PATHOLOGICAL STUDY OF THE SYNOVIAL TISSUE OF PATIENTS WHO UNDERWENT OPEN CARPAL TUNNEL RELEASE

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ABSTRACT

Objective: To determine whether a biopsy of the synovia of the carpal tunnel is able to identify systemic diseases that were not diagnosed by clinical examination and laboratory tests. Methods: Anatomical pathology P examinations of synovial tissue were performed in 46 patients that underwent open carpal tunnel release. Anatomical pathology examination with hematoxylin-eosin staining determined the intensity of the inflammatory process and the authors proposed a new classification of the injury according to the intensity of the inflammatory process. Results: The anatomical pathology examination showed that 56.6% were classified as grade I (unchanged inflammation), 32.6%, grade II (leukocyte infiltration

and moderate fibrosis), 4.3%, grade III (leukocyte infiltrate and intense fibrosis, the presence of fibrin and vascular neoformation) and 6.5%, grade IV (changes described above associated with the presence of local calcification and giant cells). Two patients with amyloidosis were classified as grade I and II and no stockpiles of amyloid material were found on their slides. Two patients with hyperparathyroidism and another with chronic kidney failure were classified as grade IV. Conclusion: Synovial biopsy of the carpal tunnel did not make early diagnosis of potential systemic diseases in patients with carpal tunnel syndrome, and adds extra costs to the procedure.

Keywords: Carpal tunnel syndrome. Biopsy. Nerve crush.

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INTRODUCTION

Carpal Tunnel Syndrome (CTS) results from compression of the median nerve inside the carpal channel.

Paget¹ described compression of the median nerve in the carpal tunnel after trauma. Marie and Foix² described the histopathology of carpal tunnel syndrome, correlating it with mechanical compression of the median nerve. Learmonth³ performed the first decompressive surgery. Phalen⁴ described a test for evaluation of median nerve compression. Semple and Cargil⁵ compared the results of several types of surgical incisions, concluding that there is no influence in terms of results, excepting with transversal incisions. Sunderland⁶ described the physiopathology of carpal tunnel syndrome, in which he attributed to the initial lesion an intrafunicular anoxia caused by obstruction of the intraneural venous return, produced by a pressure increase in the carpal tunnel of any nature.

Carpal tunnel syndrome occurs more frequently in patients between 40 and 60 years of age, and is more common among

women.⁷⁻⁹ Any condition that increases the content or that decreases the continent of the carpal tunnel may start off the symptoms. Among the systemic conditions that can lead to CTS, we emphasize rheumatoid arthritis, kidney failure, hypothyroidism, diabetes mellitus, amyloidosis, pregnancy, Raynaud's phenomenon and obesity. Local causes such as Colles' fracture, edema resulting from trauma or infection and tumors, ¹⁰ are among the agents that trigger the disease. CTS can also be associated with De Quervain's tenosynovitis, rhizarthrosis and trigger finger.

Paresthesia in the median nerve territory is the most frequent syndrome, and in more severe cases, wakes the patient up during the night. The clinical diagnosis is confirmed by the compression test of the examiner's thumb on the topography of the carpal tunnel¹¹ and by Tinel's¹² and Phalen's⁴ signs. The elevation test of the affected upper limb, insufflating the sphygmomanometer cuff on the arm between systolic and diastolic pressures, reproduces the symptoms. Atrophy of the thenar eminence is present

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in the more severe and long-lasting cases. It is considered that electroneuromiography presents sensibility of 90% and specificity of 60%. Ultrasonography, magnetic resonance and computed tomography are imaging exams reserved for cases in which the etiology arises from an expansive process.

The treatment of CTS should be based on its etiology. Local factors should be removed and systemic factors, corrected.

Patients with slight symptoms respond well to non-surgical measures such as non-hormonal anti-inflammatory drugs (NHAI), physiotherapy and orthosis. Some authors advocate the local infiltration of corticosteroids in cases in which thenar eminence does not present atrophy. Clinical intractability leads to the need for surgical treatment, which can be open or endoscopic. Anterior displacement of the median nerve, its adherence to the transverse carpal ligament and reflex sympathetic dystrophy are some of the possible complications of surgical treatment.

The aim of this study is to evaluate the need to perform routine anatomopathological examination on patients with CTS and to analyze the impact of biopsy performance on the cost of the procedure.

MATERIAL AND METHOD

During the period from January 1996 to November 2002 forty-six surgeries were performed for CTS treatment in our service. The surgical option was by the open route, with "L" shaped ulnar access²² (Figure 1), executing the opening of the transverse carpal ligament, external epineurotomy of the median nerve and biopsy of the synovia of the carpal tunnel. All the procedures were performed by the same surgeon.

The results of the anatomopathological exams were analyzed and classified according to the intensity of the inflammatory pro-



Figure 1 - L-shaped ulnar access.

cess, following the classification proposed by the authors. The slide staining method was with hematoxylin-eosin. This classification involves:

Grade I: unchanged inflammation. (Figure 2)

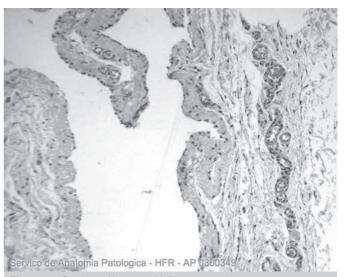


Figure 2 - Grade 1 - Absence of inflammatory changes

Grade II: modest leukocyte infiltrate and fibrosis, besides hyalinosis*. (Figure 3)

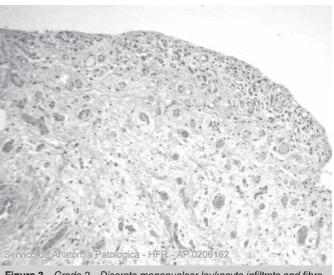


Figure 3 – Grade 2 – Discrete mononuclear leukocyte infiltrate and fibrosis, besides hyalinosis.

Grade III: intense leukocyte infiltrate and fibrosis, presence of fibrin and vascular neoformation. (Figure 4)

Grade IV: the alterations described above associated with the presence of local calcification and giant cells. (Figure 5)

The study was approved by the Committee of Ethics in Research of our institution.

^{*} Deposition of unstructured, amorphous, translucent and acidophilus material in the interstice.

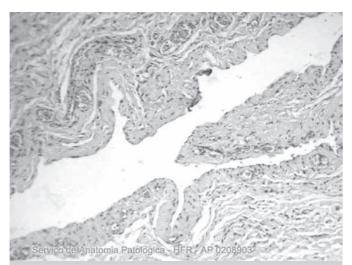


Figure 4 – Grade 3 – Intense mononuclear leukocyte infiltrate and fibrosis, presence of fibrin and vascular neoformation.

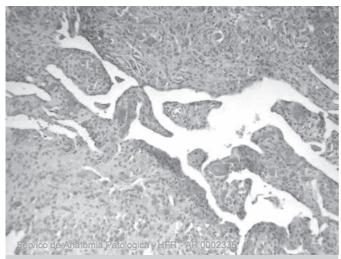


Figure 5 – Grade 4 – The same as grade 3 and the presence of local calcification and of giant cells.

RESULTS

Distribution by gender was 86.9% for females and 13.1% for males. The right side was affected in 64.5% and the left, in 35.5%. Age ranged from 21 to 86 years, with mean age of 51 years.

In the anatomopathological evaluation, 56.6% of the slides were classified as grade I, 32.6% as grade II, 4.3% as grade III and 6.5%, as grade IV. (Figure 6)

Two patients with amyloidosis were categorized in grades I and II and no deposition of amyloid material was found on the slides of these patients. Two patients that presented hyperparathyroidism and another with chronic kidney failure were classified as grade IV.

DISCUSSION

Surgical indication was based on clinical intractability and on the exclusion of corrigible systemic causes that could cause compressive symptoms.

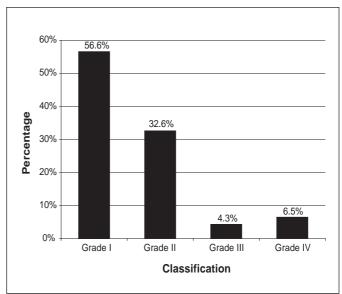


Figure 6 – Results encountered according to the intensity of the inflammatory process.

The anatomopathological examination of the synovia of the carpal tunnel of patients treated surgically was performed as routine. The goal was to identify at an early stage pathologies not diagnosed clinically or by laboratory exams.

All the patients operated in the six-year period were included in the study and their distribution in relation to gender, age and affected side was similar to that found in literature.⁷⁻⁹

The route of access used for the performance of the surgery was the "L" shaped ulnar access, 22 modification of the route proposed by Tubiana, 23 which crosses the palmar crease in a zigzag pattern.

An anatomopathological classification of the synovial tissue of the carpal tunnel was used to facilitate the analysis of the biopsy results, based on the intensity of the local inflammatory process. Grade I represents the absence of inflammatory alterations. Grade II shows discrete leukocyte infiltrate and fibrosis, besides hyalinosis. At grade III, there are intense fibrosis and leukocyte infiltrate, besides vascular neoformation. At grade IV, besides the lesions described, the presence of local calcification and giant cells is also observed.

Most of the patients (56.6%) did not present inflammatory process in the synovial tissue (grade I). Leukocyte infiltrate was observed in 43.4%. Patients with known diagnosis of hyperparathyroidism showed the presence of mononuclear inflammatory infiltrate, multinuclear giant cells and calcification focuses, characterizing moderate chronic synovitis, and were classified as grade IV. The same finding was observed in individuals with chronic kidney failure. In the patients with amyloidosis, the presence of amyloid deposits was not observed on the slides. One of the patients did not present

was not observed on the slides. One of the patients did not present inflammatory infiltrate (grade I) and another, slight inflammatory infiltrate (grade II).

Schuind et al.²⁴ classified lesion of the synovial tissue of the carpal tunnel at three levels, according to the intensity of fibrosis and the presence of zones of necrosis. In his series, which consisted of 34 samples, it was observed that 50% of the patients presented fibrous hypertrophy with localized areas of necrosis.

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Our microscopic findings, observed with the same stain (hematoxylin-eosin - HE), did not demonstrate areas of necrosis. We verified the presence of leukocyte infiltrate and, in some cases, vascular neoformation, which was not observed by Schuind et al.²⁴ Due to these observations, it was not possible to correlate our findings with the classification proposed by Schuind et al.²⁴ which prompted us to create a new classification.

No early diagnosis was obtained of systemic diseases not diagnosed clinically or by laboratory exams, through anatomopathological findings of synovial tissue of the carpal tunnel, which authorizes us to not perform the exam as routine.

Analyzing the treatment cost, excluding medical fees, an increase of around 10% was verified in performing the simulation of procedures carried out with synovial tissue biopsy in relation to those performed without biopsy.

CONCLUSION

We concluded that the performance of biopsy as routine is not necessary in patients undergoing surgical treatment for compressive carpal tunnel syndrome. Biopsy performance implies additional costs to the procedure and does not produce benefits in relation to the early diagnosis of systemic pathologies.

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