Video-assisted thoracoscopy for the diagnosis of diffuse parenchymal lung disease*

A videotoracoscopia no diagnóstico das doenças difusas do parênquima pulmonar

Renato Tadao Ishie, João José de Deus Cardoso, Rafael José Silveira, Lucas Stocco

Abstract

Objective: To evaluate the role of video-assisted thoracoscopy in the diagnosis of diffuse parenchymal lung diseases. Methods: The medical charts of patients suspected of having diffuse parenchymal lung disease were retrospectively reviewed, as were the results of the anatomopathological examination of lung biopsy specimens collected through video-assisted thoracoscopy. Results: Of the 48 patients included in the study, 25 (52.08%) were female and 23 (47.92%) were male. The mean age was 58.77 years (range, 20-76 years). A total of 54 biopsy fragments were submitted to anatomopathological examination: 24 (44.44%) from the lingula; 10 (18.52%) from the left lower lobe; 7 (12.96%) from the right middle lobe; 6 (11.11%) from the right lower lobe; 5 (9.26%) from the left upper lobe; and 2 (3.71%) from the right upper lobe. The mean duration of thoracic drainage was 2.2 days. Adverse events included conversion to thoracotomy, in 2 patients (4.17%), and residual pneumothorax, in 1 (2.08%). The definitive diagnosis was made in 46 patients (95.83%), with idiopathic interstitial pneumonia being the predominant diagnosis (in 54.18%). The most common diagnoses were usual interstitial pneumonia (in 29.27%), nonspecific interstitial pneumonia (in 16.67%) and hypersensitivity pneumonia (in 12.50%). Conclusions: Lung biopsy through video-assisted thoracoscopy is a safe, effective and viable procedure for the diagnosis of diffuse parenchymal lung diseases.

Keywords: Lung diseases, interstitial; Thoracoscopy; Diagnosis.

Resumo

Objetivo: Analisar o papel da videotoracoscopia no diagnóstico das doenças difusas do parênquima pulmonar. Métodos: Os prontuários de pacientes com suspeita de doenças difusas do parênquima pulmonar e os resultados do exame anatomopatológico das amostras de biópsia pulmonar por videotoracoscopia foram analisados retrospectivamente. Resultados: Dos 48 pacientes incluídos no estudo, 25 (52,08%) eram do sexo feminino, e 23 (47,92%) eram do sexo masculino. A idade média foi de 58,77 anos, variando entre 20 e 76 anos. Foi realizado o exame anatomopatológico de 54 fragmentos de biópsia pulmonar: 24 (44,44%) da lingula; 10 (18,52%) do lobo inferior esquerdo; 7 (12,96%) do lobo superior esquerdo; 6 (11,11%) do lobo inferior direito; 5 (9,26%) do lobo superior direito; e 2 (3,71%) do lobo superior direito. O tempo médio de drenagem torácica foi de 2,2 dias. Como eventos adversos, houve conversão para toracotomia em 2 pacientes (4,17%) e pneumotórax residual em 1 (2,08%). O diagnóstico definitivo foi obtido em 46 (95,83%) casos, com predomínio das pneumonias intersticiais idiopáticas (54,18%). Os diagnósticos mais frequentes foram pneumonia intersticial usual (29,27%), pneumonia intersticial não-específica (16,67%) e pneumonia por hipersensibilidade (12,50%). Conclusões: A videotoracoscopia com biópsia pulmonar é um procedimento eficaz, seguro e viável para o diagnóstico das doenças difusas do parênquima pulmonar.

Descritores: Doenças pulmonares intersticiais; Toracoscopia; diagnóstico.

Introduction

Diffuse parenchymal lung diseases (DPLDs) constitute a heterogeneous group of lung diseases, including more than two hundred different interstitial diseases, characterized by varying degrees of inflammation and fibrosis. These non-neoplastic disorders primarily affect the lung interstitium, although the alveolar space, bronchioles and pulmonary vessels can also be affected. In addition to the histological aspects, these disorders present similar

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The process of diagnosing a DPLD is dynamic. The diagnostic reasoning is based on the joint analysis of clinical, radiological and anatomopathological aspects.\(^{[1]}\)

The radiological presentation of a DPLD typically varies. Routine chest X-rays present low sensitivity and specificity,\(^{[4]}\) typically identifying the presence of pulmonary disease and indicating the need for additional tests that are more complex. A common presentation is the development of honeycomb cystic areas.

Among imaging tests, an HRCT scan of the chest is the best suited to the evaluation of a DPLD. Such scans can be used not only to identify the presence of disease but also to evaluate its extent and characterize its pattern. In addition, an HRCT scan of the chest helps reduce the number of differential diagnoses and define the site from which the biopsy specimen is to be taken, as well as allowing the clinical course of the disease and the response to therapy to be evaluated.\(^{[5]}\) However, HRCT lacks diagnostic specificity and therefore, in most DPLD patients, does not preclude the need for histological confirmation,\(^{[6]}\) as shown in Figure 2.

Pulmonary function tests are often recommended. These tests can estimate severity and prognosis, as well as monitoring response to therapy and disease progression.\(^{[7]}\) The recommended tests include spirometry, DLCO determination and evaluation of the degree of reduction in oxygenation during exercise.

**Figure 1** - Classification of diffuse parenchymal lung diseases.\(^{[1]}\) DPLD: diffuse parenchymal lung disease.
through the use of blood gas analysis or pulse oximetry. Usually, DPLDs present a characteristic functional pattern, with restrictive ventilatory failure and reduced diffusing capacity.

Bronchoalveolar lavage is a test that allows the analysis of cells in the airway and alveolar space, as well as of soluble substances from the extracellular mucus layer. Bronchoalveolar lavage is a useful technique for the investigation of pneumoconiosis, opportunistic infections, suspected malignancy, some hematologic diseases and alveolar proteinosis, as well as diseases related to lung transplantation or drug-induced diseases.

Frequently, the definitive diagnosis of DPLD can be established only through anatomopathological examination of the material obtained by lung biopsy. In addition to diagnostic confirmation, this procedure provides information regarding disease activity, disease progression and response to therapy. The options for lung biopsy include bronchoscopy with transbronchial biopsy, open lung biopsy and lung biopsy through video-assisted thoracoscopy.

Bronchoscopy with transbronchial biopsy is useful in cases in which the disease presents peribronchial, peribronchiolar or diffuse distribution. One limitation of this procedure is the small quantity of lung tissue obtained in the biopsy, which is why it is not recommended for the investigation of idiopathic interstitial pneumonia. In addition, its accuracy in the diagnosis of DPLD in immunocompetent patients is only 7-37%.

The gold standard for the diagnosis of DPLD is surgical lung biopsy, which should be used whenever it is not possible to establish a definitive diagnosis based on the available clinical and radiological data. It can be performed as an open procedure or through video-assisted thoracoscopy.

Open lung biopsy has a high diagnostic yield (92%), as well as low rates of morbidity and mortality (2.5% and 0.3%, respectively). The most common technique is minimal inframammary thoracotomy, through which lung biopsy is performed, using either manual or mechanical sutures.

Video-assisted thoracoscopy is considered a minimally invasive technique. It provides excellent visualization of the intrathoracic structures and allows the collection of a greater number of lung samples, when necessary. Since it is a less invasive procedure, video-assisted thoracoscopy has come to be used as the principal means of diagnosing DPLD, and the number of surgical
lung biopsies has increased. However, its use must be evaluated in terms of safety and diagnostic resolution. Therefore, the objective of the present study was to analyze the role of this method, which is currently widely used, in the diagnosis of DPLD.

**Methods**

The medical charts of patients being monitored in order to diagnose DPLD were evaluated, as were the results of the anatomopathological examination of lung biopsy specimens collected through video-assisted thoracoscopy. All of the eligible patients had been monitored in the Department of Thoracic Surgery of the Nereu Ramos Hospital in the city of Florianópolis, located in the state of Santa Catarina, between July of 1999 and July of 2007. The inclusion criteria were as follows: being under outpatient follow-up treatment in order to diagnose DPLD; not having received a diagnosis by noninvasive evaluation; and not having received a histopathological diagnosis in the transbronchial biopsy, when performed.

The exclusion criteria were as follows: requiring mechanical ventilation in an intensive care unit and being oxygen-dependent.

There were 48 patients who met the criteria adopted, and the charts of those patients were therefore selected for study.

Data regarding gender and age of the patients were analyzed. Age (in years) was defined as that at the time of lung biopsy. The distribution of biopsy sites and the anatomopathological diagnoses obtained were also analyzed. Regarding the surgical procedure, the variables studied were surgical complications, duration of thoracic drainage in the postoperative period and postoperative complications.

**Results**

A total of 48 patients met the study criteria and were included. Of those, 25 (52.08%) were female and 23 (47.92%) were male. Patient ages ranged from 20 to 76 years (mean, 58.77 years).

A total of 54 lung biopsy samples were sent for anatomopathological analysis: 39 (72.22%) from the left lung and 15 (27.28%) from the right lung. Table 1 shows the biopsy site distribution.

The mean duration of thoracic drainage in the postoperative period was 2.2 days (range, 1-4 days).

Regarding intraoperative complications, 2 patients (4.17%) required 4-5 cm auxiliary incisions. Only 1 patient (2.08%) presented a postoperative complication (residual pneumothorax after chest tube removal).

The anatomopathological results of the video-assisted thoracoscopic lung biopsies are listed in Table 2.

<table>
<thead>
<tr>
<th>Table 1 - Lung biopsy site distribution.</th>
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<tbody>
<tr>
<td>Biopsy site</td>
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</tr>
<tr>
<td>Lingula</td>
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<td>Left lower lobe</td>
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<td>Middle lobe</td>
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<td>Right lower lobe</td>
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<th>Table 2 - Distribution of the diagnoses and findings made by video-assisted thoracoscopic biopsy.</th>
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<tr>
<td>Diagnosis/finding</td>
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<td>---------------------------------------------</td>
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<tr>
<td>DPLDs of known etiology</td>
</tr>
<tr>
<td>Hypersensitivity pneumonia</td>
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<tr>
<td>Pulmonary tuberculosis</td>
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<tr>
<td>Paracoccidioidomycosis</td>
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<tr>
<td>Amiodarone-induced interstitial lung injury</td>
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<tr>
<td>Idiopathic interstitial pneumonia</td>
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<td>Usual interstitial pneumonia</td>
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<tr>
<td>Nonspecific interstitial pneumonia</td>
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<tr>
<td>Lymphocytic interstitial pneumonia</td>
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<tr>
<td>Cryptogenic organizing pneumonia</td>
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<tr>
<td>Granulomatous DPLDs</td>
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<td>Wegener’s granulomatosis</td>
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<tr>
<td>Sarcoidosis</td>
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<td>Other forms of DPLD</td>
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<td>Eosinophilic pneumonia</td>
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<td>Eosinophilic granuloma</td>
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<td>Papillary adenocarcinoma</td>
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<td>Alveolar proteinosis</td>
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<td>Airway-centered interstitial fibrosis</td>
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<td>Other findings</td>
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<tr>
<td>Lung honeycombing</td>
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<td>Normal lung tissue</td>
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<td>Total</td>
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DPLD: diffuse parenchymal lung diseases.
Discussion

The DPLDs constitute a heterogeneous group of lung diseases characterized by varying degrees of inflammation and fibrosis. In some DPLDs, significant morbidity and unfavorable evolution, comparable to those of neoplastic diseases, are seen. Therefore, an efficient and safe method for the diagnostic confirmation of DPLD is needed. Currently, lung biopsy through video-assisted thoracoscopy is widely used for this purpose.

The objective of the present study was to analyze the role of this technique in the diagnosis of DPLD. To that end, the medical charts of 48 patients who underwent lung biopsy through video-assisted thoracoscopy were analyzed.

In our study sample, there was a slight predominance of females (52.08%) over males (47.92%). The same was found in another study, in which 55.13% of the patients were female and 44.87% were male. However, other authors have found a slight predominance of males. Most of our patients (89.59%) were over 41 years of age (mean, 58.77 years). This finding is similar to those reported in the studies cited above.

A total of 54 lung biopsy samples were obtained from 48 patients. The samples were representative and sufficient to establish the diagnosis in 46 patients (95.83%). In 1 patient (2.08%), histopathological confirmation was not possible due to extensive honeycombing with distortion of the lung architecture. In another case, the histopathological analysis revealed normal lung parenchyma.

In 42 patients (87.50%), a single biopsy was performed. In 6 (12.50%), multiple biopsies, of 2 samples each, were performed.

In the multiple biopsies analyzed, the biopsy sites were the right upper lobe/lingula in 2 cases, the right middle lobe/right upper lobe in 1 case and the right middle lobe/right lower lobe in 1 case. In the remaining 2 cases, it was not possible to define the site of the second biopsy due to lack of information in the medical charts. In the multiple biopsies, the definitive diagnosis was made in 5 cases. The remaining case is the one in which the definitive diagnosis was not possible due to honeycombing. When the anatomicopathological results of the two samples from the same patient were compared, the diagnoses were concordant.

In the present study, a definitive diagnosis was established in all 42 cases in which a single biopsy was performed. When multiple biopsies were performed, the results of the two samples were the same. This is in agreement with the findings of another study, in which it was concluded that a single sample obtained from a region pre-selected using radiology is sufficient for the diagnosis. Yet another study, although not demonstrating greater diagnostic efficacy with the use of multiple biopsies, recommended that such biopsies be performed whenever possible. Various authors have stated that multiple biopsies increase the probability of establishing the definitive diagnosis.

The justification for performing multiple biopsies is probably the difficulty in choosing the segment to be biopsied, since extensive areas of fibrosis make it difficult to identify the specific characteristics of the underlying disease.

The data obtained in the present study, although limited, show that a single sample is sufficient for the diagnosis. Nevertheless, pre-planning, with the aid of HRCT, is necessary for choosing the appropriate site to be biopsied. In addition, multiple biopsies would probably increase the diagnostic accuracy of the test, although the high cost of endoscopy can be a limiting factor for this option in Brazil.

The most common biopsy site was the lingula, accounting for nearly half of the samples obtained. In the left lung, the most common site was the middle lobe. The choice of these sites for biopsy is controversial. Some authors recommend that these sites be avoided since they are common sites of nonspecific infectious processes, inflammation, cicatrization and vascular congestion. This would affect the quality of the sample, which would present more fibrosis and vascular alterations than those collected from other pulmonary areas. However, various studies have demonstrated that biopsy samples obtained from the lingula or from the middle lobe have the same diagnostic yield as those obtained from other lung segments.

The samples obtained from the middle lobe were sufficient for the diagnosis, except for that obtained in 1 case, in which there was pronounced honeycombing. It is likely that, in that case, the lung presented an advanced degree of diffuse fibrosis, since multiple biopsies...
Sarcoidosis was found in only 1 case (2.08%). In contrast, in the literature, sarcoidosis is described as a common DPLD. In a study involving 3,152 patients, sarcoidosis was the most common disease (in 33.72% of the cases), followed by idiopathic pulmonary fibrosis (in 27.41% of the cases). This is due to the fact that sarcoidosis, due to its characteristic peribronchial or peribronchiolar distribution, is preferably diagnosed by transbronchial biopsy. Therefore, in suspected cases of sarcoidosis, surgical lung biopsy is used only when the diagnosis cannot be made by transbronchial biopsy.

In 12.08% of our cases, the histological finding was normal lung parenchyma. In the subsequent diagnostic investigation of this patient, gastroesophageal reflux disease was diagnosed. This might have been due to an atypical clinical presentation, the differential diagnosis being made only after the surgical lung biopsy.

Many studies have compared video-assisted thoracoscopy and open lung biopsy in the diagnosis of DPLD. These techniques have been compared in terms of diagnostic efficacy and safety in surgical lung biopsy.

As discussed above, video-assisted thoracoscopy provided adequate lung tissue samples with high diagnostic efficacy. The definitive diagnosis was made in 95.83% of the cases. This finding is in agreement with those of another study, in which the diagnosis was established in 98.39% of the cases. Open lung biopsy produces findings similar to those of video-assisted thoracoscopy.

In the patients studied, there was a higher prevalence of diseases that belong to the idiopathic interstitial pneumonia group (54.18%), especially usual interstitial pneumonia (29.17%) and nonspecific interstitial pneumonia (16.67%). Hypersensitivity pneumonia was the third most common (12.50%).

One group of authors also described a higher prevalence of idiopathic interstitial pneumonia, accounting for 38.58% of the 744 cases studied. Among those, usual interstitial pneumonia was also the most common. However, hypersensitivity pneumonia accounted for only 5.11% of the total, and the number of cases of nonspecific interstitial pneumonia was not communicated by the authors. Other studies have described a lower prevalence of nonspecific interstitial pneumonia than that found in the present study; in one study, the prevalence was reported to be 4%.

In the literature, the prevalence of hypersensitivity pneumonia ranges from 1.5% to 14%. Similarly, the prevalence of pneumoconiosis ranges from 4% to 10.4%.

In the present study, the absence of pneumoconiosis and the frequency of hypersensitivity pneumonia can be explained by the environmental conditions to which the patients were exposed, that is, the characteristics of a region that has a relatively low concentration of industries and is more agricultural, with greater exposure to organic dust. In addition, the diagnosis of pneumoconiosis might have been established by means of other diagnostic methods, there being no need for confirmation by lung biopsy.

Regarding surgical complications, 2 patients (4.17%) required 4–5 cm auxiliary incisions. In
1 patient, there was difficulty in maintaining single-lung ventilation, whereas, in the other, it was not possible to introduce the camera and visualize the lung parenchyma due to extensive pleural adherences. This was also described by another group of authors, who recommended video-assisted thoracoscopy to be avoided in these situations.

Only 1 patient (2.08%) presented a postoperative complication (residual pneumothorax after chest tube removal, with no need of additional pleural drainage). In the patient sample studied, there were no deaths in the immediate postoperative period.

In the literature, the rate of postoperative complications varies. A study presented similar results, with only 1 case (2.94%) of pneumothorax as an adverse event. In other studies, complications were reported in 3.6% of the cases, and morbidities were described in 19.1% of the patients. In that study, the most common complications were pneumothorax, need for postoperative mechanical ventilation and hematoma in the incision site. In the literature, the mortality rate over a 60-day postoperative period was found to be 4.4%.

The comparison of the two techniques of surgical lung biopsy in terms of safety revealed no significant differences between open lung biopsy and video-assisted thoracoscopy.

In the present study, variables such as intraoperative blood loss, need for analgesics in the postoperative period and length of hospital stay were not evaluated. Various authors state that video-assisted thoracoscopy is superior to open lung biopsy in terms of those variables. However, there are studies stating that there are no differences. Therefore, further studies are needed in order to elucidate this point.

Currently, video-assisted thoracoscopy is the most widely used tool for performing lung biopsy in the diagnosis of DPLD. Since it is considered a minimally invasive technique, its use has increased. Therefore, the present study was needed in order to evaluate its role. Our results demonstrate that video-assisted thoracoscopy is highly efficient in diagnostic resolution. In addition, video-assisted thoracoscopy has proven to be safe for this purpose, since postoperative complications occurred in only 2.08% of the cases. The analysis of the results of our study revealed that, as long as there is an investigation and appropriate pre-planning, as well as careful selection of candidates, video-assisted thoracoscopy is an option with a good success rate when surgical lung biopsy is indicated.

References


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