Effect of mesenchymal stem cells on the regeneration of structures associated with temporomandibular joint: narrative review

Efeito de células-tronco mesenquimais na regeneração das estruturas associadas à articulação temporomandibular: revisão narrativa

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ABSTRACT

BACKGROUND AND OBJECTIVES: Temporomandibular disorders are the problems involving the masticatory muscles and/or the temporomandibular joint and, among them, the bone and joint disc degenerative processes stand out. However, an effective treatment for these cases has not yet been identified in the literature. Thus, the primary objective of this study was to evaluate the reparative potential of mesenchymal stem cells on degenerative changes in structures associated with the temporomandibular joint in humans and animal models.

CONTENT: This narrative review included intervention trials in humans and animals that presented as an outcome variable the repair of joint discs and/or temporomandibular joint. The following databases were used: Pubmed, LILACS, Scielo and Google Scholar. Titles and abstracts were analyzed for the pre-selection of articles potentially eligible for inclusion in this review. The information collected from each article was included in a specific spreadsheet for this purpose containing the year of publication, article title, author’s name, study location, type of study, methodology, results, and conclusions. Two human studies and four animal studies were selected to compose the narrative review. In all studies presented, the presence of stem cells was able to improve the clinical, histological, and morphological parameters of the temporomandibular joint.

CONCLUSION: The use of stem cells seems to be effective in treating degenerative changes in temporomandibular joint associated structures in both animal and human models. However, due to the small number of studies and their heterogeneity, the results presented should be evaluated sparingly.

Keywords: Stem cells, Temporomandibular joint disc, Temporomandibular joint disorders.

RESUMO

JUSTIFICATIVA E OBJETIVOS: As desordens temporomandubulares constituem-se em um termo coletivo de problemas que envolvem os músculos mastigatórios e/ou a articulação temporomandibular. Dentre esses, destacam-se os processos degenerativos ósseos e do disco articular, contudo, ainda não foi identificado na literatura um tratamento eficaz para esses casos. Dessa forma, o objetivo desse estudo foi avaliar o potencial reparador das células-tronco mesenquimais sobre as alterações degenerativas das estruturas associadas à articulação temporomandibular em humanos e em modelos animais.

CONTEÚDO: Foram incluídos ensaios de intervenção em humanos e em animais que apresentassem como variável desfecho o reparo dos discos articulares e/ou da articulação temporomandibular. Foram realizadas buscas nas seguintes bases de dados: Pubmed, LILACS, Scielo e Google Acadêmico. Os títulos e resumos foram analisados para a pré-seleção dos artigos potencialmente elegíveis para sua inclusão. As informações coletadas de cada artigo foram incluídas em planilha específica para essa finalidade contendo o ano de publicação, título do artigo, nome do autor, local do estudo, tipo de estudo, metodologia, resultado e conclusões. Foram selecionados 2 estudos em humanos e 4 estudos em animais para compor este estudo. Em todas essas pesquisas apresentadas, a presença de células-tronco foi capaz de melhorar parâmetros clínicos, histológicos e morfológicos da articulação temporomandibular.

CONCLUSÃO: O uso de células-tronco parece ser eficaz no tratamento das alterações degenerativas das estruturas associadas à articulação temporomandibular. Todavia, devido ao reduzido número de estudos e sua heterogeneidade, os resultados apresentados devem ser avaliados com parcimônia.

Descritores: Células-tronco, Disco da articulação temporomandibular, Transtornos da articulação temporomandibular.

INTRODUCTION

Temporomandibular disorders (TMD) are the problems involving the masticatory muscles and/or the temporomandibular joint (TMJ). Pain, clicking and jaw limitation form
the classic triad of TMD symptoms, which may be related to the masticatory muscles, the pre-auricular area or both. The most common subtypes include muscular and joint pain, mainly the joint disc displacements and the degenerative joint diseases. Epidemiological studies have shown that 20-50% of the population presented at least one symptom associated to TMD.

When the conservative alternatives for the treatment of joint alterations are not effective, the disease may progress, resulting in the modification of the bone structures and the joint disc. The mechanism of the pain phenomenon is not yet completely clear, but it’s known that when the synovial membrane of the TMJ is damaged, lots of inflammatory cytokines are produced and secreted into the synovial liquid, promoting the degenerative and painful process, which can largely vary for each patient.

The joint surfaces of the TMJ are composed of the temporal bone, the mandibular fossa, the joint tubercle, and the mandibular condyle (MC). The scaly part of the temporal bone is part of the formation of the zygomatic arch and the TMJ, including in its extension the mandibular fossa and the joint tubercle, which acts as a bulkhead for the mandibular condyle during movement.

From the anatomical point of view, the MC is mediolaterally longer than in the anteroposterior direction, forming an ellipse in the transversal plane. The fibrous connective tissue extends to the periphery of the disc, fixing the joint disc to the mandibular condyle below and to the temporal bone above. Anteriorly and posteriorly, the MC connects to the TMJ disc through the capsular ligaments, while mediolaterally, it connects to the disc via collateral ligaments. This arrangement ensures close contact between the disc and the MC during joint movement.

The TMJ disc is featured as a fibrocartilage and is located between the MC and the temporal bone, providing the mandibular rotation and traverse movements. The extracellular matrix of the joint disc is composed by a mesh of type I collagen, which represents 80-90% of its dry weight, and glycosaminoglycans (GAGs), which represent up to 10% of its weight. Approximately two thirds of its cells are fibroblasts, while one third have a morphology similar to chondroblasts.

Although the functions and mechanical properties of the TMJ joint disc are already well described, its biomechanical characteristics in inflammatory and degenerative processes are still unclear. In this context, the study highlights that both the joint disc and synovial membrane can undergo degenerative processes after chronic inflammatory processes.

Chondrocyte death caused by apoptosis or necrosis is considered a central feature in clinical or experimental osteoarthritic pathology. Several therapies were applied with the intention of recovering the joint injury presented in animal models, but they were not able to stimulate local chondrocyte proliferation.

Mesenchymal stem cells (MSC) are multipotent cells present in a wide variety of tissues. They are a source of tissue originating from the mesoderm, such as bone, cartilaginous and adipose tissue. These cells are capable of adopting a fibroblast morphology and, under special conditions, differentiate into adipocytes, chondrocytes and osteocytes.

Evidence also suggests the existence of MSCs associated with synovial fluid obtained from patients undergoing TMJ arthrocentesis. Moreover, recent studies suggest that the use of stem cells in animal models of inflammatory pain, neuro-pathic pain and pain associated with cancer produce powerful analgesic effects. From the perspective that an effective treatment for cases of degeneration of TMJ-associated structures and the tissue forming and analgesic potential of MSCs has not yet been identified in the literature, the primary objective of this study was to assess the repair potential of MSCs on degenerative changes in structures associated with TMJ in human and animal models.

### CONTENTS

A literature review using the Pubmed, LILACS, Scielo and Google (gray literature) databases was performed. The search strategy was divided to make the selection of articles more sensitive. For the search in humans, the following descriptors were used: ("temporomandibular joint" [MeSH Terms] OR ("temporomandibular"[All Fields] AND “joint”[All Fields]) OR "temporomandibular joint” [All Fields]) AND ("stem cells"[MeSH Terms] OR ("stem"[All Fields] AND "cells"[All Fields]) OR "stem cells” [All Fields]) AND “humans” [MeSH Terms]. For the search on animals, the following were used: ("temporomandibular joint"[MeSH Terms] OR ("temporomandibular"[All Fields] AND “joint”[All Fields]) OR "temporomandibular joint” [All Fields]) AND ("stem cells” [MeSH Terms] OR ("stem”[All Fields] AND "cells” [All Fields]) OR "stem cells” [All Fields]) AND "animals” [MeSH Terms:noexp].

Manual search strategies were performed in the reference list of publications included in the study. Titles and abstracts were analyzed for pre-selection of articles that were potentially eligible for inclusion in the study.

The articles which were included performed animal or human intervention trials and presented as an outcome variable the repair of TMJ discs and/or bone structures using MSCs. Literature review studies, in vitro studies, subcutaneous studies and those that did not present the outcome variable were excluded. No restrictions regarding the period of publication of the manuscript were included, except in the case of Google Scholar, where the search was conducted in the last 3 years in order to identify the most recent studies. The information collected for the qualitative analysis were authors, year, place of publication, methodology, results and conclusions.

527 articles using the stipulated descriptors to evaluate the effectiveness of MSC use in the regeneration of ATM-related structures in humans were found. Of these, 52 were found through Pubmed and 475 through Google Scholar. No articles were found in LILACS and Scielo databases. After reading the titles and abstracts, 3 articles were selected for full reading, as shown in figure 1. Of these, only 2 were included for qualitative analysis, totaling 41 patients (28 women and 13 men), with ages ranging from 23 to 47 years old. The studies originated from Brazil and Italy.
For animal studies, 330 articles were found, 65 in Pubmed and 265 in Google Scholar. No articles were found in LILACS and Scielo databases. After reading the titles and abstracts, 14 articles were selected for full reading, as shown in figure 2. Of those, 10 were excluded, leaving 4 articles for full reading. The studies were conducted in Finland, Egypt, China, United States, and Italy. The data was tabulated and presented in table 1 (human studies) and table 2 (animal studies).

**Table 1. Presentation of the studies on human included in the review**

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<th>Authors and country</th>
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<td>de Souza Tesch et al.14 Brazil</td>
<td>One 27-year-old male patient presenting resorption of the mandibular condyles, mainly on the right side. Tissue used to obtain stem cells: nasal septum. Evaluation of the temporomandibular joint structures (TMJ) by computed tomography (CT). Variables evaluated: joint pain, maximum mouth opening, joint noises. Follow-up time: 1 week; 2 weeks; 1 month; 3 months; 6 months; 1 year.</td>
<td>All evaluated clinical parameters showed significant improvements, without the presence of adverse events. Through CT, the presence of new cortical and subcortical bone formation in the right TMJ, reduction of the joint space and partial recovery of the anatomy of the mandibular condyle and temporal bone could be observed.</td>
<td>Promising results with the use of stem cells.</td>
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<td>Carboni et al.15 Italy</td>
<td>Forty patients (28 women and 12 men) with intra-articular alterations, divided into 2 groups: test group (arthrocentesis + stem cells) and control group (arthrocentesis + saline solution). Age varied between 23 and 47 years old. Tissue used to obtain stem cells: adipose. Evaluation of TMJ structures through nuclear magnetic resonance (NMR). Variables assessed: joint pain, headache, cervical pain, tinnitus, brachialgia, dizziness, joint noises, maximum mouth opening. Follow-up period: 1 week; 1 month; 3 months; 6 months.</td>
<td>Improved parameters in both groups. The test group showed better clinical results regarding pain reduction and mouth opening amplitude. Regarding the repair of the joint structures, only in the test group the almost complete repair of joint disc morphology and capsular ligaments was observed.</td>
<td>The use of stem cells seems to be a promising alternative for the treatment of TMJ disorders.</td>
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DISCUSSION

After the critical reading of the studies, it was possible to identify the promising results for the MSCs in humans and animal models on the clinical, histological, and morphological perspectives. The tissue engineering has been searching for substances that could substitute the removed discs, including autogenous substances as the temporal muscle dermis and fascia, as well as the synthetic materials produced from silicone and polytetrafluoroethane. Over the last decades, the autologous chondrocytes implantation (ACI) has developed rapidly. However, the limited speed of cell proliferation and chondrocyte differentiation during in vitro cultures has restricted the use of ACI.

On the other hand, MSCs have become the object of increasing study in this area due to their high proliferative capacity, lower cost, and lower morbidity to the donor site. Its attainment has been associated with the tooth pulp, adipose tissue, umbilical cord and, more recently, the nasal septum. During the in vitro growth, MSCs form colonies called fibroblastic colony forming units (CFU-F). The phenotypic characterization of these cells reveals the expression of CD44, CD29, CD105, CD73 and CD166 and the absence of hematopoietic lineage markers such as CD34 and CD14. A sheep. Preparation of stem cell supports mediated using CAD-CAM. The stem cells were derived from the bone marrow of the iliac crest and the platelets were obtained from the sheep’s venous blood. Evaluation time: 4 months.

The rabbits were sacrificed after the third week and the joints were histologically processed. The rabbits were comprised of the TMJs.

The animals were divided into 2 subgroups according to the type of treatment. One subgroup (n=10) received an intra-articular injection (IA) of saline solution (PBS) and the other (n=10) received an IA injection of PBS plus stem cells. Group III (n=20) received an oil emulsion injection before treatment with PBS or PBS plus stem cells as in group II.

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Several therapeutic alternatives have been proposed for the recuperation of the function and the improvement of the clinical parameters in individuals that presented joint alterations. In this context, the joint disc changes are classified into 4 stages with progressive characteristics: a) Joint disc displacement. Stage 1: joint disc displacement with reduction; Stage 2: joint disc displacement with reduction and intermittent closed locking; Stage 3: joint disc displacement without reduction (closed locking); Stage 4: joint disc displacement without reduction with evidence of joint disc perforation or degenerative joint disease. Thus, it became clear that therapeutic approaches must be related to the severity of the alteration presented by the patient and its clinical

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<td>Ahtiainen et al.16</td>
<td>Finland</td>
<td>Ten animals submitted to the removal of the joint disc bilaterally. Discsinserted in the anterior portion of the condyle of the mandible.</td>
<td>Increase of 7 to 14 times in the expression of the molecules of aggercan and type I and type II collagen in the test group after 6 and 12 months, respectively. Regarding bone morphology, it was possible to observe, in the test group, a greater calcification in the mandibular condyle and temporal bone, as well as a reduction in the number of microcyst. In histological analysis, the joint cartilage in the test group seemed more regular when compared to the control group.</td>
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<td>Ciocca et al.17</td>
<td>Italy</td>
<td>A sheep. Preparation of stem cell supports mediated using CAD-CAM. The stem cells were derived from the bone marrow of the iliac crest and the platelets were obtained from the sheep’s venous blood. Evaluation time: 4 months.</td>
<td>The analysis of bone formation showed a statistically significant difference between the test group and the control group (p&lt;0.05). The bone growth values showed a significant difference between mandibular condyles with and without the presence of stem cells (p&lt;0.05). The mandibular condyle of the test group showed greater bone formation.</td>
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<td>Zhang et al.18</td>
<td>China</td>
<td>Rats of the lineage C57BL/6J Bone marrow cells marked with green fluorescent protein (GFP-CTM) were injected weekly into the TMJ, starting after 3 weeks of unilateral anterior crossbite stimulation (UACS) and continuing for 4, 8 and 12 weeks. Another group stopped receiving injections for 4 weeks after 8 weeks of injections. Evaluated measurements: DAP3, CD163 and ki67 expression.</td>
<td>UACS caused bone matrix loss and ankylosis. The weekly injections of stem cells largely restored these changes. The implanted stem cells expressed a high level of CD163 protein but showed no remarkable cell proliferation. The termination of the supply of exogenous stem cells reversed the restorative effects.</td>
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<td>Zaki et al.19</td>
<td>Egypt</td>
<td>Fifty rabbits were divided into three groups: Group I (n=10) did not receive treatment. Group II (n=20) was divided into 2 subgroups according to the type of treatment. One subgroup (n=10) received an intra-articular injection (IA) of saline solution (PBS) and the other (n=10) received an IA injection of PBS plus stem cells. Group III (n=20) received an oil emulsion injection before treatment with PBS or PBS plus stem cells as in group II. The rabbits were sacrificed after the third week and the joints were histologically processed. The TMJs of subgroup III, treated with combined therapy, showed improvement in all tested parameters.</td>
<td>Stem cells can be used safely and effectively to repair degenerative changes in the TMJs.</td>
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implication, with MSCs joint infiltration being another promising treatment for these diseases. Regarding the evidences presented in the two human studies, the improvement of clinical parameters could be observed, as well as the presence of bone repair assessed through imaging examinations. However, the number of studies and patients is still quite small and should be expanded in order to provide more robust and concrete results regarding the use of MSCs. In addition, gold standards should be established for the definition of diagnostic criteria in order to standardize the studies and their derived results.

As for the researches performed in animals, characterized as pre-clinical studies, it was possible to observe similar behavior to that in humans regarding the recovery of morphology of structures affected by TMJ osteoarthritis models. However, it’s worth noting that the different studies presented quite heterogeneous models. In this context, the study reported that studies performed in a single animal species are not able to provide standard experimental results with repeated TMJ changes, and more reliable animal models are still necessary. The exact same mechanism by which the presence of MSCs is able to improve the degenerative changes present in the TMJ is still little known. However, there is a belief that growth factors such as tumor growth factor β1 (TGF-β1) and the family of bone morphogenetic proteins (BMPs) are directly involved in this process. Moreover, the authors have demonstrated that the Wnt pathway regulation promotes the repair of cartilages mediated by the presence of MSCs.

As for the reduction of pain and inflammatory process, several studies report pain relief with the administration of MSCs in rodent models after systemic or local injection. Percutaneous injection of MSCs also caused long-term relief in a pilot study of lumbar discogenic pain in humans. Furthermore, another study demonstrated that the use of MSCs may be a new approach to the treatment of neuropathic pain, due to its significant reduction of pain. These results further contribute to the understanding of pain relief mediated using MSCs.

Regarding the animal studies, only one presented sample calculation for the performance of the experiments. In addition, the necessity for laboratory apparatus for the differentiation of the MSCs still hinders the approach of the technique within the dentistry practice, since the establishment of more simple protocols that could be used in the dentistry routine are necessary.

**CONCLUSION**

The use of MSCs seems to be effective in treating degenerative changes in TMJ-associated structures in both animal and human models. Due to the small number of studies and their heterogeneity, the results presented should be evaluated sparingly. More research is still necessary to produce more robust evidences for the implementation of this new therapy.

**REFERENCES**