

CLINICAL STUDY

Therapeutic effects of arthrocentesis in treatment of temporomandibular joint disorders

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ABSTRACT

OBJECTIVE: Analyse, compare and evaluate the effects of the treatment arthrocentesis with a control group of non-steroid drugs treated patients.

MATERIAL AND METHODS: 1752 out-patients (1293 females, 459 males) were examined within the years 2013–2017. We evaluated the following criteria: gender, age, visual analog scale for pain (VAS), inter-incisal distance and reducing intake of orally administered analgesics.

RESULTS: The pain level of the group of patients with arthrocentesis had stabilized at mark 2.5 on the pain scale, unlike in the control group after treatment with non-steroidal anti-inflammatory drugs, pain had stabilized at mark 9 on the pain scale. The inter incisal distance amongst the first group was 37 mm and in the control group only 27 mm, after completed treatment. The amount of applied analgesics in the group with arthrocentesis decreased to 100 mg daily, while in the control group, the dose was adjusted to 700 mg daily.

CONCLUSION: In the observed study, arthrocentesis was effective in reducing pain, amount of analgesics per day and improving the mobility of temporomandibular joint (TMJ). The results of this study we use in the ongoing project, that focuses on progressive and innovative methodology of endoscopically assisted arthrocentesis. (Tab. 4, Fig. 5, Ref. 39). Text in PDF www.elis.sk.

KEY WORDS: internal joint disorders, arthrocentesis, analgesics.

Introduction

The craniofacial area is one of the best innervated areas of the human body and the place where pains occur quite often – from ordinary headaches to unusual and difficult-to-explain pains such as trigeminal neuralgia. As the pains do not always arise from the joint itself, many authors believe that a wider term, such as craniomandibular disorders (23), should be introduced. Bell (3) proposed the term ‘temporomandibular’ disorders (TMD). This is a comprehensive term for a variety of clinical difficulties relating either to the muscles of the jaw, or the joint and associated structures, respectively both, i. e. muscles, joint and associated structures (7, 12, 19, 26, 34).

Generally, pain or hypomobility of the TMJ due to disc dislocation, infectious, traumatic or metabolic arthritis or adhesions

is considered a good indication for arthrocentesis. Most often it appears to be used in patients with anterior disc dislocation without reposition. By means of arthrocentesis, a transient expansion of the joint space can be achieved; alternatively, it may produce decompression in an area of increased intra-articular pressure, for example during exudative inflammatory processes. By means of lavage pus and infectious agents of inflammation, metabolic wastes or products of degradation processes are washed out (5, 14, 24, 36). Arthrocentesis of the TMJ is frequently identified as a lavage of TMJ and is conventionally concluded without viewing the joint. The principal task of arthrocentesis is to release adhesions, to wash out with inflammatory mediators and besides, to affect directly with medical treatments (6, 20). The success of this method is attributed to the initial distension of the joint space, where it can disrupt intra-articular adhesions between the disc and the hole, and flush out the waste products of inflammation (7).

Material and methods

At the center for Temporomandibular Joint Diseases at the Department of Stomatology and maxillofacial surgery Jessenius Faculty of Medicine and University Hospital in Martin, 1752 out-patients (1293 females, 459 males) were examined within the years 2013–2017.

Among individual research tasks, intensity of pain was objectified by means of a visual analogue scale of pain before treatment and in monitored periods after treatment lasting three, six, and

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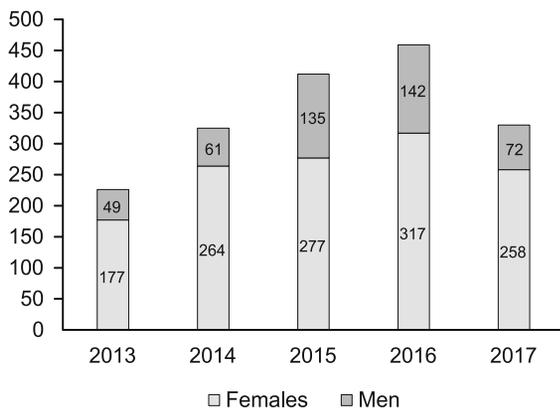


Fig. 1. Division of patients by year and sex.

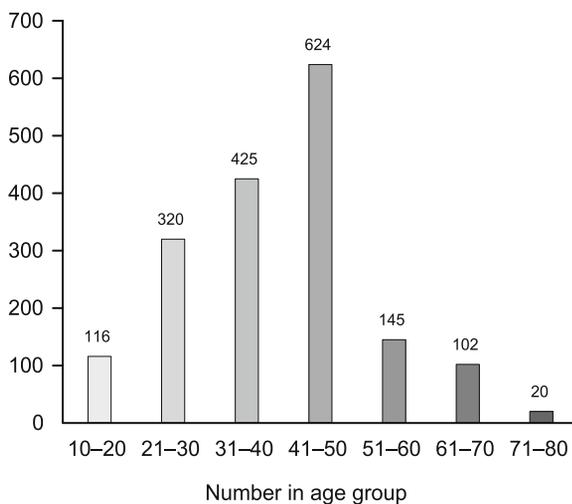


Fig. 2. Division of patients according to their age.

twelve months. Patients subjectively rated their pain on a scale numbered from 1 to 10. In the monitored periods, the inter incisal distance between the upper and lower teeth in the frontal section was assessed. An important factor was also the consumption of analgesics during monitored periods.

Arthrocentesis of the TMJ was done in 419 patients (348 females, 71 males). In 233 cases, the right side and in 186 cases the left side were affected and subsequently treated.

The control group was composed of patients who were orally administered non-steroid anti-inflammatory drugs. Arthrocentesis was performed in an aseptic operating room. It was an in-office procedure and required only simple technical equipment (Figs 1 and 2).

With the patient’s mouth open, an injection needle with a diameter of 1.0 to 1.2 mm was inserted into the posterior point of the upper joint space. The needle was obliquely and mediocranially inserted forward until the tip came into contact with the rear wall of the articular eminence. If there was inflammation of the joint, about 5 ml of local anaesthetic was applied under slight pressure to achieve dilation of the joint. A second needle with the same diameter was then inserted into the marked anterior point.

The joint is flushed with at least 100 ml of saline or Ringer’s solution. Generally, the first needle serves as an inlet needle and the second one as a drain needle. Lavage is done very slowly via a syringe.

After lavage of the joint space, the anterior needle was taken out and the posterior needle served for application of the medical preparations. Removal of the needles was followed by mandible exercise in order to promote the disruption of adhesions. In the group of patients diagnosed for arthrocentesis and in control patients, the level of pain was tested by means of a visual analogue 10-level pain scale (VAS) and the length of treatment was followed (in months) (Fig. 3) during monitored periods of 3, 6 and 12 months.

Results

In the group of patients after arthrocentesis, there was significant pain relief. After three months of follow-ups the value was close to mark 1 on the pain scale. In the control patients after three months of follow-ups, the degree of pain stabilized at mark 4 on the pain scale. After six months of the monitored period, the degree of pain suffered by the group of patients with arthrocentesis had not changed, while with the control group of patients, there was a significant deterioration with pain rising to mark 8 on the pain scale. Twelve months after completed treatment, the pain level of the group of patients with arthrocentesis had stabilized at mark 2.5 on the pain scale. With the control patients, twelve months after treatment, their pain had stabilized at mark 9 and either other medicaments had to be applied or the daily dose of non-steroidal anti-inflammatory drugs had to be doubled (Fig. 3).

During the follow-up period in both groups of patients, the inter incisal distance between the upper and lower tooth sets in the frontal aspect was assessed.

Before treatment, in a group of patients with indicated arthrocentesis, this distance was about 27 mm, and in the control group, approximately 26 mm. Three months after completed treatment, these values did not significantly differ from each other and remained at 27 mm, and 26 mm respectively. A significant change occurred six months into the monitored period, when in the group of patients with arthrocentesis, this value stabilized at 36 mm, while in the control group of patients the distance was only 28 mm. Twelve months after completed treatment, the inter incisal distance amongst the first group was 37 mm and in the control group only 27 mm (Fig. 4).

The fact that the application of targeted treatment for patients with arthrocentesis contributed to a significant reduction in the consumption of analgesics during treatment and after it is of key importance. The analgesics that the patients in both groups received daily were either Ibuprofen (400 mg) or Paracetamol (500 mg). While in the group of patients treated with arthrocentesis had a daily analgesic intake before treatment of 800 mg daily, the control patients had a daily intake of 850 mg.

After completed treatment and during the following three months, the amount of applied analgesics in the treatment group of patients decreased to 100 mg daily, while in the control group of patients, the dose was adjusted to 700 mg daily.

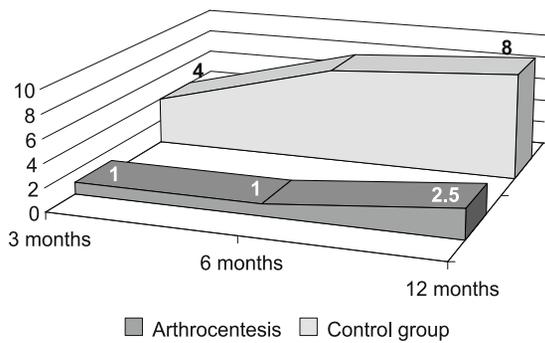


Fig. 3. Determination of degree of pain (VAS) after treatment.

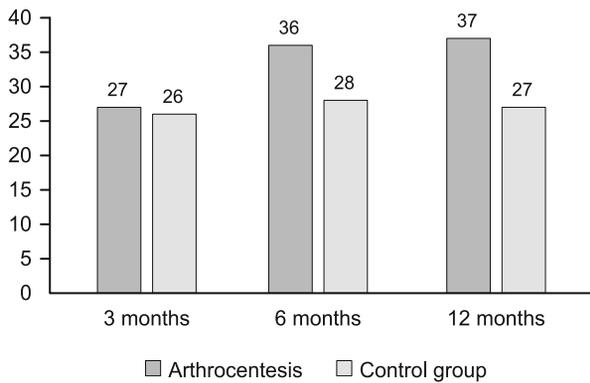


Fig. 4. Range of interincisal distance (mm).

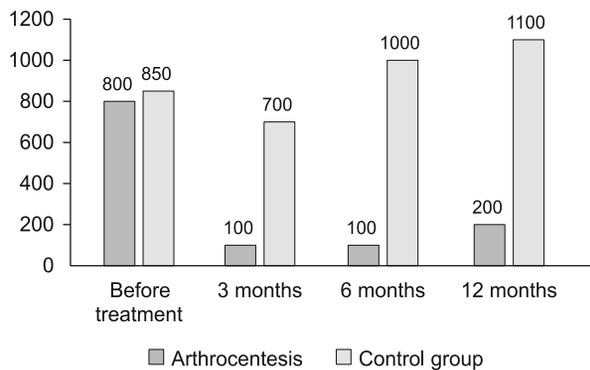


Fig. 5. Daily consumption of analgesics (mg).

During the six months following the end of treatment, in the treatment group of patients there was no change in increase of daily use of analgesics and this amount stabilised at a level of 100 mg. In the control group of patients, there was a rebound increase in daily consumption of analgesics, which reached 1000 mg.

Twelve months after completed treatment in the former group, the daily consumption of analgesics stabilized at 200 mg, whereas in the control group the daily consumption of analgesics increased to 1100 mg (Fig. 5).

Statistic assessment

Below each chart of measured values, arithmetic means are given as indicators of position and standard deviations as indica-

tors of variability. For each type of treatment, in table 1, p-values of the test of dissipation of variances (F-test) and correlation of mean values (Student's unpaired t-test) are listed. This test verifies a null hypothesis according to which the mean value of the treated group of patients is the same as the mean value of the control group. The test has two alternatives of counting of the test criterion and a p-value, which are determined by testing of dissipation values in both groups. Conformity of variances is tested by Fisher's F-test. The statistical significance of differences of mean values and variances of the variables are expressed in table 1 with a p-value of the t-test. p-values less than 0.05 indicate that studied mean values and variances are statistically significantly different (Tabs 1-4).

Discussion

A variety of intra-articular disorders respond well to arthrocentesis (26, 28). The success of this method is attributed to the initial distension of the joint space, where it can disrupt intra-articular

Tab. 1. Arthrocentesis – Control group.

Variable	p	
	F-test	t-test
vas	0.0248	0.9253
vas 3	< 0.0001	< 0.0001
vas 6	< 0.0001	< 0.0001
vas 12	< 0.0001	< 0.0001
i-i	< 0.0001	< 0.0001
i-i 3	< 0.0001	< 0.0001
i-i 6	0.0840	< 0.0001
i-i 12	< 0.0001	< 0.0001
sa	< 0.0001	< 0.0001
sa 3	< 0.0001	< 0.0001
sa 6	< 0.0001	< 0.0001
sa 12	< 0.0001	< 0.0001

vas – visual analogue scale (of pain), i-i – inter incisal distance, sa – consumption of analgesics

Tab. 2. Arthrocentesis.

Variable	p
vas	< 0.0001
vas 3-12	< 0.0001
i-i	< 0.0001
i-i 3-12	< 0.0001
sa	< 0.0001
sa 3-12	< 0.0001

vas – visual analogue scale (of pain), i-i – inter incisal distance, sa – consumption of analgesics

Tab. 3. Control group.

Variable	p
vas	< 0.0001
vas 3-12	< 0.0001
i-i	< 0.0001
i-i 3-12	< 0.0001
sa	< 0.0001
sa 3-12	< 0.0001

vas – visual analogue scale (of pain), i-i – inter incisal distance, sa – consumption of analgesics

Tab. 4. Arithmetic means and standard deviations.

Arthrocentesis	vas	vas 3	vas 6	vas 12	i-i	i-i 3	i-i 6	i-i 12	sa	sa 3	sa 6	sa 12
Mean	6.00	0.969	0.969	1.313	27.00	27.00	36.00	37.00	800	100	99.76	199.76
Deviation	1.117	0.304	0.174	0.527	0.789	0.660	1.142	0.894	86.40	39.43	44.96	79.02
Control group	vas	vas 3	vas 6	vas 12	i-i	i-i 3	i-i 6	i-i 12	sa	sa 3	sa 6	sa 12
Mean	6.01	4.01	7.98	8.99	26.08	26.00	27.94	27.00	851.5	700	1000	1101
Deviation	0.927	0.785	0.765	0.362	1.447	1.279	1.301	1.206	47.38	72.13	71.07	138.17

vas – visual analogue scale (of pain), i-i – inter incisal distance, sa – consumption of analgesics

adhesions between the disc and the hole, and flush out the waste products of inflammation (2, 5, 7, 12).

Frost and Kendell reported that arthrocentesis can be considered as a treatment choice between non-surgical treatment and arthroscopic surgery. Goudot et al studied comparatively about formation of pain and function after arthroscopy and arthrocentesis of the TMJ and reported that both arthroscopy and lavage are beneficial methods for function progress and decrease of pain (1, 8, 17). In the studies, it has been reported up to 91 % effective rate belonging to arthrocentesis to treat patients with anterior disc displacement without reduction (16). To determine the indication of case is important, because arthrocentesis can be inefficient in the patients who has bony changes, fibroankylosis and perforation of the disc (37). Treatment with arthrocentesis is very effective and the results are comparable to arthroscopy or arthrotomy. The success of arthrocentesis is indicated in 70–80 % of cases, which is comparable to arthroscopy (more than an 80 % success rate) (9, 11, 15, 25, 27).

Today, arthrocentesis of the temporomandibular joint is used not only in cases of acute closed lock but also in the treatment of various temporomandibular disorders. Thus, the most frequent indication is an acute anterior displacement of the articular disc without reduction or hypomobility of the joint with occurrences of disc adhesions (a stuck disc). It is possible to select arthrocentesis as a palliative procedure for patients with an acute episode of degenerative or rheumatoid arthritis and also for patients with a painful displacement of the disc with reduction, which rarely responds to conservative treatment. Treatment success is prominent in cases of acute patients or patients with a history of short-term problems (4, 30, 32).

Opinions on the amount of solution used for lavage of the joint space are not uniform, but it was discovered that for therapeutic lavage during arthrocentesis to be effective in washing away most undesirable substances from the joint space, at least 100 ml of solution should be used. Generally, it is recommended to use 200 ml of solution (25, 26, 34).

Exercise and treatment of the TMJ with arthrocentesis significantly contribute to improving the mobility of intra-articular adhesions. Some authors recommend performing these activities during arthrocentesis. We prefer the patient at rest during arthrocentesis and manipulation and mobilization exercises are carried out after removing the needles. It is assumed that the movement of the needles during exercise might cause traumatization of joint heads. Some authors consider diagnostic or therapeutic

intra-capsular injection as a variant of arthrocentesis (13, 26, 31, 33). They fill the joint space with a solution injected with just one single needle and thereby temporarily extend the joint space. This procedure is effective in cases of dislocation of the joint disc or adhesions (10, 21, 36, 38, 39).

Conclusion

The results of this study we use in the ongoing project that focuses on progressive and innovative methodology of endoscopically assisted arthrocentesis. The main idea is to use and combine the advantages of arthrocentesis and arthroscopy. The innovative nature of the project lies in the use of a single cannula during the arthrocentesis to introduce a 0.8 mm thick optical fiber into the intra-articular space, allowing visualization, diagnosis of the degree of tissue damage and subsequent treatment. After removing of the optical device, we aspirate and then inject the saline solution with the same cannula in a total dose of 100 ml. The total duration of the procedure will be from 20 to 40 minutes, depending on the patient's cooperation and the range of damage inner structures or adhesion of joint tissues.

The procedure can be performed in local anesthesia, as compared to arthroscopy, this leads to elimination of health risks of general anesthesia, costs associated with pre-operative examinations and hospitalization. At the same time, we anticipate benefits in patient management, reducing treatment times and patient sick leave, reducing analgesics consumption.

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