Study of the inflammatory process induced by injection of carrageenan or formalin in the rat temporomandibular joint

Estudo do processo inflamatório induzido pela injeção de carragenina ou de formalina na articulação temporomandibular de ratos

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ABSTRACT: The aim of this study was to evaluate the effects of the injection of two phlogistic agents, carrageenan and formalin, in the rat TMJ, and the inflammatory process induced by these substances. In this study, a total of 45 adult rats were distributed in two experimental groups and a control group. The animals were sacrificed after three hours, 24 hours, three days, seven days, and 15 days after a single injection of each substance. Histological data initially demonstrated an inflammatory process represented by acute infiltration, which later became mixed, and finally chronic in both experimental groups. Hyperplasia of the synovial membrane was observed after three days, being intense at seven days, and present after 15 days only in the formalin group. Local saline injection in the control group caused no inflammatory reaction. It was concluded that a single local injection of carrageenan or formalin was enough to induce inflammatory reaction in the TMJ and periarticular soft tissues, and that the resulting processes were similar, but more persistent in the formalin group.

DESCRIPTORS: Temporomandibular joint; Inflammation; Carrageenan, adverse effects; Formalin, adverse effects.

INTRODUCTION

Disorders of the temporomandibular joint (TMJ) frequently present an inflammatory component. However, an understanding of the pathogenesis of inflammation in the TMJ has been based on data from other synovial joints. A few experimental studies have evaluated only the immediate events of the inflammatory process in the TMJ.

Special features of the TMJ, such as its dense fibrous tissue covering, may influence its response as a synovial joint. Also, concentrations of some inflammatory substances in the TMJ differ from those in other joints, probably due to the denser innervations in the former. A satisfactory orofacial model should permit the study of inflammation...
in general, as well as of mechanisms that may be specific to TMJ tissues.

The purpose of the present study was to evaluate an experimental model in an attempt to analyze the effects of the injection of two phlogistic agents – carrageenan and formalin – in the rat TMJ.

MATERIALS AND METHODS

The study animals were 45 three-month-old female Wistar rats weighing 190 to 250 g. They were housed three per cage at 23°C in a 12 h light/dark cycle, and received an ordinary diet of rodent feed (Labina, Agribands Purina, Paulinia, Brazil) and water. The study was approved by the local research ethics committee.

The animals were anaesthetized by an intraperitoneal injection of xylazine hydrochloride (Rompum, Bayer, Porto Alegre, Brazil), 10 mg/kg of body weight and ketamine hydrochloride (Ketalar, Parke-Davis, Rio de Janeiro, Brazil), 25 mg/kg of body weight. The right preauricular area was shaved and cleansed with a povidone-iodine solution (Riodeíne, Rioquímica, São José do Rio Preto, Brazil). The animals were distributed in two experimental groups and a control group with 15 rats each. The rats received a local injection in the right TMJ of 0.02 ml of either 1% carrageenan (Sigma-Aldrich, St. Louis, MO, USA) in saline solution (carrageenan group), 5% buffered formalin solution (Laboratório Fórmula e Ação, São Paulo, Brazil) (formalin group), or 0.9% saline solution (Áster Produtos Médicos, Sorocaba, Brazil) (control group). Each solution was introduced by placing the tip of a 29-gauge needle (Becton Dickison, Curitiba, Brazil) just posteriorly to the zygomatic process of the temporal bone. The needle tip was then moved medioanteriorly towards the superior joint space.

The animals were sacrificed three hours, 24 hours, three days, seven days, and 15 days after the local injection of the solutions. Their heads were removed and fixed in 10% formalin solution (Quimex, Cotia, Brazil). The specimens were decalcified in 20% formic acid (Merck, Darmstadt, Germany) and coronal sections were obtained. Seven µm thick sections of the right TMJ were cut and stained with hematoxylin and eosin (Merck, Darmstadt, Germany). Injured and control sites were studied at the same magnification. The aim was to describe cellular infiltration and its type, periarticular soft tissue oedema, exudate extravasation, congested vessels, alterations in the joint capsule, synovial lining hyperplasia, and changes in the articular surfaces.

RESULTS

The animals made a recovery after the phlogistic agent injection. Food and water intakes were considered normal 24 hours after the injection, and the animals did not experience any weight loss. Edema was macroscopically observed on the lateral region of the face of the animals of the experimental groups in the first 24 hours. The histopathological findings are presented according to the substance and sacrifice time.

Carrageenan group

Three hours

Moderate neutrophilic exudation in the condylar neck region was observed, as well as edema and fibrin-hemorrhagic exudate. Lymphocytes were seen in the lumen of congested blood vessels (Figure 1A).

Twenty-four hours

Intense inflammatory exudation and edema were observed in the condylar neck region. Acute inflammatory cells extended among adjacent muscular fibers, dissociating them; an inflammatory process was also present in the joint capsule, with congested blood vessels and neutrophilic sub-synovial exudation (Figure 1B).

Three days

Mixed inflammatory infiltration was observed in the condylar neck region, extending among dissociated muscular fibers. Also, serofibrinous exudate and congested blood vessels could be seen in the condylar neck. In the articular capsule, mild subsynovial mononuclear infiltration was observed, as well as a slight proliferation of the synovial lining. Numerous inflammatory cells could be seen in the joint space (Figure 1C).

Seven days

Mild mononuclear infiltration in the condylar neck region and among muscular fibers in a restricted region near the condyle was observed. A few congested blood vessels were noticed. Marked proliferation of the synovial lining was present, and villous hyperplasia could be observed in some

areas of the joint. One specimen showed degenerative changes in the lateral pole of the condyle, with signs of bone resorption (Figure 1D).

**Fifteen days**

None of the specimens showed any signs of inflammation or changes in the articular tissues.

**Formalin group**

**Three hours**

Edema and mild neutrophilic exudation were observed in the condylar neck region, extending along the muscular fibers around the joint (Figure 2A).

**Twenty-four hours**

Marked inflammatory exudation and edema in the condylar neck region were observed. Cell infiltration was of the mixed type and scattered among adjacent muscular fibers, dissociating them. In the same region, serofibrinous-hemorrhagic exudate was observed. The inflammatory process in the joint capsule showed a great number of congested blood vessels and subsynovial inflammatory infiltration, while the synovium appeared to be normal.

**Three days**

Predominantly mononuclear cellular infiltration was seen in the condylar neck region, extend-
ing among dissociated muscular fibers. Also, se-rofibrinous-hemorrhagic exudate and congested blood vessels could be seen in the condylar neck. Subsynovial mononuclear cellular infiltration and congested blood vessels were observed in the articular capsule. Discrete hyperplasia of the synovial lining could be seen (Figure 2B).

Seven days

Intense inflammatory infiltration, predominantly mononuclear, was observed around the articular capsule, but it was only mild among muscular fibers. Serofibrinous exudate and congested blood vessels could be seen in the periarticular region. Marked hyperplasia of the synovial lining was present, and villous changes could be observed. One animal showed hyalinization and thickening of the condylar articular surface, with absence of the proliferative zone and atrophy of the fibrocartilage (Figures 2C and 2D).

Fifteen days

Mild mononuclear cellular infiltration dispersed among collagen fibers in the condylar neck region was observed. Some congested blood vessels were present and villous changes in the synovium were still evident (Figure 2E).

Control group

The injection of normal saline solution did not produce changes at any sacrifice time. The articular tissues presented characteristics of normality (Figure 3).

DISCUSSION

The present study showed that a single local injection of carrageenan or formalin results in an inflammatory reaction in the TMJ and periarticular tissues, with hyperplasia of the synovial membrane. The results were similar, although more persistent when formalin was used. Formalin has been used as a classic model to produce experimental inflammation\textsuperscript{11,20}. Carrageenan can induce a highly reproducible local antigenic inflammatory response\textsuperscript{17}. Carrageenan has been widely used to evaluate the anti-inflammatory effect of certain drugs\textsuperscript{4,11,15}.

Edema was the initial finding after the injection of either substance. A similar response has been reported with the use of these substances in the rat footpad or knee\textsuperscript{4,17}. In our experiment, edema was histologically observed between three and 24 hours, and practically disappeared after three days. This is in accordance with the findings reported in the related literature, which have shown intense edema after three hours, mild edema within three days and no edema at day seven\textsuperscript{14}. 

FIGURE 2 - Formalin group. A: mild neutrophilic exudation extending along muscular fibers three hours later; B: serofibrinous-hemorrhagic exudate, congested blood vessels and cellular infiltration in the condylar neck region three days later. c = condyle, d = articular disc (H&E, 100 X). (Continued on page 103).
Edema induced by carrageenan injection is gradual, measurable after one hour, maximum after three hours and lasts for several days. Our results also showed that cellular migration caused by carrageenan injection was moderate after three hours, but more intense than that caused by formalin injection. After 24 hours, both substances induced an intense inflammatory infiltration, which later became mild in the condylar neck region in the carrageenan group, but remained intense around the capsule in the formalin group, extending among muscular fibers in both groups. After 15 days, a mild inflammatory infiltration dispersed in the condylar neck region was observed only in the formalin group. Similar findings have been reported after carrageenan injection in the rat footpad.

Fibrinous-hemorrhagic/serofibrinous exudate was initially observed in both groups, being present for up to seven days. No exudate was observed after 15 days. Superficial fibrin deposition in synovitis cases has also been observed. Inflammatory cells enmeshed in fibrin webs have been demonstrated 24 hours after PLA\textsubscript{2} injection in the rat knee joint. Inflammatory alterations in the joint capsule, such as the presence of congested blood vessels and subsynovial inflammatory infiltration, were observed in both experimental groups between 24 hours and three days. Similar results have been reported after PLA\textsubscript{2} injection in the rat knee, with the higher number of congested blood vessels occurring after 24 hours, with maximum cellular infiltration observed after 96 hours.

Hyperplasia of the synovial lining was noticed in both experimental groups after three days, when mild cellular proliferation was present. It was intense after seven days, when it was possible to identify villous areas. However, after 15 days, it was present only in the formalin group. A histologic study of the rat knee joint has shown that 24 hours after injection of the irritating agent the synovial lining became markedly folded, and hyperplasia of the synovium was seen after 48 hours. Similar findings have been reported when synovitis was induced by trauma to the rat TMJ, and fibrous synovial thickening was noticeable 10 weeks after trauma. In addition, intense hyperplasia of the
synovial membrane occurred when anterior disc displacement was induced in the rabbit TMJ, and the hyperplastic synovial cells were present at six weeks postoperatively. It has been shown that the synovial tissue contains cells that are potentially angiogenic. Vessels from the peripheral capillary plexus and synovial membrane cells proliferate to provide the vascular origin for the connective tissue healing of the lesion.

Signs of degenerative alterations were observed on the condyle of one specimen from each group. In the carrageenan group there was a case of partial resorption of the condyle, while in the formalin group there was a case with hyalinization and thickening of the articular surface. The possibility of substances initially causing necrosis followed by degenerative alterations should be considered. However, such alterations occurred in only one specimen of each group of substance, which suggests a need for further studies. Similar findings, albeit more thorough, have been reported after indirect trauma in the rat TMJ. No similar findings with carrageenan or formalin have been reported in the related literature, probably due to the short periods of observation. On the other hand, some authors have reported that alterations such as bone erosion and loss of cartilage may occur in chronic inflammation, as caused by rheumatoid arthritis due to proliferative invasion of the synovial lining and the formation of pannus.

CONCLUSION

Both phlogistic agents used in this experiment were able to induce a local inflammatory reaction after only a single injection of each substance. However, despite the similarity between the inflammatory reactions caused by carrageenan or by formalin, including intensity and kind of cell infiltration, formalin induced more persistent inflammation. Thus, in this study, it was also possible to verify the response of the TMJ to these aggressive agents, which included initial acute inflammatory infiltration followed by chronic infiltration. Moreover, the response resulted in cellular proliferation of the synovial lining, hyperplastic and villous areas being observed, and here the reaction to formalin was again more persistent. This model may therefore be useful in further studies of the inflammatory response of the TMJ and make the development of new anti-inflammatory treatments possible.

REFERENCES


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