

# The Effect of Rotator Cuff Repair on Natural History

## A Systematic Review of Intermediate to Long-Term Outcomes

Peter N. Chalmers, MD, Hunter Ross, BA, Erin Granger, MPH, Angela P. Presson, PhD,  
Chong Zhang, MS, and Robert Z. Tashjian, MD

*Investigation performed at the University of Utah, Salt Lake City, Utah*

**Background:** Rotator cuff disease can have a progressive natural history of increasing tear size and worsening function. It remains unknown whether rotator cuff repair alters this natural history.

**Methods:** A systematic review of the intermediate to long-term (minimum 5-year) results of operative rotator cuff repair and no repair of rotator cuff injuries was performed to compare (1) patient-based outcomes, (2) future surgical intervention, (3) future tear progression or recurrence, and (4) tear size. The no-repair group included both conservative treatment and surgical treatment without repair. After the application of selection criteria, 29 studies with 1,583 patients remained. Meta-regression was conducted to adjust for baseline age, sex, tear size, and duration of follow-up.

**Results:** Comparison of the repair and no-repair groups revealed no significant differences in terms of age ( $p = 0.36$ ), sex ( $p = 0.88$ ), study level of evidence ( $p = 0.86$ ), or Coleman methodology score ( $p = 0.8$ ). The duration of follow-up was significantly longer for the no-repair group ( $p = 0.004$ ), whereas baseline tear size was significantly larger in the repair group ( $p = 0.014$ ). The percentage of patients requiring additional surgery was significantly higher in the no-repair group after adjustment for age, sex, duration of follow-up, and tear size (9.5% higher in estimated means between groups [95% confidence interval, 2.1% to 17%];  $p = 0.012$ ). The likelihood of a recurrent defect (repair group) or extension of the prior tear (no-repair group) was not different between groups after adjustment for age, sex, duration of follow-up, and tear size ( $p = 0.4$ ). There were no differences between the repair and no-repair groups in terms of the Constant score after adjustment for age, sex, duration of follow-up, and tear size ( $p = 0.31$ ). The final tear size was significantly larger in the no-repair group than the repair group (967 mm<sup>2</sup> higher in estimated means between groups [95% confidence interval, 771 to 1,164 mm<sup>2</sup>];  $p < 0.001$ ).

**Conclusions:** At intermediate to long-term follow-up, rotator cuff repair was associated with decreased final tear size and decreased need for future surgery after adjusting for age, sex, duration of follow-up, and tear size. The likelihood of a recurrent defect after rotator cuff repair did not differ from that of tear extension after nonoperative treatment. Thus, rotator cuff repair may not alter natural history.

**Level of Evidence:** Therapeutic Level IV. See Instructions for Authors for a complete description of levels of evidence.

Rotator cuff disease can be progressive<sup>1-5</sup>. Although not all tears increase in size<sup>6-8</sup>, many partial tears become full-thickness tears<sup>1</sup> and many full-thickness tears increase in size<sup>1,3,5</sup>. Substantial pain and disability, pseudoparalysis<sup>9,10</sup>, and a characteristic set of chondral degenerative changes called *rotator cuff tear arthropathy*<sup>5,11,12</sup> can develop as a result. As the natural history of rotator cuff disease may not be benign and the clinical results of rotator cuff repair can be good, rotator cuff

repair often has been considered to be a reasonable treatment for rotator cuff tears<sup>13-16</sup>. Indeed, arthroscopic rotator cuff repair is one of the most commonly performed orthopaedic procedures in the United States and continues to increase in incidence<sup>17</sup>.

However, in the short term, most patients with a full-thickness rotator cuff tear will have excellent results with nonoperative treatment<sup>18</sup>. In addition, several randomized

**Disclosure:** This study was supported in part by the National Center for Research Resources and the National Center for Advancing Translational Sciences, National Institutes of Health, through Grant 5UL1TR001067-02 (formerly 8UL1TR000105 and UL1RR025764). On the **Disclosure of Potential Conflicts of Interest** forms, which are provided with the online version of the article, one or more of the authors checked "yes" to indicate that the author had a relevant financial relationship in the biomedical arena outside the submitted work and "yes" to indicate that the author had a patent and/or copyright, planned, pending, or issued, broadly relevant to this work (<http://links.lww.com/JBJSOA/A29>).

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TABLE I Study Characteristics

Variable	Repair Cohorts (N = 32)	No-Repair Cohorts (N = 13)	P Value	Test
Level of evidence ( <i>no. of cohorts</i> )			0.86	Fisher
I	3 (9%)	2 (15%)		
II	2 (6%)	1 (8%)		
III	9 (28%)	2 (15%)		
IV	18 (56%)	8 (62%)		
Approach ( <i>no. of cohorts</i> )			—	—
Arthroscopic	16 (50%)	—		
Open	16 (50%)	—		
Coleman methodology score* ( <i>points</i> )	62.5 ± 11.8	61.5 ± 11.3	0.80	T

\*The values are given as the mean and the standard deviation.

clinical trials with short-term results have suggested that rotator cuff repair does not provide either a clinically or statistically significant benefit over nonoperative treatment<sup>4,19,20</sup>. Nonoperative treatment does not reduce tear size or alter the natural history of rotator cuff disease<sup>1-5</sup>. Rotator cuff repair can result in an intact bone-tendon interface<sup>16,21,22</sup>, and these results can be maintained in the long term<sup>14,23-26</sup>. However, it remains unknown whether rotator cuff repair alters the natural history of rotator cuff disease.

The purpose of the present study was to conduct a systematic review of all published clinical studies with a minimum duration of follow-up of 5 years after rotator cuff repair and/or nonoperative treatment of rotator cuff disease in order to compare (1) strength and range of motion, (2) functional and patient-based outcomes, (3) the need for future surgical intervention, (4) the likelihood of future tear progression or recurrence, and (5) tear size. We hypothesized that rotator cuff repair would lead to increased strength and motion, improved outcomes, decreased need for future surgical intervention, no change in the likelihood of future tear progression or recurrence, and decreased final tear size when compared with nonoperative therapy.

## Materials and Methods

The present study was a systematic review of the literature. A search was performed with use of PubMed, Cochrane, and Embase databases. The search terms included *rotator cuff repair*, *rotator cuff tear*, *rotator cuff conservative*, *rotator cuff non-operative*, *rotator cuff nonoperative*, *outcomes*, *long-term*, and *long term*. The search was conducted in November 2016. The exclusion criteria were a minimum duration of follow-up of <5 years, lack of either physical examination findings or clinical data at the time of the latest follow-up, case reports, technique articles, review articles, a sample size of <10, reconstruction with a graft, tendon transfers, arthroplasty studies, and studies published in languages other than English. We manually screened the references of each included study to ensure that no studies were missed. The tables of contents of the last 2 years of *The Journal of Bone & Joint Surgery*, *The American Journal of Sports Medicine*, *Clinical Orthopaedics and Related Research*, *Arthroscopy*, and *Knee Surgery, Sports Traumatology, Arthroscopy* were manually searched as well. The librarian at our institution was consulted with regard to the search algorithm. Finally, authors and study data were cross-checked to prevent data duplication, and longer-term data were preferentially included.

TABLE II Demographics

Variable	Repair		No Repair		P Value
	No. of Cohorts*	Value†	No. of Studies*	Value†	
Total no. of patients/shoulders	32	1,294	13	289	—
Age	30	58.6 yr (56.4 to 60.8 yr)	13	56.5 yr (52.7 to 60.4 yr)	0.36
Male sex	32	66.9% (61.2% to 72.3%)	13	67.6% (60% to 74.8%)	0.88
Dominant side	10	70.3% (59.8% to 80.8%)	6	67% (49.1% to 85%)	0.76
Duration of follow-up	25	9.6 yr (8.6 to 10.7 yr)	12	14.9 yr (11.5 to 18.3 yr)	0.004

\*The values are given as the number of studies in which the value was reported. †The values are reported as the estimated mean, with the 95% CI in parentheses.

TABLE III Level of Evidence, Treatment, Sample Size, and Tear Size for Each Cohort

Study	Level of Evidence	Treatment	Sample Size (no. of patients/shoulders)	Tear Size
Bell et al. <sup>58</sup> (2013)	IV	Repair	49	Large
Bidwai et al. <sup>59</sup> (2016)	I	No repair	15	Medium
Bidwai et al. <sup>59</sup> (2016)	I	Repair	18	Medium
Björnsson et al. <sup>60</sup> (2010)	III	No repair	10	Partial
Björnsson et al. <sup>60</sup> (2010)	III	No repair	3	Full
Cuff et al. <sup>61</sup> (2016)	III	Repair	28	Massive
Denard et al. <sup>62</sup> (2012)	III	Repair	62	Massive
Denard et al. <sup>62</sup> (2012)	III	Repair	45	Massive
Dodson et al. <sup>63</sup> (2010)	IV	Repair	15	Large
Galatz et al. <sup>64</sup> (2001)	IV	Repair	33	Large
Goutallier et al. <sup>25</sup> (2009)	III	Repair	30	Large
Gulotta et al. <sup>24</sup> (2011)	II	Repair	106	Large
Inderhaug et al. <sup>65</sup> (2017)	IV	Repair	147	Massive
Jaeger et al. <sup>66</sup> (2016)	IV	No repair	22	Partial
Jaeger et al. <sup>66</sup> (2016)	IV	No repair	17	Full
Jaeger et al. <sup>66</sup> (2016)	IV	No repair	17	Rotator cuff tear arthropathy
Kartus et al. <sup>41</sup> (2006)	IV	No repair	26	Partial
Kijima et al. <sup>67</sup> (2012)	II	No repair	43	Full
Kluger et al. <sup>51</sup> (2011)	III	Repair	72	Large
Kluger et al. <sup>51</sup> (2011)	III	Repair	35	Large
Lucena et al. <sup>68</sup> (2015)	III	Repair	25	Medium
Lucena et al. <sup>68</sup> (2015)	III	Repair	25	Medium
Marrero et al. <sup>69</sup> (2011)	IV	Repair	24	Medium
Miyazaki et al. <sup>23</sup> (2015)	III	Repair	35	Massive
Moosmayer et al. <sup>4</sup> (2014)	I	No repair	39	Small
Moosmayer et al. <sup>4</sup> (2014)	I	Repair	52	Medium
Moosmayer et al. <sup>4</sup> (2014)	I	Repair	12	Medium
Nich et al. <sup>70</sup> (2009)	IV	Repair	33	Medium
Nich et al. <sup>70</sup> (2009)	IV	Repair	4	Medium
Norlin et al. <sup>71</sup> (2008)	IV	Repair	89	Tendinosis
Norlin et al. <sup>71</sup> (2008)	IV	Repair	45	Partial
Norlin et al. <sup>71</sup> (2008)	IV	Repair	5	Partial
Norlin et al. <sup>71</sup> (2008)	IV	Repair	12	Small
Norlin et al. <sup>71</sup> (2008)	IV	Repair	11	Medium
Paxton et al. <sup>72</sup> (2013)	IV	Repair	15	Massive
Porcellini et al. <sup>73</sup> (2011)	IV	Repair	67	Massive
Ranebo et al. <sup>42</sup> (2017)	IV	No repair	24	Full
Ranebo et al. <sup>42</sup> (2017)	IV	No repair	45	Partial
Saraswat et al. <sup>14</sup> (2015)	II	Repair	59	Medium
Sperling et al. <sup>74</sup> (2004)	IV	Repair	29	Large
Stephens et al. <sup>75</sup> (1998)	IV	No repair	11	Partial
Stephens et al. <sup>75</sup> (1998)	IV	No repair	17	Complete
Stuart et al. <sup>76</sup> (2013)	IV	Repair	15	Partial
Zandi et al. <sup>77</sup> (2006)	IV	Repair	74	Medium
Zumstein et al. <sup>26</sup> (2008)	IV	Repair	23	Massive

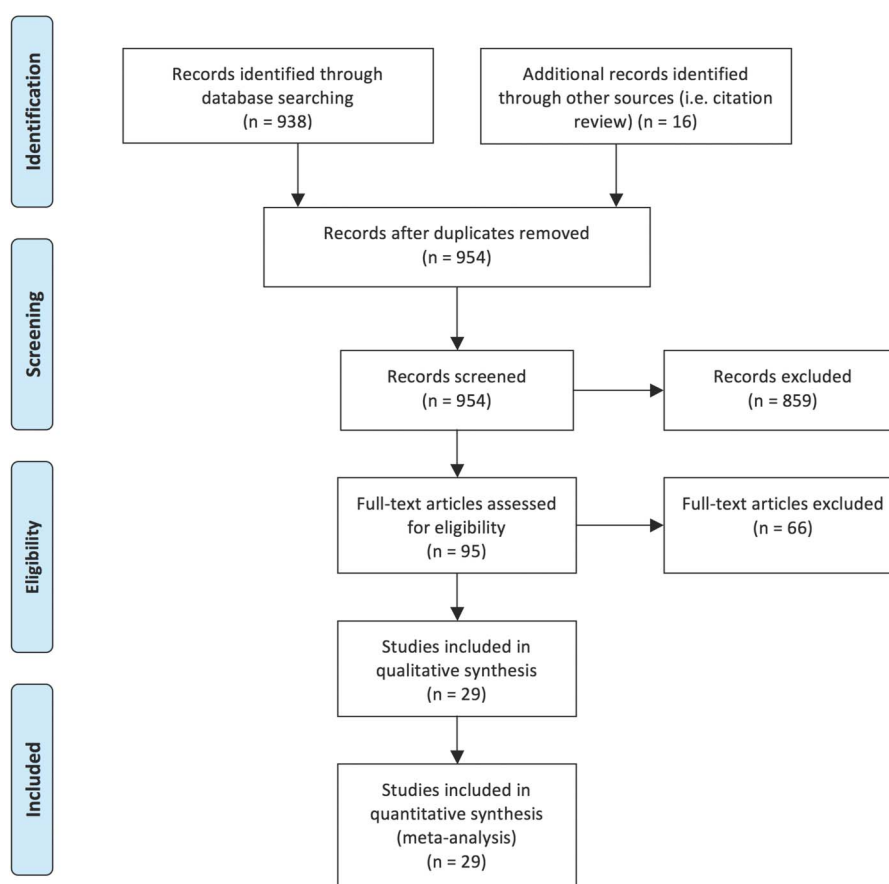


Fig. 1 PRISMA diagram showing the result of application of the study algorithm to the number of studies included, with the number of studies removed with application of each exclusion criterion displayed.

We adhered to the Preferred Reporting Items for Systematic reviews and Meta-analyses (PRISMA) guidelines<sup>25</sup>. Studies with levels of evidence ranging from I to IV were included. Studies were divided into those in which a repair of the rotator cuff was performed and those in which no repair was performed. The no-repair group included nonoperative treatments such as physical therapy and steroid injections, surgical subacromial decompression without repair, and surgical debridement with repair. These no-repair treatments were combined as the purpose of the

study was to determine whether repair alters natural history. Repair techniques, including open and arthroscopic approaches and single-row and double-row techniques, were combined to allow comparison. Two authors were involved in the decision process regarding the inclusion and exclusion of studies.

#### Data Collection

The study-related data that were collected included first author, year of publication, journal, level of evidence, number of

TABLE IV Distribution of Pre-Treatment Tear Types in No-Repair and Repair Cohorts

Tear Type	No-Repair Cohort (N = 289)	Repair Cohort (N = 1,294)
Partial	114 (39%)	154 (12%)
Small	39 (13%)	12 (1%)
Medium	15 (5%)	337 (26%)
Large	0 (0%)	369 (29%)
Massive	0 (0%)	422 (33%)
Full	104 (36%)	0 (0%)
Rotator cuff tear arthropathy	17 (6%)	0 (0%)

**TABLE V Effect of Repair (Versus No Repair) on Primary Outcomes with Adjustment for Covariates\***

Outcome	No. of Cohorts	Coefficient (95% CI)†	P Value
Percent requiring additional surgery	34	-0.095 (-0.17 to -0.021)	0.012
Percent with subsequent increase in tear size	22	0.529 (-0.693 to 1.751)	0.4
Constant score at latest follow-up	14	21.197 (-20.01 to 62.403)	0.31

\*Age, sex, duration of follow-up, and pre-treatment tear size. †Coefficients can be interpreted as estimated mean differences between groups after adjustment for covariates.

patients, minimum duration of follow-up, mean duration of follow-up, and treatment technique used (Table I). The demographic data that were collected included the number of patients in whom the dominant side was affected and the duration of symptoms before treatment (Table II). The clinical data that were collected (both preoperatively and at the time of the latest follow-up) included the number of patients who required further surgery, the number of patients with a documented increase in tear size, abduction strength, range of motion; absolute Constant score<sup>27</sup>, American Shoulder and Elbow Surgeons (ASES) score<sup>28</sup>, Disabilities of the Arm, Shoulder and Hand (DASH) and QuickDASH (an abbreviated version of the DASH) scores<sup>29</sup>, visual analogue scale (VAS) score for pain, University of California Los Angeles (UCLA) score<sup>30</sup>, Simple Shoulder Test (SST) scores<sup>31</sup>, and Western Ontario Rotator Cuff score<sup>32</sup>. The radiographic data that were collected included the number of patients at each radiographic Hamada stage<sup>12</sup>, the number of patients at each Goutallier fatty infiltration stage<sup>33</sup>, and acromiohumeral distance<sup>34</sup>. All strength measurements were converted to kilograms from pounds and newtons. The tear sizes before treatment and at the time of the latest follow-up were also collected as reported in each study on the basis of either magnetic resonance imaging (MRI) or ultrasound. All tear sizes were converted to square millimeters. When the length and width rather than the area were stated, these two 1-dimensional measurements were combined to calculate tear size, with the assumption being that the tears were rectangular, and when only a single dimension was stated, it was assumed to represent both the length and the width of a square tear; such calculations were necessary only in 3 of the 23

included repair cohorts. In addition, each cohort was classified as including partial, small, medium, large, or massive tendon tears (according to the authors' description of the cohorts or the measurements included in the studies) with use of the Cofield system<sup>35</sup>. Studies that did not describe tear size are classified as "full" and "rotator cuff tear arthropathy" as described by the authors of these studies. When possible, cohorts were split into multiple parts to allow for finer definitions of preoperative tear size; i.e., a study with both small and medium tear cohorts in which outcomes were reported for both cohorts would be split into 2 cohorts (1 containing small tears and 1 containing medium tears) for the purposes of our analysis (Table III). Studies that included patients with tendinosis with no discrete tear at the time of inclusion were excluded. Study quality was graded with use of the Coleman methodology score<sup>36</sup>.

#### Statistical Analysis

Study characteristics, including level of evidence, surgical approach (arthroscopic or open), and Coleman methodology score were summarized as the count (and percentage) or the mean and the standard deviation and were compared between repair and no-repair cohorts with use of the Fisher exact test or t test as appropriate. Age, male percentage, dominant-side percentage, and number of years of follow-up were pooled across studies with use of a random-effects model with inverse variance weighting. A chi-square Q test for heterogeneity was used to test for differences between repair and no-repair groups. Mixed-effects meta-regression models were used to compare the repair and no-repair groups in terms of the

**TABLE VI Effect of Repair (Versus No Repair) on Secondary Outcomes**

Outcome at Latest Follow-up	No. of Cohorts	Coefficient* (95% CI)	P Value
ASES score	10	1.67 (-17.21 to 20.55)	0.86
VAS pain score	8	1 (-25.23 to 27.23)	0.94
Elevation	9	-14.87° (-49.31° to 19.57°)	0.4
Elevation strength	6	1.13 kg (-5.17 to 7.43 kg)	0.72
Tear size	6	-967.37 mm <sup>2</sup> (-1,163.89 to -770.84 mm <sup>2</sup> )	<0.001

\*Coefficients can be interpreted as estimated mean differences between groups after adjustment for covariates.

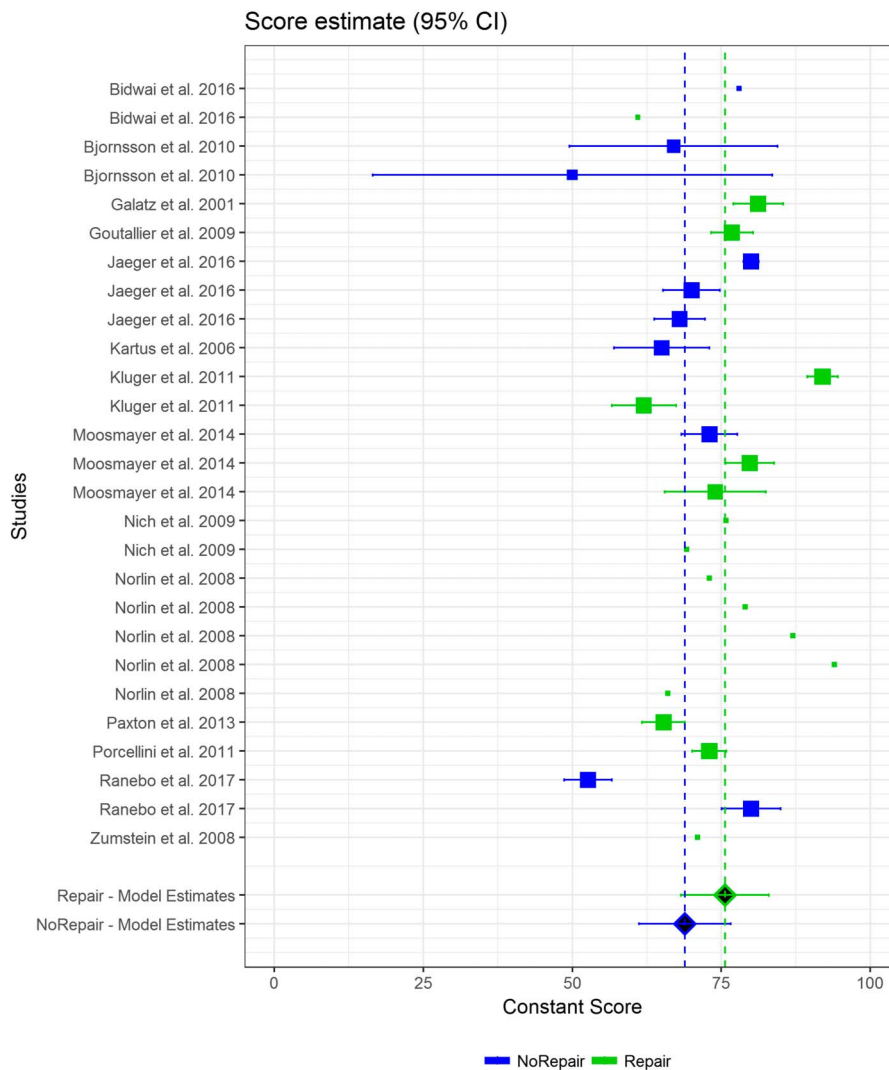


Fig. 2

Forest plot showing the final follow-up Constant scores for the repair (green) and no-repair (blue) cohorts. The horizontal bars show the 95% CIs for individual cohorts. The vertical dashed lines show the grand means.

percentage of patients or shoulders requiring further surgery, the percentage with an increase in tear size or recurrence of a defect, and the post-treatment operative Constant score, with adjustment for age, male percentage, number of years of follow-up, and preoperative tear size. Mixed-effects meta-regression also was used to compare the repair and no-repair groups with regard to postoperative ASES, VAS pain score, elevation, elevation strength, and tear size (while controlling for preoperative measures). Meta-analysis and meta-regression were conducted with use of the R package version 3.4 for meta-analysis<sup>37</sup>. Studies in which results were presented as summaries for different subgroups such as repair status or treatment type were included in the analysis as separate studies. If variance or standard deviation was not given, standard deviation was calculated from the standard error, 95% confidence interval (CI), or range<sup>38</sup>, as available. The level of significance was set at  $p < 0.05$ , and all tests were 2-tailed.

## Results

The initial search revealed 938 abstracts. After the application of our study-selection algorithm, 29 studies remained (Fig. 1); of those, 8 evaluated the outcomes of treatment without repair and 23 evaluated the outcomes of repair. The studies included 2 randomized clinical trials, 3 prospective cohort series, 7 retrospective cohort series, and 17 retrospective case series. There were no significant differences in Coleman methodology score between the individual repair cohorts ( $p = 0.8$ ). The overall repair group that was assessed in the present study included a total of 1,294 patients with a mean duration of follow-up of 9.6 years (95% CI, 8.6 to 10.7 years) (Table II). Of the patients in the repair group, 722 (56%) underwent an arthroscopic repair and 572 (44%) underwent open repair. Of the patients in the repair group, 45 (3.5%) were managed with a double-row technique, 461 (35.6%) were managed with a single-row technique, 505 (39.0%) were

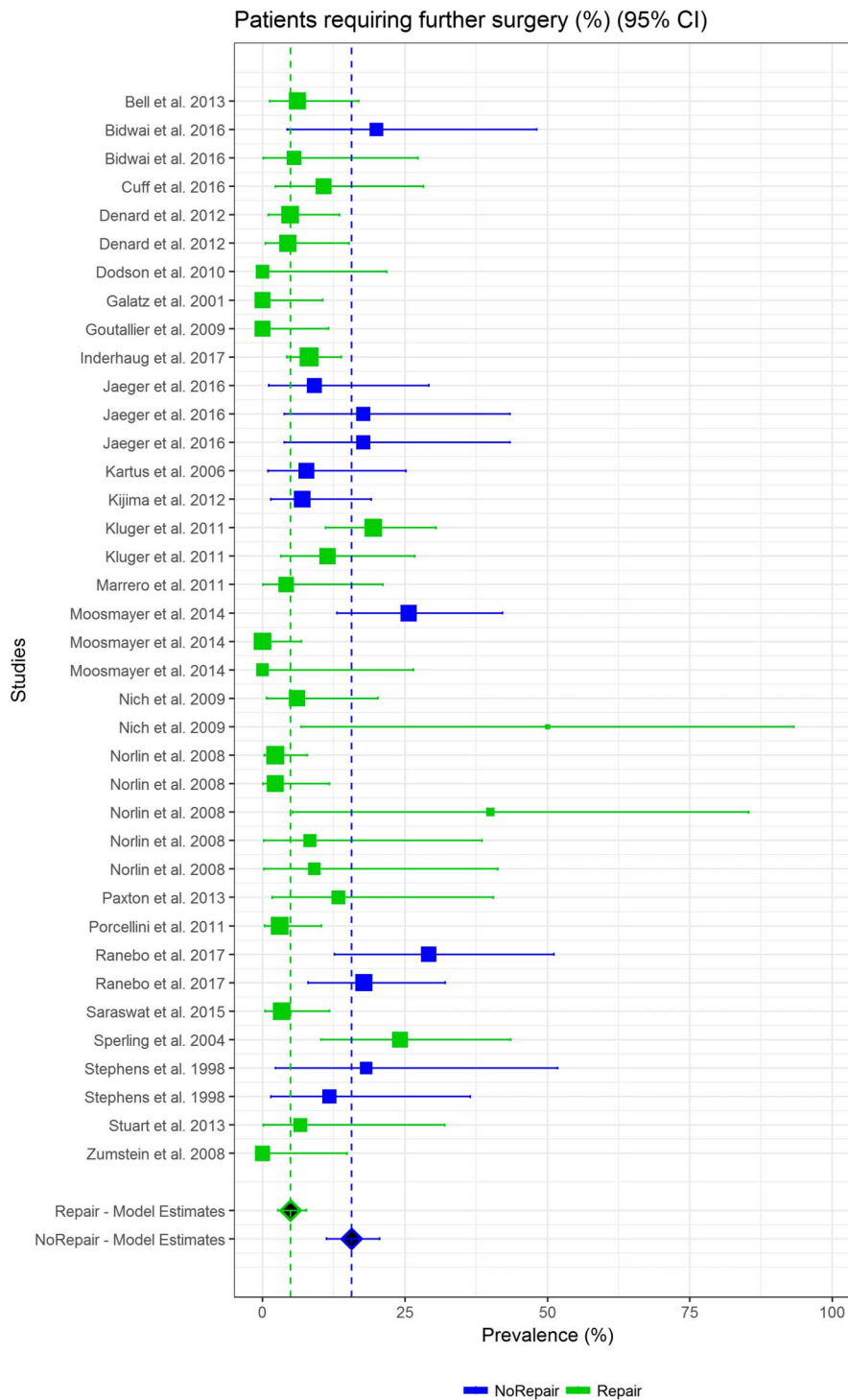


Fig. 3 Forest plot showing the mean percentage of patients requiring additional surgery for the repair (green) and no-repair (blue) cohorts. The horizontal bars show the 95% CIs for individual cohorts. The vertical dashed lines show the grand means.

managed with a transosseous technique, and 283 (21.9%) were managed with an unspecified technique. The no-repair group included a total of 289 patients with a mean duration of follow-up of 14.9 years (95% CI, 11.5 to 18.3 years). After the

studies were split into tear-size and treatment cohorts, there were 32 individual cohorts within the overall repair group and 13 individual cohorts within the overall no-repair group. The repair and no-repair cohorts did not differ with respect to age



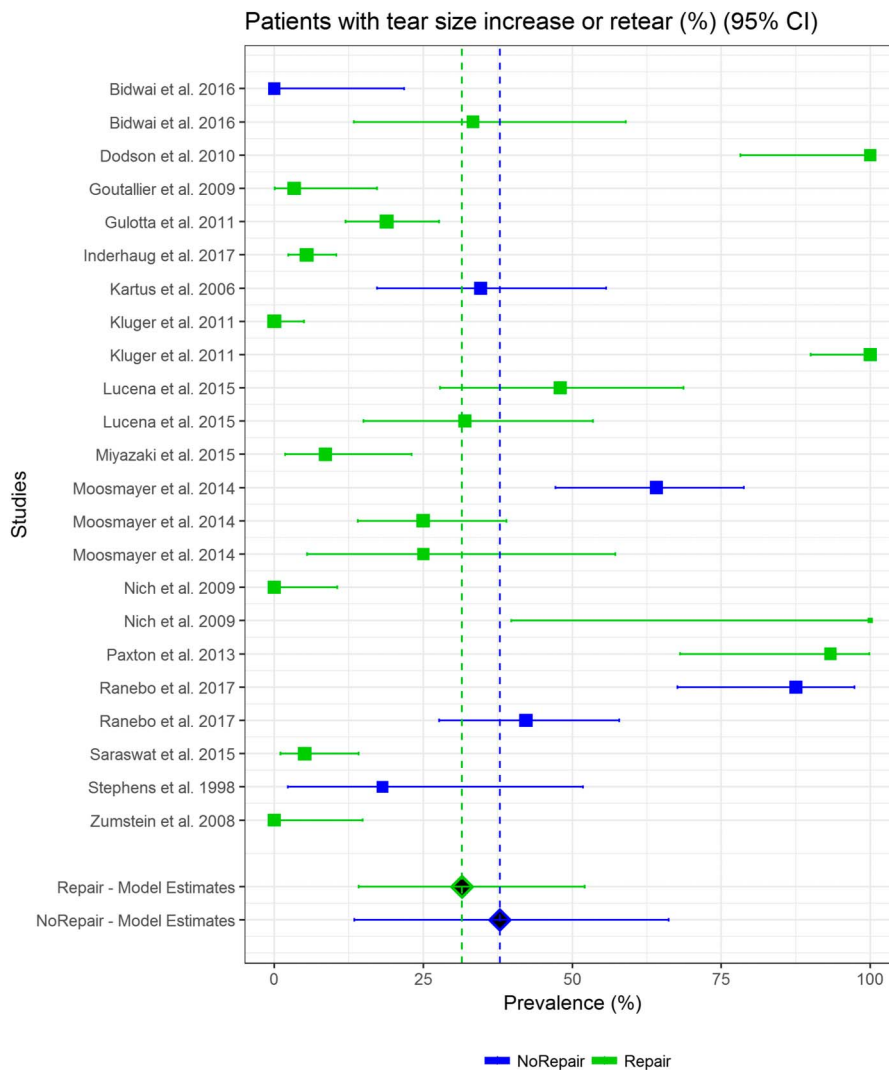


Fig. 4

Forest plot showing the percentage of patients sustaining either a recurrent defect or an enlargement of the tear from pre-treatment to the latest follow-up for the repair (green) and no-repair (blue) cohorts. The horizontal bars show the 95% CIs for individual cohorts. The vertical dashed lines show the grand means.

( $p = 0.36$ ), sex ( $p = 0.88$ ), or the percentage of patients in whom the tear was on the dominant side ( $p = 0.76$ ) (Table II). The duration of follow-up was significantly longer in the no-repair group ( $p = 0.004$ ). The baseline tear size was significantly larger in the repair group ( $p = 0.014$ ) (Table IV). Fewer than 3 repair studies and fewer than 3 no-repair studies evaluated pre-treatment strength, pre-treatment active forward elevation, pre-treatment Constant score, pre-treatment ASES score, pre-treatment VAS score, and radiographic outcomes, and thus no analyses were conducted on these variables.

There were no differences between the groups in terms of physical examination findings or strength at the time of the latest follow-up. Specifically, there were no differences in terms of elevation range of motion ( $p = 0.4$ ) or elevation strength ( $p = 0.72$ ) (Table V). There also were no differences between the groups in terms of functional and patient-based outcomes,

including the ASES score ( $p = 0.86$ ) or VAS pain score ( $p = 0.94$ ) at the time of the latest follow up (Table VI). In addition, the final Constant score did not differ between the groups after adjusting for age, sex, duration of follow-up, and tear size ( $p = 0.31$ ) (Fig. 2 and Table VI).

The percentage of patients requiring additional surgery was significantly higher in the no-repair group after adjustment for age, sex, duration of follow-up, and tear size (9.5% higher in estimated means between groups [95% CI, 2.1% to 17%];  $p = 0.012$ ) (Fig. 3 and Table VI). The percentage of patients with a recurrent defect in the repair group did not differ from the percentage of patients with an increase in tear size in the no-repair group after adjusting for age, sex, duration of follow-up, and tear size ( $p = 0.4$ ) (Fig. 4 and Table V). The final tear size was significantly larger in the no-repair group than the repair group (967 mm<sup>2</sup> greater [95% CI, 771 to 1,164 mm<sup>2</sup> greater],  $p < 0.001$ ) (Table VI).



## Discussion

Rotator cuff tears may increase in size and can lead to pain, disability, and ultimately pseudoparalysis and rotator cuff tear arthropathy<sup>1-5,9-12</sup>. Because rotator cuff repair can result in a continuous bone-tendon interface in the long term<sup>14,16,21-26</sup>, rotator cuff repair may be able to forestall this natural history. The purpose of the present study was to conduct a systematic review of all published clinical studies with a minimum duration of follow-up of 5 years after rotator cuff repair and/or nonoperative treatment of rotator cuff disease in order to evaluate (1) strength and range of motion on physical examination, (2) functional and patient-based outcomes, (3) the need for future surgical intervention, (4) the likelihood of future tear progression or recurrence, and (5) final tear size.

The present study demonstrated no differences between rotator cuff repair and no repair with respect to strength and range of motion. This finding is in concordance with the 3 randomized clinical trials that have been performed to date, each of which demonstrated no differences between rotator cuff repair and no repair with respect to strength and range of motion<sup>4,19,20</sup>. Following rotator cuff repair, shoulders in which an intact tendon is achieved have greater strength than those in which an intact tendon is not achieved<sup>39</sup>. In addition, strength has been correlated with tear size<sup>5,40</sup>. Given our finding that final tear size was larger in the no-repair group as compared with the repair group, improved strength would be expected in the repair group, but this effect may have been obscured by heterogeneity in strength measurement between studies.

We found that, compared with no repair, rotator cuff repair did not improve outcomes as measured with the Constant score even after adjustment for age, sex, duration of follow-up, and tear size. These findings are roughly congruent with those of the 3 randomized clinical trials that have been conducted to date<sup>4,19,20</sup>. One of those studies demonstrated no clinically or statistically significant differences between groups<sup>19</sup>, 1 demonstrated a statistically but not clinically significantly better outcome in terms of the Constant score for rotator cuff repair resulting in an intact tendon<sup>4</sup>, and 1 demonstrated both a clinically and statistically significantly better outcome in terms of the Constant score for rotator cuff repair resulting in an intact tendon<sup>20</sup>.

Our study demonstrated that rotator cuff repair appears to protect the shoulder from the need for future operative intervention after adjusting for age, sex, duration of follow-up, and tear size. As tear size increases following nonoperative treatment, some patients may become increasingly symptomatic and may be considered for arthroscopic debridement, subacromial decompression, biceps tenotomy or tenodesis, rotator cuff repair, tendon transfer, or ultimately reverse total shoulder arthroplasty. In addition, tear size<sup>4,41,42</sup>, age, and muscular atrophy<sup>33,42</sup> all continue to increase following nonoperative treatment, thereby decreasing the likelihood of achieving an intact tendon<sup>21,22,33,38,43-55</sup>. As a result, nonoperative treatment both increases the likelihood of future surgery and may decrease the likelihood of success if that surgery is a repair.

The present study indicates that rotator cuff repair does not decrease the likelihood of sustaining a future tear after adjusting for age, sex, duration of follow-up, and tear size but does decrease final tear size. Rotator cuff repair does not alter the underlying tendon biology that causes rotator cuff tearing, and therefore the likelihood of a recurrent defect after rotator cuff repair may be similar to the likelihood of tear progression with nonoperative treatment. A "recurrent defect" after rotator cuff repair thus may be understood not as a surgical failure but instead as a continuation of the underlying, unaltered, biological degeneration that leads to rotator cuff pathology. However, our study demonstrated that final tear size was significantly smaller after rotator cuff repair than after treatment without repair.

Our study has several limitations. First, the data were drawn from studies with different designs, and thus heterogeneity between studies limits the conclusions that can be drawn. In addition, surgical repairs and postoperative rehabilitation have changed between the publication of the first study in 2001 and that of the most recent study in 2017. Second, as with any meta-analysis, the quality of the conclusions that can be drawn is limited by the quality of the original data, which are drawn from studies of varying levels of evidence. Third, the included studies were limited to those published in English, which may introduce bias. However, each of these limitations affect both the repair and no-repair cohorts, which may mitigate their influence on our results. Fourth, a variety of repair and no-repair treatment methods were included. There is continuing debate as to whether single-row or double-row repair provides superior outcomes or a higher likelihood of an intact tendon<sup>56</sup>. Fifth, there are certainly other variables that would have been valuable to compare, such as radiographic progression toward rotator cuff tear arthropathy as indicated by Hamada stage, muscular atrophy or tendon quality at baseline and at the latest follow-up, and which specific subsequent procedures were necessary. Unfortunately, these details were not available in the included studies and thus we could not analyze them. Sixth, tear size was measured on both MRI and ultrasound scans. Finally, although no difference existed in baseline demographic data between the repair and no-repair cohorts, unmeasured residual bias likely existed between the cohorts. For instance, the baseline tear size was larger in the repair group. To mitigate this effect, we controlled for tear size in our analyses of the primary outcomes. However, for many other variables (strength, motion, Constant score, etc.), insufficient evidence existed within the pre-treatment data to allow comparison. We were able to control for age, sex, and tear size, which are the 3 variables that have been shown to most strongly correlate with outcome<sup>57</sup>. Only a randomized controlled trial will be able to overcome this limitation.

In conclusion, at intermediate to long-term follow-up, rotator cuff repair was associated with decreased final tear size and decreased need for future surgery but was not associated with higher final standardized outcomes after adjusting for age, sex, duration of follow-up, and tear size. The likelihood of a recurrent defect after rotator cuff repair did not differ from the

likelihood of tear extension with nonoperative treatment, and thus rotator cuff repair may not alter natural history. ■

Peter N. Chalmers, MD<sup>1</sup>  
Hunter Ross, BA<sup>1</sup>  
Erin Granger, MPH<sup>1</sup>  
Angela P. Presson, PhD<sup>1</sup>

Chong Zhang, MS<sup>1</sup>  
Robert Z. Tashjian, MD<sup>1</sup>

<sup>1</sup>Departments of Orthopaedic Surgery (P.N.C., H.R., E.G., and R.Z.T.), and Epidemiology (A.P.P. and C.Z.), University of Utah, Salt Lake City, Utah

E-mail address for P.N. Chalmers: p.n.chalmers@gmail.com

ORCID iD for P.N. Chalmers: [0000-0002-1275-0285](https://orcid.org/0000-0002-1275-0285)

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