A controlled clinical pilot trial to study the effectiveness of ice as a supplement to the exercise programme for the management of lateral elbow tendinopathy

P Manias, D Stasinopoulos

Background: The use of ice as a supplement to an exercise programme has been recommended for the management of lateral elbow tendinopathy (LET). No studies have examined its effectiveness.

Objectives: To investigate whether an exercise programme supplemented with ice is more successful than the exercise programme alone in treating patients with LET.

Methods: Patients with unilateral LET for at least four weeks were included in this pilot study. They were sequentially allocated to receive five times a week for four weeks either an exercise programme with ice or the exercise programme alone. The exercise programme consisted of slow progressive eccentric exercises of wrist extensors and static stretching of the extensor carpi radialis brevis tendon. In the exercise programme/ice group, the ice was applied after the exercise programme for 10 minutes in the form of an ice bag to the facet of the lateral epicondyle. Patients were evaluated at baseline, at the end of treatment, and three months after the end of treatment. Outcome measures used were the pain visual analogue scale and the dropout rate.

Results: Forty patients met the inclusion criteria. At the end of treatment there was a decline in visual analogue scale of about 7 units in both groups compared with baseline (p<0.0005, paired t test). There were no significant differences in the magnitude of reduction between the groups at the end of treatment and at three months follow up (p<0.0005, independent t test). There were no dropouts.

Conclusions: An exercise programme consisting of eccentric and static stretching exercises had reduced the pain in patients with LET at the end of the treatment and at the follow up whether or not ice was included. Further research to establish the relative, absolute, and cost effectiveness as well as the mechanism of action of the exercise programme is needed.
confirm the LET diagnosis, performed all baseline and follow up assessments, and obtained informed consent.

Patients over 18 years old with lateral elbow pain were examined and evaluated in a private outpatient physiotherapy clinic in Ithaki between January 2003 and June 2004. All patients lived in Ithaki, Greece, were native Greek speakers, and were either self referred or referred by their doctor or physiotherapist.

Patients were included in the study if, at the time of presentation, they had been evaluated as having clinically diagnosed LET for at least four weeks. Patients were included in the trial if they reported (a) pain on the facet of the lateral epicondyle when palpated, (b) less pain during resistance supination with the elbow in 90° of flexion rather than in full extension, and (c) pain in at least two of the following four tests:

1. Tomsen test
2. Resisted middle finger test
3. Mill’s test
4. Handgrip dynamometer test

Patients were excluded from the study if they had one or more of the following conditions: (a) dysfunction in the shoulder, neck and/or thoracic region; (b) local or generalised arthritis; (c) neurological deficit; (d) radial nerve entrapment; (e) limitations in arm functions; (f) the affected elbow had been operated on; (g) had received any conservative treatment for the management of LET in the four weeks before entering the study.

All patients received a written explanation of the trial before entry into the study and then gave signed consent to participate. They were allocated to two groups by sequential allocation. For example, the first patient with LET was assigned to the exercise programme/ice group, the second patient with LET to the exercise programme alone group, and so on.

All patients were instructed to use their arm during the course of the study but to avoid activities that irritated the elbow such as shaking hands, grasping, lifting, knitting, handwriting, driving a car, and using a screwdriver. They were also told to refrain from taking anti-inflammatory drugs throughout the course of study. Patient compliance with this request was monitored using a treatment diary.

The exercise programme consisted of slow progressive eccentric exercises of the wrist extensors and static stretching exercises of the ECRB tendon. Three sets of 10 repetitions of slow progressive eccentric exercises of the wrist extensors at each treatment session were performed, with one minute rest interval between each set. Static stretching exercises of the ECRB tendon were repeated six times at each treatment session, three times before and three times after the eccentric exercises, with a 30 second rest interval between each repetition. Eccentric exercises of the wrist extensors were performed with the elbow on the bed in full extension, the forearm in pronation, the wrist in an extended position (as high as possible), and the hand hanging over the edge of the bed. From this position, patients flexed their wrist slowly while counting to 30, then returned to the starting position with the help of the other hand. Patients were told to continue with the exercise even if they experienced mild pain. However, they were told to stop the exercise if the pain became disabling. When patients were able to perform the eccentric exercises without experiencing any minor pain or discomfort, the load was increased using free weights. Static stretching exercises of the ECRB tendon were performed with the help of the therapist (PM). The therapist placed the elbow of the patient in full extension, the forearm in full pronation, and the wrist in flexion and ulnar deviation according to the patient’s tolerance. This position was held for 30–45 seconds each time and then released. The exercise programme was given five times a week for four weeks and was individualised on the basis of the patient’s description of pain experienced during the procedure. In the exercise programme/ice group, the ice was applied after the exercise programme for 10 minutes in the form of an ice bag to the painful area (facet of lateral epicondyle).

Pain and dropout rate were measured in this study. Each patient was evaluated at the baseline (week 0), at the end of treatment (week 4), and three months (week 16) after the end of treatment in order to see the intermediate effects of the treatment.

Pain was measured on a visual analogue scale (VAS), where 0 (cm) was “least pain imaginable” and 10 (cm) was “worst pain imaginable”. The pain VAS was used to measure the patient’s worst level of pain over the 24 hours before each evaluation, and this approach has been shown to be valid and sensitive.

The dropout rate was also used as an indicator of treatment outcome. Dropouts were categorised as follows: (a) withdrawal without reason; (b) did not return for follow up; (c) request for an alternative treatment.

The change from baseline was calculated for each follow up. Differences between groups were determined using the independent t test. The difference within groups between baseline and end of treatment was analysed with a paired t test. A 5% level of probability was adopted as the level for statistical significance. SPSS 11.5 statistical software was used for the statistical analysis.

RESULTS

Sixty two patients eligible for inclusion visited the clinic within the trial period. Twelve were unwilling to participate in the study, and 10 did not meet the inclusion criteria.
Ice for lateral elbow tendinopathy

At baseline there were more women in the groups (14 more in total). The mean age of the patients was about 40 years, and the duration of LET was about four months. LET was in the dominant arm in 85% of patients. There were no significant differences in mean age (p > 0.0005, independent t test) or the mean duration of symptoms (p > 0.0005, independent t test) between the groups. Patients had received a wide range of previous treatments (table 1). Drug therapy had been tried by 55%. All patients were manual workers.

Baseline pain on VAS was 8.70 (95% confidence interval 8.42 to 8.98) for the whole sample (n = 40) (table 2). There were no significant differences between the groups for baseline pain (p > 0.05 independent t test; table 2). At week 4 there was a decline in VAS of about 7 units in both groups compared with the baseline (p < 0.0005, paired t test; table 3). There were no significant differences in the magnitude of reduction between the groups at week 4 and week 16 (p > 0.0005 independent t test; table 3).

There were no dropouts and all patients successfully completed the study.

### DISCUSSION

The results obtained from this pilot trial are novel, as to date there have been no data comparing the effectiveness of an exercise programme with ice and an exercise programme alone for the reduction of pain in LET.

The ice may decrease the extravasation of blood and protein from new capillaries found in tendinopathy as well as decreasing the metabolic rate of the tendon. Both mechanisms promote healing of LET. In addition, ice can be used for symptomatic relief of pain. However, the findings of this trial indicate that ice as a supplement to the exercise programme offers no benefit in patients with LET. Therefore the reduction in pain at the end of the treatment and at the follow up was due to the exercise programme consisting of eccentric and static stretching exercises.

Standard eccentric exercises offer adequate rehabilitation for tendon disorders, but many patients with tendinopathies do not respond to this prescription alone. The load of eccentric exercises was increased according to the patients’ symptoms because the opposite has shown poor results.

Eccentric exercises were performed at a low speed in every treatment session because this allows tissue healing.

Exercise programmes appear to reduce the pain and improve function, reversing the pathology of LET, as supported by experimental studies on animals. The way that an exercise programme achieves the goals remains uncertain as there is a lack of good quality evidence to confirm that physiological effects translate into clinically meaningful outcomes and vice versa.

There are two types of exercise programme: home exercise programmes and exercise programmes carried out in a clinical setting. A home exercise programme is commonly advocated for patients with tendinopathies such as LET because it can be performed any time during the day without requiring supervision from a physiotherapist. Our clinical experience, however, has shown that patients fail to comply with the regimen of home exercise programmes. This problem can be solved by exercise programmes performed in a clinical setting under the supervision of a physiotherapist. For the purposes of this report, “supervised exercise programme” will refer to such programmes.

This exercise programme has been used in previous clinical trials on LET. However, it was the sole treatment in only two previous trials. A home exercise programme was the sole treatment in one of these two, and was administered in a totally different manner from the supervised exercise programme used in the present controlled clinical trial and the study of Stasinopoulos and colleagues. The differences were not only in the environment in which the exercise programmes were administered, but also in the development of the treatment protocol (type of exercises, intensity, frequency, duration of treatment). There is clearly a need for a clinical trial that would compare the effects of the supervised exercise programme treatment protocol, consisting of eccentric and static stretching exercises, with the home exercise programme treatment protocol used by Pienimaki et al.

Previous trials have found that a home exercise programme reduced the pain in patellar and Achilles tendinopathy. However, it was performed for about three months in all previous studies. In contrast, in the present controlled clinical trial and the study of Stasinopoulos and colleagues, a supervised exercise programme was administered for a month. Thus it seems that the supervised exercise programme may give long term clinical results in a shorter period of time than the home exercise programme. The most likely explanation for this difference is that a

### Table 1 Previous treatments of participants

<table>
<thead>
<tr>
<th></th>
<th>Exercise programme and ice</th>
<th>Exercise programme</th>
</tr>
</thead>
<tbody>
<tr>
<td>Drugs</td>
<td>12 (60)</td>
<td>11 (55)</td>
</tr>
<tr>
<td>Physiotherapy</td>
<td>4 (20)</td>
<td>4 (20)</td>
</tr>
<tr>
<td>Injection</td>
<td>4 (20)</td>
<td>5 (25)</td>
</tr>
</tbody>
</table>

Values are number (%).

### Table 2 Pain over the 24 hours before each evaluation

<table>
<thead>
<tr>
<th></th>
<th>Week 0</th>
<th>Week 4</th>
<th>Week 16</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exercise</td>
<td>8.60 (8.22 to 8.98)</td>
<td>1.70 (0.99 to 2.41)</td>
<td>1.50 (0.94 to 2.06)</td>
</tr>
<tr>
<td>programme and ice</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Exercise</td>
<td>8.80 (8.35 to 9.25)</td>
<td>1.90 (1.08 to 2.72)</td>
<td>1.60 (0.83 to 2.37)</td>
</tr>
<tr>
<td>programme</td>
<td></td>
<td></td>
<td></td>
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</tbody>
</table>

Values are mean (95% confidence interval) visual analogue scores where 0 = least pain imaginable and 10 = worst pain imaginable.

### Table 3 Change in pain over the 24 hours before each evaluation from baseline

<table>
<thead>
<tr>
<th></th>
<th>Exercise programme and ice</th>
<th>Exercise programme</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Week 4</td>
<td>-6.90</td>
<td>-6.90</td>
<td>&gt;0.0005</td>
</tr>
<tr>
<td>Week 16</td>
<td>-7.10</td>
<td>-7.10</td>
<td>&gt;0.0005</td>
</tr>
</tbody>
</table>

Values are mean visual analogue scores where 0 = least pain imaginable and 10 = worst pain imaginable. p Values for independent t test on change in VAS from baseline are shown.
supervised exercise programme achieves a higher degree of patient compliance. Studies to compare the effects of these two exercise programmes are required to confirm the findings of the present controlled clinical trial.

The present findings suggest that an exercise programme consisting of eccentric and static stretching exercises is adequate treatment for patients with LET. However, this trial does have some shortcomings. Firstly, a power analysis was not performed. Secondly, the samples were small and therefore the study was susceptible to lack of internal validity. Thirdly, although this study was not a randomised controlled trial because a genuine randomisation procedure was not followed, the use of sequential allocation to allocate patients to treatment groups allowed a true cause and effect relation to be demonstrated. Fourthly, the absence of a placebo/no treatment group meant that the study could not allow for normal fluctuations in the patients' symptoms, although LET is not a self limiting condition.47 Fifthly, the compliance of patients was not monitored when they were away from the clinic. Finally, the lack of blinding of patients, therapists, and assessor may be a reason for the effectiveness of the exercise programme. However, the blinding of patients and therapists would be problematic, if not impossible, of the exercise programme. However, the blinding of patients should be incorporated into the analysis of the effectiveness of the exercise programme in a future trial.

CONCLUSIONS

Ice as a supplement to an exercise programme offers no benefit to patients with LET. The exercise programme, consisting of eccentric and static stretching exercises, had reduced the pain in patients with LET at the end of the treatment and at follow up. Controlled studies are needed to establish the effects and the mechanism of action of such an exercise programme in LET. A cost effectiveness analysis should be incorporated into the analysis of the effectiveness of the exercise programme in a future trial.

REFERENCES


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What is already known on this topic

• Many clinicians use ice as a supplement to an exercise programme consisting of eccentric and static stretching exercises for the management of lateral elbow tendinopathy

What this study adds

• The findings of this trial indicate that supplementing the exercise programme with ice offers no benefit to patients with LET
• Therefore the reduction in pain at the end of the treatment and at the three month follow up was due to the exercise programme alone