolaryngol. 1969;68(6):526-36.
27. Hillel ND. The study of laryngeal muscle activity in normal human subjects and in patients with laryngeal dystonia using multiple fine-wire electromyography. Laryngoscope. 2001;111(4 Pt 2 Suppl 97):1-47.
28. Kimaid PA, Crespo AN, Quagliato EM, Wolf A, Viana MA, Resende LA. Laryngeal electromyography: contribution to vocal fold immobility diagnosis. Electromyogr Clin Neurophysiol. 2004;44(6):571-4.
29. Sanders I, Wu BL, Mu I, Li Y, Biller HF. The innervation of the human larynx. Arch Otolaryngol Head Neck Surg. 1993;119(9):934-9.
30. Koufman JA, Postma GN, Cummins MM, Blalock PD. Vocal fold paresis. Otolaryngol Head Neck Surg. 2000;122(4):537-41.
31. Aronson AE, Bless DM. Voice Disorders of Structural Origin. In: Aronson AE, Bless DM, editors. Clinical Voice Disorders. 4th ed. New York: Thieme Med Pub; 2009. p.24-38.
32. Koufman JA. Bowing of the Vocal Cords [Site na Internet].Disponível em: http://www.bgsm.edu/voice/bowing.html. Acessado em 20 de abril de 2001.