The Impact of Minimally Invasive Treatment for Rotator Cuff Calcific Tendinitis on Self-Reported Work Ability and Sick Leave

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Purpose: To examine the impact of rotator cuff calcific tendinitis on patients’ self-reported work ability and sick leave, to compare work ability and sick leave with shoulder function after minimally invasive treatment, and to assess which prognostic factors influence the change in work ability. Methods: A prospective cohort was analyzed in this study. The primary outcome measure was the single-question work ability score (0-10 points). Secondary outcome measures were quality and quantity of work, sick leave, functional outcome, and radiographic resorption. Potential predictive factors (treatment method, age, sex, resorption of the calcific deposit, physical work load, and work status) were tested in a statistical model. Follow-up was at 6 months and 1 year. Results: The study cohort consisted of 67 patients. The mean age was 49.6 ± 6.4 years and 45 (67%) were female. Physical workload was categorized as light (58%), medium (24%), and heavy (18%). Work ability score improved from a mean of 6.1 ± 2.8 to 8.5 ± 2.0 points after 1 year. Treatment with minimally invasive treatment techniques was associated with a reduction in partial or full-time sick leave from 28% to 6%. The mean days of sick leave a month declined from 3.3 to 0.8 days. Functional disability was greater in patients with partial or full-time sick leave. The physical workload turned out to be the most important patient associated factor predicting change in work ability. Conclusions: This study supports the hypothesis that rotator cuff calcific tendinitis has a significant impact on work ability and sick leave. Minimally invasive treatment resulted in a clinically relevant improvement in work ability score and decline in sick leave. In particular, patients with medium and high physically demanding work for the shoulder benefit from minimally invasive treatment to improve their work ability.

Level of Evidence: Level II, prospective comparative study.

Shoulder problems are common in the Netherlands, with an incidence in the primary care of around 19 per 1000 person-years. Shoulder disorders represent various clinical diagnoses, varying from International Classification of Diseases codes M75.0 to M75.5. Rotator cuff calcific tendinitis (RCCT; M75.3) represents a specific subgroup of shoulder patients with calcific deposits in the tendons. The prevalence of RCCT is between 2.7% and 10% in patients without shoulder pain and up to 40% in symptomatic patients. Clinical symptoms are generally described as activity-related pain similar to subacromial pain syndrome (SAPS). Synthes and Link Lima, outside the submitted work.

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The treatment initially consists of physiotherapy, analgesics, and a subacromial infiltration with corticosteroids. When primary treatment fails, minimally invasive therapies like high-energy extracorporeal shockwave therapy (ESWT) and ultrasound-guided needling (UGN) can be considered as an alternative for a surgical intervention. Multiple prospective studies and reviews have analyzed the functional outcome after treatment for patients with RCCT.26 Klik of tik om tekst in te voeren. However, little is known about the patients' work ability and sick leave before and after minimally invasive treatment for RCCT. Since this condition primarily affects people in their working age, it is of importance to know what the treatment effect is on work ability and sick leave. These are questions that too often remain unanswered in clinical studies. The purposes of this study were to examine the impact of RCCT on patients' self-reported work ability and sick leave, to compare work ability and sick leave with shoulder function after minimally invasive treatment, and to assess which prognostic factors influence the change in work ability. We hypothesized that RCCT would have a significant impact on work ability and sick leave and that this would be correlated with a greater functional disability. With regard to the prognostic factors, we hypothesized that work status (salaried or self-employed), work load, and successful resorption of the deposit would be the most important predictors for change in work ability.

Methods
The study was designed as a prospective cohort study. The patients were included in a published clinical randomized controlled trial comparing high-energy ESWT with UGN in patients with RCCT with the Constant score as main outcome measure.26 Klik of tik om tekst in te voeren. For the current study, both groups were analyzed as a cohort to answer our research questions concerning work ability and sick leave. Patients were included between May 2014 and December 2017. The study was registered in the Dutch clinical trial registration (NL4304/NTR4448) and approved by both the medical ethics review committee (METC, number NL44205.094.13) and the institutional review board (IRB number 2013.26, Spaarne Gasthuis, Hoofddorp, the Netherlands). Informed consent forms were signed by all participating patients.

Study Population
Sixty-seven consecutive patients referred to the outpatient orthopaedic clinic with clinical signs of non-traumatic anterograde-lateral sided shoulder pain when the arm was elevated were included. Patients were eligible for inclusion whether self-employed or with salaried work. Inclusion criteria for participation in this study were age >18 years, clinical sign of SAPS, standardized radiographs showing a calcific deposit in the rotator cuff tendons with a diameter of at least 5 mm in size, morphologic type I and type II deposits corresponding to the classification of Gärnner and Simons.8 Klik of tik om tekst in te voeren. (type I, sharply outlined and densely structured; type II, sharply outlined and inhomogeneous or homogenous with no defined border), symptoms for more than 4 months, a completed and unsuccessful nonsurgical treatment program including non-steroidal anti-inflammatory drugs, physiotherapy, and a subacromial infiltration with a corticosteroid. Exclusion criteria were the following: unemployment, ultrasonic signs of a partial or full rotator cuff tendon, clinical or radiographic signs of a resorption phase, calcific deposits in multiple tendons of the rotator cuff, osteoarthritis of the glenohumeral or acromioclavicular joint, adhesive capsulitis, previous shoulder surgery, ESWT or UGN to the affected shoulder, instability of the shoulder, rheumatoid arthritis, neurologic disorders or dysfunction of the upper limb, and the inability to give informed consent.

Inclusion
The medical history was taken and a clinical examination of the shoulder was performed. Standardized shoulder radiographs (anteroposterior, outlet-, axial-, and acromioclavicular view) and an ultrasound examination of the rotator cuff were obtained. Eligible patients were provided with written and oral information about the trial and had at least 1 week to consider participation. Patients who were willing to participate were contacted by the coordinating investigator (J.L.) for further evaluation and inclusion.

Treatment Procedures
The ESWT group was treated with 4 sessions of high-energy ESWT with a 1-week interval between sessions. Each session consisted of 2000 piezoelectric pressure pulses at a frequency of 4 Hz with a total energy flux density of 0.351 mJ/mm² resulting in a total energy amount of 2808 mJ. The Piezowave2 system (Richard Wolf GmbH, Knittlingen, Germany) was used as ESWT device. The UGN group was treated with a single procedure, performed in an outpatient clinical setting by 1 orthopaedic shoulder surgeon (A.v.N.) assisted by an experienced ultrasonographer. A double-needle UGN technique with aspiration and lavage of the calcific deposit was used. After the UGN procedure, one of the needles was placed in the subacromial bursa under ultrasound guidance and a mixture of 4 cc bupivacaine 0.5% (Pfizer Inc, New York, NY) and 1 cc Depo-Medrol 40 mg/mL (Pfizer Inc) was injected.

After treatment, both groups followed a standardized physical therapy program including active and passive
exercise mobilization techniques (centric and eccentric rotator cuff—strengthening exercises in combination with scapular stabilization) to increase power and range of motion and prevent muscular deficit or imbalance. Oral analgesics were administered for a maximum of 7 days postintervention when necessary. The use of additional over-the-counter analgesics was not systematically monitored.

**Work Ability and Sick Leave**

At baseline, all patients were asked if they were self-employed or had a salaried job. Subsequently, to assess the self-reported work ability, the single-item work ability score (WAS) question concerning the “current work ability compared with the lifetime best,” with a score of 0 (“completely unable to work”) to 10 (“work ability at its lifetime best”) was used as the primary outcome of this study.10-12

Clinical and Radiographic Evaluations

The following patient characteristics were collected: sex, age (years), body mass index, comorbidities, duration of symptoms, and hand dominance. At baseline and after 6 months and 1 year, the Constant-Murley score,16 the Disabilities of the Arm, Shoulder and Hand questionnaire (DASH),17 and a visual analog score (VAS), for average pain over a 1-week period were used for the clinical assessment. The minimal clinically important difference for the Constant-Murley score (8.3 points), DASH (10.2 points), and VAS pain (2.1 points) were determined based on available literature.18 The size, morphology, and amount of resorption (full, more than 50%, less than 50%, no change) of the calcific deposit were analyzed by standard shoulder radiographs at baseline and after 6 months. The length of the deposit was measured in terms of the maximum size of the longest axis in any direction. The radiographs were graded by another physician, who was blinded for the allocated treatment.

**Statistical Analysis**

Statistical analyses were performed by using SPSS, version 26.0 (IBM Corp., Armonk, NY). Continuous data are presented as means with standard deviations or 95% confidence intervals, and categorical variables as frequencies with accompanying proportions. Change of work-related and clinical outcome between follow-up moments was assessed with paired t-tests or McNemar tests or Wilcoxon signed rank tests where appropriate. To identify predictive factors for change in WAS, all potential predictive factors (treatment method [ESWT vs UGN], age, sex, resorption [less or more than 50%], physical work load [light vs medium/heavy], and work...
status [salaried vs self-employed]) were initially tested by univariate linear regression analyses. In case of significant association (adjusted significance level of 0.10), these factors were entered in a multivariate regression model.

Results

Baseline Characteristics

Between May 2014 and December 2017, a total of 67 patients were included (Table 1). One patient was lost to follow-up after 6 months. The mean age was 49.6 ± 6.4 years and 45 (67%) were female. The mean duration of symptoms was 3.2 ± 3.0 years, and the supraspinatus was the most frequently (87%) affected tendon. Most patients were employees (78%). Physical workload was categorized as light (58%), medium (24%), and heavy (18%). Nineteen patients reported sick leave (28%), of whom 9 patients (13%) were on permanent sick leave before treatment. Twenty-one percent of the patients stated that their shoulder symptoms were work-related.

Table 2. Work-Related and Clinical Outcome

| Sick Leave, n (%)            | Baseline (n = 67) | Six Months (n = 66) | One Year (n = 64) | P Value |
|------------------------------|-------------------|---------------------|-------------------|---------|
| None                         | 48 (71.6)         | 54 (81.8)           | 60 (93.8)         | −       |
| Partial                      | 10 (14.9)         | 7 (10.6)            | 2 (3.1)           | −       |
| Full time                    | 9 (13.4)          | 5 (7.6)             | 2 (3.1)           | −       |
| Days of sick leave a month, mean (95% CI) | 3.3 (1.8-4.8) | 2.2 (0.9-3.6) | 0.7 (0.0-1.5) | −       |

| Self-reported work-ability questionnaire, mean (95% CI) |
|--------------------------------------------------------|
| Quality                                                |
| 7.7 (7.0-8.4)                                          |
| 8.3 (7.6-9.0)                                          |
| 9.2 (8.7-9.6)                                          |
| <.001                                                 |
| Quantity                                               |
| 7.7 (7.0-8.4)                                          |
| 8.3 (7.6-9.0)                                          |
| 9.1 (8.7-9.6)                                          |
| <.001                                                 |
| Work-ability                                           |
| 6.1 (5.5-6.8)                                          |
| 7.8 (7.1-8.4)                                          |
| 8.5 (8.0-9.0)                                          |
| <.001                                                 |
| Functional limitations                                 |
| 4.7 (3.9-4.4)                                          |
| 7.4 (6.7-8.1)                                          |
| 8.0 (7.3-8.6)                                          |
| <.001                                                 |
| Clinical outcome measures, mean (95% CI)               |
| CMS                                                    |
| 66.9 (63.9-69.8)                                       |
| 81.3 (77.1-85.5)                                       |
| 87.2 (84.2-90.1)                                       |
| <.001                                                 |
| DASH                                                   |
| 37.4 (33.6-41.1)                                       |
| 20.1 (15.7-24.4)                                       |
| 14.1 (10.5-17.6)                                       |
| <.001                                                 |
| VAS (pain)                                             |
| 5.9 (5.5-6.3)                                          |
| 3.6 (2.9-4.2)                                          |
| 3.0 (2.3-3.7)                                          |
| <.001                                                 |

CI, confidence interval; CMS, Constant-Murley Score; DASH, Disabilities of Arm, Shoulder and Hand score; SD, standard deviation; VAS, visual analog score.

Work-Related and Clinical Outcomes

Table 2 presents the work-related outcomes of this study. Between 6 months and 1 year, 2 patients lost their job because of nonshoulder-related reasons. WAS improved from a mean score of 6.1 ± 2.8 to 8.5 ± 2.0 after 1 year. The change from baseline scores for work-related factors improved significantly for all 4 subcategories: work ability, quality of work, quantity of work, and functional limitations. Of the 9 patients who reported to be on full-time sick leave, 5 patients were still on full-time sick leave after 6 months and 2 after 1-year follow-up. The percentage of patients reporting sick leave was reduced from 19 patients (28%) to 4 patients (6%), with a decline in sick leave from a mean of 3.3 days a month to 0.7 days a month.

Overall, the Constant-Murley score, DASH, and VAS pain scores at final follow-up improved with clinically relevant differences in comparison with the baseline scores (Table 2). Radiographic resorption was complete in 53%, almost complete in 18%, minimally changed in 9%, and unchanged in 20%. Patients who were with partial- or full-time sick leave at baseline had

Table 3. Prognostic Variables for Work Ability

|                      | Six Months                          | One Year                            |
|----------------------|-------------------------------------|-------------------------------------|
|                      | β-coefficient (95% CI) | P Value          | β-coefficient (95% CI) | P Value          |
| Univariate           |                                     |                                   |                      |
| Age                  | −0.07 (−0.17; 0.04) | .20                               | −0.04 (−0.14; 0.06) | .41               |
| Sex                  | 0.43 (−1.02; 1.89) | .56                               | 0.53 (−0.79; 1.84) | .43               |
| High-energy ESWT vs UGN | 0.09 (−1.28; 1.47) | .90                               | 0.74 (−0.50; 1.99) | .24               |
| Resorption of the calcific deposit | 0.64 (−0.87; 2.15) | .40                               | 1.08 (−0.27; 2.43) | .11               |
| Workload (light vs medium/heavy) | 1.43 (0.08; 2.79) | .04                               | 1.55 (0.33; 2.76) | .01               |
| Work status (self-employed vs salaried) | 0.31 (−1.53; 2.15) | .74                               | 0.54 (−0.92; 2.00) | .46               |
| Final model           |                                     |                                   |                      |
| Workload (light vs medium/heavy) | 1.43 (0.08; 2.79) | .04                               | 1.55 (0.33; 2.76) | .01               |

NOTE. Boldface indicates statistical significance. ESWT, extracorporeal shockwave therapy; UGN, ultrasound-guided needling.
significantly lower Constant-Murley scores (58.5 ± 9.4 vs 70.2 ± 11.4, \( P < .001 \)) and DASH scores (48.5 ± 15.3 vs 32.9 ± 13.2, \( P < .001 \)) when compared with the group without sick leave. After 6 months (\( P = .05 \)) and 1 year (\( P = .006 \)), this difference remained significant for the DASH score.

**Prognostic Variables for the WAS**

Table 3 presents the results of the univariate analyses of each potentially prognostic factor for the change in WAS, as well as the final model. After 6-month and 1-year follow-up, only the physical workload, defined as light versus medium and heavy combined, was a predictive factor for a significant change in WAS (\( P = .01 \)) with a \( \beta \)-coefficient of 1.43 (95% confidence interval 0.08-2.79) in favor of the high-workload group. All other factors were not significantly associated with the change in WAS. Fig 1 graphically shows that the change from baseline scores for WAS in the medium physical workload group (1.0 after 6 months and 1.8 after 1 year) was significantly lower than the scores in the high physical workload group (2.9 after 6 months and 3.5 after 1 year).

**Discussion**

The principal finding of this study is that RCCT has a significant impact on patients’ work ability and sick leave. Treatment with minimally invasive treatment techniques was associated with a reduction in partial or full-time sick leave from 28% to 6%. The mean days of sick leave a month declined from 3.3 to 0.8 days. WAS improved from a mean score of 6.1 ± 2.8 to 8.5 ± 2.0 after 1 year. The physical workload turned out to be the most important patient-associated factor predicting change in WAS. In particular, the patients with medium-to-high physically demanding work improved the most. This is important information for the clinicians when discussing treatment options with their patients, and it might encourage patients with high physical demanding work to choose minimally invasive therapies.

Not much is known about the impact of RCCT on work ability and sick leave. Since this condition primarily affects patients of working age, information regarding these outcome measures is of great importance for both clinicians and patients. However, none of the 16 randomized controlled clinical trials analyzed in a previous meta-analysis discussed work-related outcome measures. Although data on RCCT are scarce, multiple studies have been published on work-related risk factors for SAPS. For work-related specific shoulder disorders, the biomechanical factor seems to be the most important, but psychosocial factors might also contribute. Van Rijn et al. concluded in their systematic review that forceful exertion in work, highly repetitive work, awkward postures, and high psychosocial job demand are associated with the occurrence of shoulder disorders. This was confirmed by a more recent meta-analysis by van der Molen et al. stating that there is moderate GRADE level evidence for a 2-fold chance of developing SAPS when being exposed to arm elevation and shoulder load during work.
In this study, the physical demands were classified based on the evidence-based exposure criteria for the work-relatedness of SAPS by the Dutch Center for Occupational Diseases. We found that a greater percentage of patients in medium and high physical demand jobs were on (partial) sick leave at baseline and reported a significant lower WAS with low functional performance scores. The present data support the hypothesis that in particular patients with medium or high physically demanding work for the shoulder might benefit from minimally invasive treatment.

The other factors that were tested in our predictive model were not significantly associated with the change in WAS. Engebretsen et al.\textsuperscript{20} found, in their RCT comparing radial ESWT and supervised exercise in patients with SAPS, that 12 or fewer years of education is the most consistent predictive factor for absenteeism and low functional outcome scores after 1 year. While they did not report the workload of these patients, there is probably a strong relationship between a lower education level and performance of higher physically demanding work. Seil et al.\textsuperscript{22} reported no difference in return to work between difference physical work load categories in a retrospective cohort study analyzing surgical outcomes for RCCT. Klik om tekst in te voeren. They concluded that the only difference in time to return to work was due to the presence of disability claims.

A strength of this study is that it contains 1-year follow-up data on both validated work-related outcome measures and clinical outcome measures. Moreover, the patients worked in a wide variety of occupational settings, which makes the results more generalizable than a selective sample of workers. Although the numbers in this study were relatively small, data from 95\% of the patients were available at 1-year follow-up and differences statistically tested.

Work-related outcome measures should be included more frequently in orthopaedic surgery research, as these parameters are relevant in the treatment of working-age patients and are frequently not reported in clinical trials. The use of a single-item measure has been validated in previous studies and has numerous advantages: it is short in length, requires less time to complete, and is more likely to be completed by the employee in comparison with multiple-item questionnaires.\textsuperscript{3-5} Klik om tekst in te voeren. Furthermore, it has been shown not to decrease the validity of the work ability information collected in comparison with multiple-item questionnaires.

Self-reported variables such as self-reported likelihood of the work-relatedness of the musculoskeletal disease, rating of the expected effectiveness of work-related interventions, presence of support from supervisors, and presence of modifiable job duties can play an important role in the assessment of work-related outcome measures. Furthermore, it would be interesting to validate if patient with high physical demanding jobs for the shoulder benefit more from minimal invasive treatment options than patients with low physical demanding jobs.

**Limitations**

One of the limitations of this study is that it did not include a control group and therefore we could not compare the outcome with patients receiving no treatment. Furthermore, since the study was primarily powered to look for a difference in the clinical Constant-Murley scores, it might have been under-powered for the outcome measure work ability and sick leave. Finally, since this study focused specifically on RCCT, statements made in this study are not generalizable to other conditions that cause SAPS.

**Conclusions**

This study supports the hypothesis that rotator cuff calcific tendinitis has a significant impact on work ability and sick leave. Minimally invasive treatment resulted in a clinically relevant improvement in work ability score and decline in sick leave. In particular, patients with medium and high physically demanding work for the shoulder benefit from minimally invasive treatment to improve their work ability.

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