Specific exercises for subacromial pain
Good results maintained for 5 years

Hanna C BJÖRNSSON HALLGREN 1, Lars E ADOLFSSON 1, Kajsa JOHANSSON 2, Birgitta ÖBERG 2, Anna PETERSON 1, and Theresa M HOLMGREN 1

In 2 previous publications we have demonstrated that specific exercises reduced the need for surgery in subacromial pain patients at 1-year follow-up. We have now investigated whether this result was maintained after 5 years and compared the outcomes of surgery and non-surgical treatment.

Patients and methods — 97 patients were included in the previously reported randomized study of patients on a waiting list for surgery. These patients were randomized to specific or unspecific exercises. After 3 months of exercises the patients were asked if they still wanted surgery and this was also assessed at the present 5-year follow-up. The 1-year assessment included Constant–Murley score, DASH, VAS at night, rest and activity, EQ-5D, and EQ-VAS. All these outcome assessments were repeated after 5 years in 91 of the patients.

Results — At the 5-year follow-up more patients in the specific exercise group had declined surgery, 33 of 47 as compared with 16 of 44 (p = 0.001) in the unspecific exercise group. The mean Constant–Murley score continued to improve between the 1- and 5-year follow-ups in both surgically and non-surgically treated groups. On a group level there was no clinically relevant change between 1 and 5 years in any of the other outcome measures regardless of treatment.

Interpretation — This 5-year follow-up of a previously published randomized controlled trial found that specific exercises reduced the need for surgery in patients with subacromial pain. Patients not responding to specific exercises may achieve similar good results with surgery. These findings emphasize that a specific exercise program may serve as a selection tool for surgery.
Table 2. Patients participating in the 5-year follow-up

|                        | Total | Non-operated | Operated |
|------------------------|-------|--------------|----------|
| Patients               | 91    | 49           | 42       |
| Specific/Control exercises | 47/44 | 33/16       | 14/28    |
| Sex: Men/Women         | 60/31 | 31/18       | 29/13    |
| Age at follow-up, mean (range) | 58 (38–69) | 57 (38–69) | 58 (39–69) |
| Patients lost between 1-year and the present 5-year follow-up | 4 | 3 | 1 |

*Previous randomization

as weakness in external and internal rotation and pathologic infraspinatus and subscapularis tests. All had undergone previous exercise therapy in primary care with an unsatisfactory result. Inclusion and exclusion criteria are listed in Table 1 (see Supplementary data). The specific exercise program focused on eccentric exercises for the rotator cuff and both eccentric and concentric exercises for the scapula-stabilizing musculature. The control exercise program included unloaded range of motion exercises for neck and shoulder without progression. The programs are described in detail in previous publications (Holmgren et al. 2012b, Hallgren et al. 2014). At the 3-month follow-up a shoulder surgeon blinded to the type of exercises asked the patients if they wanted to go through with surgery and in that case an ASD was performed as soon as possible. Surgery was performed by 1 of 2 experienced shoulder surgeons not involved in the study and included arthroscopic inspection of the glenohumeral joint and subacromial space, bursal and acromion resection. A supervised exercise program commonly used after ASD was performed postoperatively (Holmgren et al. 2012a). The patient’s choice of surgery or not resulted in 4 groups of patients after the 3-month assessment: specific non-operated, specific operated, control non-operated and control operated (Table 2, Figures 2, 3). A second follow-up was performed 1 year after inclusion. At all follow-ups (3 months, 1 and 5 years) the same shoulder surgeon, blinded to group assignment, recorded the Constant–Murley (C–M) score, Disability of the Arm Shoulder and Hand questionnaire (DASH) Score (Swedish version), Visual Analogue Scale (VAS) (0–100 mm) assessing pain intensity at rest, at night and at arm activity during the last 24 hours, EQ-5D, and EQ-VAS.

**5-year follow-up**

All 95 patients who participated in the 1-year follow-up were invited to a 5-year follow-up performed by a shoulder surgeon blinded to the initial group randomization. The data collection was identical to the 1-year follow-up including the patient’s choice of surgery or not and the clinical outcome measurements described above (Hallgren et al. 2014). The patients also filled in a questionnaire asking for use of health care, present shoulder symptoms, recurrence, and shoulder exercise habits during the past 4 years. Ultrasound examinations of the rotator cuff were performed by an experienced assessor, blinded to the findings at inclusion. A Siemens Acuson Sequoia 512 (Acuson, Mountain View, CA, USA) with a variable 8–10 MHz linear array transducer was used at all examinations. The status of the rotator cuff was divided into: intact, partial-thickness tear (PTT), or full-thickness tear (FTT) referring to the depth of the tendon (Björnsson et al. 2011). Tear size in mm was not measured. Tear progression was defined as progression from intact tendons at baseline to a partial- or full-thickness tear or from an initial partial- to a full-thickness tear at the 5-year follow-up. A full-thickness tear at inclusion that had enlarged to affect an adjacent, previously intact, tendon was also considered a progression.

**Statistics**

Pearson’s chi-square test was used to compare the proportion of patients choosing surgery in the originally randomized group, and also for the proportions of patients with progression of a cuff tear. Since some patients during this period needed surgery in addition to exercises, group comparison at the 5-year follow-up was performed using a paired t-test. p < 0.05 was considered significant.

**Ethics, registration, funding, and potential conflicts of interest**

Ethical approval was obtained for the 5-year follow-up from the regional committee for medical ethics in Linköping 2016-10-27 (dnr:2016/444-32). Written consent to participate in the study was collected from all patients after verbal and written information. The original trial was registered at Clinical trials: NCT01037673. The study was funded by the Linköping University Hospital and Linköping University but no other support, financial or other, was received for this study. No competing interests declared.

**Results**

**5-year follow-up**

At the 5-year follow-up 91 of the 95 invited patients could be reassessed (Figure 1). Any patient operated or re-operated had had this procedure performed at least 1 year prior to the 5-year follow-up. The proportion of patients not wanting surgery, who were satisfied with the exercise treatment, was still after 5 years larger (p = 0.001) among those originally randomized to the specific exercise group (33/47) compared with the control group (16/44). Between the 1-year and 5-year follow-ups 2 patients had chosen ASD, both initially randomized to the specific exercise group (Figure 1). All patients in the 4 different groups continued to improve in mean C–M score between the 1- and 5-year follow-ups (Table 3, see Supplementary data, Figure 2). There were no clinically relevant changes in the mean DASH scores between the 1- and 5-year follow-ups (Table 3, see Supplementary data, Figure 3).
When dividing the cohort into non-operated and operated patients from both exercise groups, the non-operated group had reached a significantly higher mean C–M score of 90 points (95% CI 82–90) compared with the operated group at 81 points (95% CI 77–85) (p = 0.002) at the 5-year follow-up (Table 4). At the 5-year follow-up non-operated patients scored better in pain at rest (p = 0.05) and at night (p = 0.02).

From baseline to 5-year follow-up the change in mean C–M score was 38 points in the non-operated group and 42 points in the operated group. A similar improvement was seen in the mean DASH score in operated (24 points) and non-operated patients (19 points) (Table 3, see Supplementary data, Figure 3). No clinically relevant changes were seen in the VAS, EQ-5D, and EQ-VAS recordings during the same time period (Table 3, see Supplementary data).

The 5-year follow-up questionnaire revealed that 7 of 49 individuals in the non-operated group had had further treatment after the 1-year follow-up, 5 a subacromial corticosteroid injection and 2 had further physiotherapy instructions. In the operated group 4 of 42 patients had received further treatment, 3 were re-operated and 1 had had osteopathy treatment. All 3 reoperations included acromioplasty, biceps tenotomy, and lateral clavicle resection. These re-operated patients had a similar 5-year outcome in all of the outcome measures as
Table 4. Mean Constant-Murley score (C–M) and standard deviation (SD) in operated (n = 42) and non-operated (n = 48) patients at 3-month, 1-year and 5-year follow-ups for the 90 patients with 5-year C–M score

| C–M score at | Group       | Mean | SD |
|--------------|-------------|------|----|
|              | Non-operated| 79   | 12 |
|              | Operated    | 45   | 20 |
|              | Total       | 63   | 24 |
| 3 months     | Non-operated| 86   | 12 |
|              | Operated    | 74   | 18 |
|              | Total       | 80   | 16 |
| 1 year       | Non-operated| 90   | 11 |
|              | Operated    | 81   | 15 |
|              | Total       | 86   | 14 |
| 5 years      | Non-operated| 90   | 11 |
|              | Operated    | 81   | 15 |
|              | Total       | 86   | 14 |

C–M score = Constant-Murley Shoulder Assessment Score 0–100 points (100 points = maximum shoulder function). 1 patient of the total cohort was not assessed at 5-year follow-up with C–M score.

Table 5. Rotator cuff status, assessed with ultrasound. Findings from baseline in the original RCT and from the 5-year follow-up divided into those treated with surgery and those without surgery up until the 5-year follow-up

| Rotator cuff status            | Total     | Operated   | Non-operated |
|-------------------------------|-----------|------------|--------------|
|                               | n = 90 ±  | n = 42     | n = 48       |
| Baseline                      |           |            |              |
| Intact                        | 64 ± 19   | 26 ± 9     | 38 ± 11      |
| Partial thickness tear        | 17 ± 13   | 9 ± 11     | 8 ± 7        |
| Full-thickness tear           | 9 ± 2     | 7 ± 2      | 2 ± 2        |
| 5-year follow-up              |           |            |              |
| Intact                        | 52 ± 13   | 19 ± 11    | 33 ± 12      |
| Partial thickness tear        | 18 ± 8    | 11 ± 7     | 7 ± 4        |
| Full-thickness tear           | 20 ± 12   | 12 ± 8     | 8 ± 4        |
| Progression c                 | 26 ± 16   | 16 ± 9     | 9 ± 7        |

*Arthroscopic subacromial decompression.
†One patient of the total cohort was not assessed with ultrasound.
‡Tear progression was defined as progression from intact tendons at baseline to partial thickness tearing or to full-thickness tearing or from partial tearing to full-thickness tearing at the 5-year follow-up. Additional full-thickness tearing in a previous intact tendon was also considered a progression of tearing.

compared with the rest of the cohort. 44 of the patients in the non-operated group reported that they had no or slight shoulder dysfunction compared with 31 in the operated group 5 years after inclusion. None of the patients in the non-operated group was worse compared with the 1-year assessment but 4 patients rated that they had the same symptoms ongoing. In the operated group 1 patient rated that he was worse and 3 persons that they still had the same symptoms as at the 1-year follow-up. In the non-operated group 28 patients had continued to perform exercises involving the shoulder compared with 17 in the operated group.

The ultrasound examination at 5 years showed that there were 38 rotator-cuff tears, including both partial- and full-thickness tears, in the cohort as compared with 26 tears at baseline. Significantly more patients (n = 16) in the operated group had progression of the tendon affection or a new tendon lesion as compared with (n = 9) the non-operated group (p = 0.002) (Table 5).

Discussion

Our main findings are that after 5 years more patients in the specific exercise group could still avoid surgery as compared with the unspecific exercise group and that patients who had not benefited from exercise treatment had a good outcome after surgery. Supervised exercise as the first line of treatment for subacromial pain is supported by results from other randomized trials and this study adds further evidence to the current recommendations (Brox et al. 1999, Haahr and Andersen 2006, Ketola et al. 2013, Haik et al. 2016). Our exercise program included both eccentric and concentric exercises for the rotator cuff and the scapula-stabilizing muscles. Pain was allowed to a certain limit and progression of load was guided by a pain-monitoring model (Thomee 1997). The rationale was that an increased range of motion, strength, and endurance would help to normalize the scapulohumeral kinematics and centralize the humeral head in the glenoid fossa during movement (Kromer et al. 2013, Maenhout et al. 2013, Struyf et al. 2013). Exercises are also hypothesized to have an inhibitory effect on central sensitization that may occur in many of the unilateral subacromial pain patients’ symptoms (Sanchis et al. 2015). Since subacromial pain has a multifactorial origin it is impossible to know which one of the components, or a combination of them, could explain the positive outcome after our specific exercise strategy (Lewis 2016). Reasons for the remaining effect in the current study, 5 years after a 3-month specific exercise intervention, are unclear. Patients may have learned to correct their shoulder kinematics to use their shoulder more functionally over the years (Curry et al. 2015). Also, a likely positive effect of the program was the “vocal treatment”, including information on their shoulder disorder, ergonomics, and posture correction (Adolfsson 2015, Lewis 2016). The mean age in the cohort was 58 (38–69) (see Table 1). The mean C–M score of the cohort was 86 which corresponds well with age and sex-adjusted C–M scores in the healthy population (Katolik et al. 2005). This reflects that the patients in the present study, on a group-level, reached a very good outcome.

Brox et al. (1999) compared surgery, supervised exercises, and placebo, and found that 25% of the patients in the placebo group reported a satisfactory result and contributed this to the natural course of the disease. In our study 16 of the 44 patients in the control exercise group chose not to be operated despite previous long-standing symptoms and an unsatisfactory result of physiotherapy in primary care. The positive result in this third of the group might be explained by multiple factors, the
natural course being one (Arroll and Goodyear-Smith 2005, Crawshaw et al. 2010).

When considering the other objective of this study, to compare surgical and non-surgical treatment, we found that the change over time in mean C–M score was well above the level for clinical relevance, reported to be between 17 and 24 points, in both operated and non-operated groups (Holmgren et al. 2014). These results are also in line with the clinically relevant pain reduction displayed in the VAS recordings and the overall patient satisfaction in both groups (Tashjian et al. 2009). The operated group showed a similar clinical improvement but this occurred after the surgical intervention.

The presence and progression of cuff tears was more often found in the operated group, a result that is in line with our previous study, where we found that patients with full-thickness tears and the lowest baseline scores were more prone to choose surgery (Hallgren et al. 2014). A structural cuff pathology may in part explain the inferior result in the operated group as rotator cuff disease may be the leading cause of prolonged shoulder pain and disability (Adler et al. 2008). Also the non-operated group included patients with progression of structural lesions that may be the result of natural aging, despite which they had an excellent 5-year result measured with several different outcomes. Multiple factors not related to pathoanatomy, such as mental health, age, genetics, comorbidities, and female sex, are found to influence outcome after treatment of subacromial pain with or without cuff tears (Curry et al. 2015, Lewis 2016). The multifactorial cause of symptoms may explain why a specific exercise strategy addressing several mechanisms is successful for the majority of patients.

As a result of the growing body of evidence supporting structured exercises as treatment of subacromial pain, ASD has become questioned (Brox et al. 1999, Haahr and Andersen 2006, Ketola et al. 2013, Haik et al. 2016). Ketola et al. (2013, 2015) concluded that patients without satisfactory symptom relief after non-operative treatment did not do any better after surgery. These conclusions are in conflict with our findings that patients treated with ASD improved substantially and with the same magnitude as the non-operated. Comparison between previous controlled studies is, however, difficult because of difference in inclusion criteria and baseline scores. The cohort in the study by Ketola et al. (2013) may have included patients with other disorders not responsive to any of the treatments used. We used strict inclusion criteria and we believe that our study group was homogeneous in terms of symptoms and all patients rated low baseline values on the Hospital Anxiety and Depression scale (HAD), a screening tool for depression and anxiety (Zigmond and Snaith 1983, Holmgren et al. 2012b). Understanding of the individual pathomechanisms is difficult but the results from other and our studies appear to confirm that a specific exercise strategy should be the initial treatment for subacromial pain with or without small rotator cuff tears (Holmgren et al. 2012b, Ketola et al. 2013, Hallgren et al. 2014). Acromioplasty can be recommended for patients without clinical signs of major cuff dysfunction and with unsatisfactory relief from specific exercise treatment.

A limitation of our study is the lack of an observational group to follow the natural course, but since all patients had been recommended some kind of exercises in primary care before inclusion in the original randomized trial we could not study the natural course. Since the investigation is based on a sample of patients with similar symptoms and radiological findings, performed at 1 hospital in a trial setting, the generalizability to all subacromial pain patients may be limited. Further, ultrasound is reportedly more accurate in detecting full-thickness tears than partial thickness tears (Cole et al. 2016). To handle this potential insecurity, we used experienced ultrasound assessors and the same equipment at all assessments. Strengths are the 5-year longitudinal data, both clinically and structural, on 91 of the 97 patients in the original cohort, making this study unique.

In summary, this 5-year follow-up supports the hypothesis that a specific exercise strategy should be the initial treatment of patients with subacromial pain. Patients not responding to specific exercises and those with more pronounced pathology may need surgery to reach a similar good result and the specific exercise program may serve as a selection tool for surgery.

Supplementary data
Tables 1 and 3 are available as supplementary data in the online version of this article, http://dx.doi.org/ 10.1080/17453674.2017.1364069

HB, TH, BÖ, KJ and LA conceived and designed the study protocol. TH and KJ designed the physiotherapy interventions. TH did the statistical analyses with assistance from a statistician (HM). HB and AP were the blinded assessors. HB drafted the manuscript, and TH, BÖ, KJ and LA contributed to the manuscript.

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