# Nonarthroplasty Surgical Treatment Options for Massive, Irreparable Rotator Cuff Tears

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Massive, irreparable rotator cuff tears (MIRCTs) provide a significant dilemma for orthopaedic surgeons. One treatment option for MIRCTs is reverse total shoulder arthroplasty. However, other methods of treating these massive tears have been developed. A search of the current literature on nonoperative management, arthroscopic debridement, partial repair, superior capsular reconstruction (SCR), graft interposition, balloon spacer arthroplasty, trapezius transfer, and latissimus dorsi transfer for MIRCTs was performed. Studies that described each surgical technique and reported on clinical outcomes were included in this review. Arthroscopic debridement may provide pain relief by removing damaged rotator cuff tissue, but no functional repair is performed. Partial repair has been suggested as a technique to restore shoulder functionality by repairing as much of the rotator cuff tendon as possible. This technique has demonstrated improved clinical outcomes but also fails at a significantly high rate. SCR has recently gained interest as a method to prohibit superior humeral head translation and has been met with encouraging early clinical outcomes. Graft interposition bridges the gap between the retracted tendon and humerus. Balloon spacer arthroplasty has also been recently proposed and acts to prohibit humeral head migration by placing a biodegradable saline-filled spacer between the humeral head and acromion; it has been shown to provide good clinical outcomes. Both trapezius and latissimus dorsi transfer techniques involve transferring the tendon of these respective muscles to the greater tuberosity of the humerus; these 2 techniques have shown promising restoration in shoulder function, especially in a younger, active population. Arthroscopic debridement, partial repair, SCR, graft interposition, balloon spacer arthroplasty, trapezius transfer, and latissimus dorsi transfer have all been shown to improve clinical outcomes for patients presenting with MIRCTs. Randomized controlled trials are necessary for confirming the efficacy of these procedures and to determine when each is indicated based on specific patient and anatomic factors.

**Keywords:** massive, irreparable rotator cuff tear; superior capsular reconstruction; graft interposition; balloon spacer arthroplasty; tendon transfer

The Orthopaedic Journal of Sports Medicine, 6(11), 2325967118805385 DOI: 10.1177/2325967118805385 © The Author(s) 2018 Massive, irreparable rotator cuff tears (MIRCTs) provide a significant dilemma for orthopaedic surgeons. These lesions are characterized by rotator cuff tears >3 cm with advanced fatty infiltration of the rotator cuff tendons, a reduced acromiohumeral interval, significant tendon retraction, and the presence of poor-quality tissue.<sup>36,62</sup> Historically, limited reliable options have existed for the treatment of these tears, and although multiple options have been described, many have been insufficient to meet patients' demands (eg, isolated arthroscopic debridement), are technically demanding and result in difficult rehabilitation (eg, tendon transfers), and have generally been met with underwhelming clinical success as a whole. One of the recent popular options for MIRCTs has been reverse total shoulder arthroplasty (rTSA), as it allows the deltoid muscle to take over the function of the irreparable rotator cuff muscles and has shown promising clinical results in multiple studies when employed for this indication.<sup>23,41,62,63</sup> This technique has become the treatment of choice among surgeons for patients with MIRCTs complicated by significant glenohumeral arthritis, anterosuperior escape, and/or

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pseudoparalysis.<sup>62</sup> However, complications after rTSA, although improving, have been reported to occur at a rate of 20% to 50% in both the recent and past literature.<sup>23,41,63</sup> Additionally, there are concerns regarding the longevity of these implants, with a revision rate of 10% to 33% and increasing complication rates with each revision.<sup>3,4,41,63</sup> Therefore, surgeons have attempted to develop novel surgical treatment methods for massive tears. It should be noted that inferior results of rTSA have been demonstrated in patients who have undergone previous shoulder surgery.<sup>20</sup> Therefore, while nonarthroplasty procedures may be useful in select patients, they may jeopardize the results of rTSA if they fail.

The purpose of this review was to provide an overview of these surgical techniques for the treatment of MIRCTs, including arthroscopic debridement, partial repair, superior capsular reconstruction (SCR), graft interposition, balloon spacer arthroplasty, trapezius transfer, and latissimus dorsi transfer, as well as to highlight the current literature on clinical outcomes after these procedures.

# SEARCH STRATEGY

A search through May 2018 was performed using various combinations of the following keywords/phrases: "massive," "irreparable," "rotator cuff tear," "surgical," "operative," "non-operative," "arthroscopic debridement," "partial repair," "superior capsular reconstruction," "graft interposition," "balloon spacer arthroplasty," "trapezius transfer," and "latissimus dorsi transfer." Surgical technique descriptions and clinical studies that described and reported on the clinical outcomes of arthroscopic debridement, partial repair, SCR, graft interposition, balloon spacer arthroplasty, trapezius transfer, and latissimus dorsi transfer for the treatment of MIRCTs were included in this review.

# NONOPERATIVE MANAGEMENT

Because of the challenges that an MIRCT presents surgically, it is recommended to have patients attempt nonoperative treatment before surgical intervention. Nonoperative management options typically include activity modification, steroid injections, and physical therapy.<sup>44</sup> Studies have shown that nonoperative management and physical therapy with an emphasis on rehabilitation of the anterior deltoid can improve clinical outcomes in patients with MIRCTs (Table 1). $^{44,64}$  Dunn et al $^{15}$  examined a group of patients who underwent nonoperative treatment for MIRCTs to define specific patient factors as predictors of treatment failure and the need for future surgery. Predictors of surgical intervention included low expectations of physical therapy, a high activity level, and not smoking. Whereas some studies<sup>44</sup> have suggested that patients with chronic MIRCTs or significant pain due to MIRCTs tend to have inferior outcomes with nonoperative management, Dunn et al<sup>15</sup> found no correlation between patient symptoms or anatomic features of the rotator cuff tear and the

need for future surgery. However, additional studies have suggested that nonoperative management can cause significant degenerative structural joint changes and progression of the tear, potentially complicating any future surgical intervention if indicated.<sup>64</sup>

# ARTHROSCOPIC DEBRIDEMENT

Arthroscopic debridement has been described as a surgical treatment option for patients with MIRCTs.<sup>31,55</sup> In 1995, Rockwood et al<sup>55</sup> described the use of this procedure in a group of patients treated from 1976 to 1988. The technique involves debridement of the torn rotator cuff by removing avascular or unstable tissue that could be caught or impinged between the humeral head and acromion during shoulder flexion.<sup>31,55</sup> Additionally, surgeons may elect to perform bursectomy of the subacromial bursa, scar tissue removal, acromioplasty, subacromial decompression, and biceps tenotomy or tenodesis in conjunction with rotator cuff debridement.<sup>31</sup> The goal of this intervention is to provide pain relief by removing the sources of mechanical irritation or inflammation. Some believe that this procedure should only be used as a salvage option for patients who seek pain relief, as no functional repair is performed.<sup>31</sup> Therefore, arthroscopic debridement is indicated in patients with an irreparable rotator cuff tear who have minimal to mild osteoarthritis with a chief complaint of pain after the failure of conservative treatment.<sup>31</sup>

Multiple studies have reported promising clinical outcomes after arthroscopic debridement of MIRCTs (Table 1).<sup>3,21,32,38,55,60</sup> Berth et al<sup>3</sup> evaluated a group of 21 patients who underwent arthroscopic debridement with subacromial bursectomy and decompression for the treatment of MIRCTs. At a mean follow-up of 16.8 months, there was a significant improvement in the mean Constant-Murley (CM) score (29.9 to 40.7; P < .01) and the mean Disabilities of the Arm, Shoulder and Hand (DASH) score (69.5 to 35.3; P < .01). However, this study demonstrated no significant change in abduction (93.5° to  $103.5^{\circ}$ ; P = .074) or external rotation (40.5° to 42.7°; P = .157) from preoperatively to final follow-up, despite a significant increase in internal rotation (49.5° to 71.6°; P < .01). Franceschi et al<sup>21</sup> performed a similar study with a group of 34 patients undergoing arthroscopic debridement, subacromial bursectomy, acromioplasty, and in select cases biceps tenotomy. The authors found a significant improvement in the mean University of California, Los Angeles (UCLA) score (7.6 to 21.4; P < .0001) and visual analog scale (VAS) score for pain (6.7 to 1.2; P < .0001) at a mean follow-up of 93.6 months. Range of motion also significantly improved over the study period in terms of forward flexion (104.1° to 132.0°; P < .001), external rotation (42.9° to 48.0°; P < .001), and internal rotation (37.8° to 46.7°; P < .001).

Many of the studies examining the clinical outcomes of arthroscopic debridement of MIRCTs also assessed preoperative factors and their influence on clinical outcomes.<sup>3,21,32,38,60</sup> Liem et al<sup>38</sup> found male patients to have significantly improved clinical outcomes relative to those of female patients (P = .008), although no significant effect

			Clinical Outcomes		
Study	No. of Patients	Mean Follow-up, mo	Preoperative	Postoperative	P Value
Nonoperative management	10	40	CM ND	CIM CO	OM ND
Zingg et al <sup><math>31</math></sup> (2007)	19	48	CM: NR	CM: 69	CM: NR
			SSV: NK	SSV: 68%	55V: NK
			FF: 115°	FF: 139°	FF: .047
			AB: 118 <sup>-</sup>	AB: 139 <sup>-</sup>	AB: .070
			IK: 76° FD: 44°	IK: 67° ED: 49°	IK: .054
Arthroscopic debridement			ER: 44	ER: 45	ER: .804
Borth at $n^3$ (2010)	91	16.8	CM: 20.0	CM: 40.7	$CM_{\rm C} < 01$
Dertil et al (2010)	21	10.8	DASH: 60 5	DASU: 25 2	$DASU_{1} < 01$
			AB: 02 50	AB: 102 50	$\Delta \mathbf{R}_{1}  0.74$
			AD. 95.5 FD: 40.5°	AD. 103.3 FD. 49.70	AD074 FD: 157
			ID: 40.5	ER. 42.7 ID: 71.6°	ER157 IP. < 01
Even eccepti et $al^{21}(9015)$	94	02.6	III. 49.5 LICI A. 7.6	III. 71.0 LICI A. 91 4	$\frac{110.}{1001} \times 0.001$
Franceschi et al (2013)	-04	95.0	UCLA: 7.0	UCLA: 21.4 VAS: 1.9	UCLA: < .0001
			VAS: 0.7 FF: 104 1º	VAS: 1.2 FF: 122.0°	VAS: < .0001
			FF: 104.1 FD: 49.0°	FF: 152.0 FD: 49.00	FF: < .001
			ER: 42.9 ID: 97.00	ER: 40.0 ID: 46.7°	ER. < .001
Howhere at $a^{32}$ (2016)	0.9	45.0	IN: 57.0 CM: 24	IN: 40.7 CM: 65	$CM_{\rm h} < 001$
Heuberer et al (2016)	20	45.0	CM: 54 SSV: 2507	CMI: 00 CCV. 7907	CM: < .001
				DACIL 02	DASU: < 001
Lign at $a1^{38}$ (2008)	91	47.0	DASH: 02	DASH: 23	DASH: $<.001$
Liem et al $(2008)$	31	47.0	ASES: 24.0	ASES: 09.8	ASES: < .001
$10^{-10}$	07		VAS: 7.8	VAS: 2.9	VAS: <.001
Veado and Rodrigues <sup>21</sup> (2015)	27	27.0	UCLA: 15	UCLA: 31	UCLA: NR
Classical repair	97	20.0	ACEC 40.0		
Chen et al $(2017)$	37	29.6	ASES: 46.0	ASES: 78.6	ASES: <.001
(1, 0) $(1, 1)$ $(1, 1)$ $(1, 1)$	00	71.1	VAS: 0.22	VAS: 1.01	VAS: <.001
Cuff et al (2016)	28	71.1	ASES: 40.0	ASES: 79.3	ASES: <.001
			SST: 5.6	SST: 9.1	SST: <.001
			VAS: 6.9	VAS: 1.9	VAS: <.001
			FF: 168°	FF: 154°	FF: .074
			ER: $38^{\circ}$	ER: 39 <sup>-</sup>	ER: >.99
$D_{1} = 11 + 10 + 10 + 16 (2007)$	60	12.0	IR: 84%	IR: 80%	IR: >.99
Duraide and Bair <sup>14</sup> (2005)	68	43.0	ASES: 41.0	ASES: 80.1	ASES: <.001
			FF: 114 <sup>-</sup>	FF: 154"	FF: NK
$(1,1,\dots,1,1)^{22}$ (2017)	00	04.0	ER: 44°	ER: $54^{\circ}$	ER: NR
Galasso et al (2017)	90	84.0	CM: 39.1	CIMI: 76.3	CM: <.001
			551: NK	SS1: 9.1	SST: NK
			$FF^{\circ}: 171^{\circ}$	FF": 174"	FF: .062
			$AB : 167^{\circ}$	$AB : 177^{\circ}$ EDC 010	AB: <.001
			$\mathbf{ER}^{*}: \mathbf{ZS}^{*}$	$EK^{2} 31^{\circ}$ $ID^{c} T7$	ER: $.022$
12	07	41.9		IR : II	1R: <.001
Kim et al $(2012)$	21	41.3	SS1: 0.1	SS1: 0.0 CM: 74 1	SS1: < .001
			UNI: 43.0		UM: < .001
$\mathbf{D}_{1} = 1 + 147 (2017)$	10	24.0	OCLA: 10.5	UCLA: 25.9	OCLA: <.001
Pandey et al. $(2017)$	13	24.0	USS: 17.8	055: 37.1 CM 70.9	OSS: .009
Shon et al <sup>59</sup> (2015)	01	10 5	CM: 43.1	UM: 70.8	CM: .01
	51	40.5	VAS: 0.13	VAS: 2.13 (1 y 1/u)	VAS: .001
			ASES: 41.97	ASES: 13.18	ASES: <.001
Superior equally recordent			10.6 1100	0.07	600.1166
Superior cupsular reconstruction Depend of $al^{13}$ (2012)	FO	100	ACTC. 49.0	1959. 77 F	ACTC - 001
Denaru et al (2018)	99	11.1	АЗĽЗ: 43.0 VAC. 5 0	АЗЦЭ: 11.0 VAS, 17	ASES: <.001
			VAD: 0.0	VAD: 1.1 SSV. 76.9	VAO; < .001
			557.30.U FF.1900	557.70.3 FF.150	55V. < .001
			FF: 130 FD: 960	FF: 100 FD: 450	FF: <.001
			ER: 50 ID: 1-2	ER: 40 ID: I 1	ER: $.000$ ID: $< 0.01$
			IU: T9	IU: L1	IU: <'001

TABLE 1 Summary of Clinical Studies on Massive, Irreparable Rotator Cuff Tears  $^a$ 

(continued)

TABLE	1	(continued)	)
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Study	No. of Patients		Clinical Outcomes		
		Mean Follow-up, mo	Preoperative	Postoperative	P Value
Lee and Min <sup>37</sup> (2018)	36	24.8	ASES: 50.3 CM: 56.3	ASES: 84.0 CM: 82.8	ASES: <.01 CM: .02
Mihata et al <sup>39</sup> (2013)	23	34.1	JOA: 48.3 ASES: 23.5	JOA: 92.6 ASES: 92.9	JOA: <.00001 ASES: <.00001
			UCLA: 9.9 FF: 84°	UCLA: 32.4 FF: 148°	UCLA: <.00001 FF: <.001
Pennington et al <sup>48</sup> (2018)	88	12	ER: 26° VAS: 4.0 ASES: 52	ER: 40° VAS: 1.5 ASES: 82	ER: <.01 VAS: .005 ASES: .005
			AB: 103°	AB: 159°	AB: .02
Graft interposition					
Audenart et $al^2$ (2006)	41	43	CM: 25.7	CM: 72.1	CM: <.001
Gupta et al <sup>30</sup> (2012)	24	36	ASES: 66.6	ASES: 88.7	ASES: .0003
			SF-12: 48.8	SF-12: 56.8	SF-12: .03
			VAS: 5.4	VAS: 0.9	VAS: .0002
			FF: 111.7°	FF: 157.3°	FF: .0002
			ER: $46.2^{\circ}$	ER: $65.1^{\circ}$	ER: .001
Noumann at $a^{145}(2017)$	60	50.2	AB: 105.0° VAS: 4.0	AB: 151.7 VAS: 1.0	AB: $.002$ VAS: < 001
Neumann et al (2017)	00	00.5	VAS: 4.0 ASES: NR	VAS: 1.0 ASES: 87.8	VAS: <.001
			FF: 140 7°	FF: 160 4°	FF < 001
			ER: 55.6°	ER: 70.1°	ER: .001
			IR: 52.0°	IR: 76.2°	IR: .001
Ranebo et al <sup>54</sup> (2018)	13	216	CM: NR	CM: 46	CM: NR
			WORC: NR	WORC: 59	WORC: NR
Venouziou et al <sup>61</sup> (2013)	14	30.2	VAS: 7.4	VAS: 1.7	VAS: .001
			ASES: 23.8	ASES: 72.3	ASES: .001
			$FF: 73.6^{\circ}$	FF: 129.3°	FF: .002
			AB: $67.5^{\circ}$	AB: 117.9°	AB: .002
			ER: $7.9^{\circ}$	ER: $43.2^{\circ}$	ER: .001
Balloon spacer arthroplasty					
Deranlot et al <sup>14</sup> $(2017)$	37	32.8	CM: 44.8	CM: 76.0	CM: <.001
			FF: 130°	FF: 160°	FF: .02
			AB: 100°	AB: 160°	AB: .01
C	15	10.0	ER: $30^{\circ}$	ER: $45^{\circ}$	ER: .0001
Gervasi et al <sup>24</sup> (2016)	15	12.0	CM: 31.9	CM: 69.8	CM: <.0001
$P_{1}^{1} = p_{1}^{1} (2017)$	4.4	19.0	ASES: 24.0	ASES: 76.0	ASES: < .0001
Flekaar et al (2017)	44	12.0	CM: 27.1	CM: 60.2	$CM_{\rm c} < 001$
Prot at $a1^{53}$ (2018)	99	14.4	$UCLA \cdot 10.9$	$UCL\Delta \cdot 15.9$	UCLA: $001$
11at et al (2010)	22	11.1	FF: 90°	FF: 106 5°	FF: 17
			ER: 34.1°	ER: 37.5°	ER: .48
			IR: L5	IR: L4	IR: .37
Senekovic et al <sup>57</sup> (2017)	24	60.0	CM: 34.2	CM: 67.4	CM: <.0001
Senekovic et al <sup>58</sup> (2013)	20	36.0	CM: 33.4	CM: 65.4	CM: <.0001
Trapezius transfer					
Elhassan et al <sup>19</sup> (2016)	33	47.0	SSV: $54\%$	SSV: 78%	$\mathrm{SSV:}<.01$
			DASH: 52	DASH: 18	DASH: <.01
			FF: 70°	FF: 120°	FF: <.01
			AB: 40°	AB: 90°	AB: <.01
<b>T</b> ,··· <b>T</b> ··· A			ER: $20^{\circ}$	ER: 50°	ER: <.01
Latissimus dorsi transfer	0.0	0.0.4	CM of f	OM COF	OM CONT
Castricini et al $(2016)$	86	36.4	CM: 35.5	CM: 69.5 CM: 74.0	CM: < 05
Castricini et al <sup></sup> (2014)	27	21	UMI: 30.0 FD: 990	UNI: 74.U FD: 200	OWI: < .05
			En: 23	EU: 90	<b>БИ: &lt;'0</b> 0

(continued)

	No. of Patients	Mean Follow-up, mo	Clinical Outcomes		
Study			Preoperative	Postoperative	P Value
El-Azab et al <sup>17</sup> (2015)	108	111.6	CM: 36.1 ASES: 30.1 VAS: 7.8 FF: 86.0° AB: 88.7°	CM: 62.0 ASES: 70.2 VAS: 2.4 FF: 133.5° AB: 127.4°	CM: <.0001 ASES: <.0001 VAS: <.0001 FF: <.0001 AB: <.0001
Gerber et al <sup>24</sup> (2013)	44	146.6	ER: 17.6° SSV: 29.0% CM: 47.3 FF: 118.0° AB: 112.1° ER: 17.9°	ER: 29.2° SSV: 70.1% CM: 63.5 FF: 132.4° AB: 122.6° ER: 32.5°	ER: <.0001 SSV: .0001 CM: <.0001 FF: .029 AB: .089 ER: .0001
Grimberg et al <sup>29</sup> (2015)	55	29.0	SSV: 26% CM: 37.0 FF: 134° AB: 67°	SSV: 71.1% CM: 65.4 FF: 157° AB: 92.5°	SSV: <.001 CM: <.001 FF: <.001 AB: <.001
Kanatli et al <sup>34</sup> (2017)	15	26.4	ER: 29° UCLA: 6.53 CM: 21.0 VAS: 7.47 FF: 58° AB: 51°	ER: 41.5° UCLA: 27.47 CM: 59.73 VAS: 2.47 FF: 130° AB: 129.7°	ER: <.001 UCLA: <.001 CM: <.001 VAS: <.001 FF: <.001 AB: <.001
Mun et al <sup>42</sup> (2018)	24	12	ER: 13.3° CM: 46 ASES: 40 VAS: 6 FF: 135° IR: L5 FB: 51°	ER: 32° CM: 69 ASES: 70 VAS: 2 FF: 166° IR: L1 FR: 68°	ER: <.001 CM: <.001 ASES: <.001 VAS: .006 FF: .016 IR: .010 FF: .062
Petricciolo et al <sup>50</sup> (2016)	33	35.7	CM: 34.6 VAS: 5 DASH: 49.7 FF: 138° ER: 7°	CM: 64.9 VAS: 1.4 DASH: 22.6 FF: 168° ER: 34°	CM: <.05 VAS: <.0001 DASH: <.001 FF: <.05 ER: <.05

TAF	BLE	1	(continu	ed)
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<sup>*a*</sup>AB, abduction; ASES, American Shoulder and Elbow Surgeons; CM, Constant-Murley; DASH, Disabilities of the Arm, Shoulder and Hand; ER, external rotation; FF, forward flexion; f/u, follow-up; IR, internal rotation; JOA, Japanese Orthopaedic Association; NR, not reported; OSS, Oxford Shoulder Score; SF-12, 12-Item Short Form Health Survey; SST, Simple Shoulder Test; SSV, Subjective Shoulder Value; UCLA, University of California, Los Angeles; VAS, visual analog scale for pain; WORC, Western Ontario Rotator Cuff Index.

<sup>b</sup>Values are presented for surgically repaired shoulder.

<sup>c</sup>Values are presented for contralateral shoulder.

was found based on age, tear size, or preoperative osteoarthritis. Franceschi et al<sup>21</sup> found that patients with a reduced preoperative acromiohumeral distance fared worse with regard to the UCLA score at final follow-up. The authors also found that high-demand manual laborers reported significantly worse clinical outcomes compared with nonmanual workers (P < .001). In contrast, 3 other studies<sup>3,32,60</sup> found no relation between age, sex, profession, or fatty infiltration status of the rotator cuff tendons and clinical outcomes.

Because functional repair is not performed with arthroscopic debridement of MIRCTs, revision rates are low. One study<sup>32</sup> reported a single case of revision to rTSA because of persistent pain and functional limitation. Berth et al<sup>3</sup> also reported a single case of revision to hemiarthroplasty because of the development of severe glenohumeral osteoarthritis. No other cases of failure or revision were reported. Additionally, there was no incidence of intraoperative complications reported.

#### PARTIAL REPAIR

Partial repair of massive rotator cuff tears was first described by Burkhart et al<sup>7</sup> in 1994. This technique involves repairing as much of the rotator cuff tissue as possible to partially restore shoulder functionality. Burkhart et al<sup>7</sup> described repairing the inferior portion of the rotator cuff by reattaching the infraspinatus and subscapularis tendons to their anatomic insertions while leaving the irreparable supraspinatus unrepaired. This creates force coupling of the deltoid and repaired rotator cuff tendons to allow for effective elevation of the arm. Although the supraspinatus is left unrepaired, complete coverage of the humeral head is unnecessary because the biomechanics of the shoulder are restored with repair of the infraspinatus and subscapularis. This technique was originally described as an open procedure, although several recent advancements<sup>5,9,12,16</sup> using the same principles have allowed surgeons to arthroscopically perform partial tendon repair as well as to repair different combinations of tendon tears. This procedure is typically indicated in patients with an irreparable supraspinatus and a reparable infraspinatus and subscapularis who lack glenohumeral arthritis and who continue to have pain and dysfunction after conservative management.<sup>9,12,35</sup>

Several studies have reported on the clinical outcomes after partial repair of MIRCTs (Table 1).<sup>11,12,16,22,36,47,59</sup> All studies found a statistically significant improvement in functional outcome scores compared with preoperatively. One study<sup>59</sup> evaluated patients at 1 year and again at >2 years postoperatively. The mean VAS, American Shoulder and Elbow Surgeons (ASES), and Simple Shoulder Test scores significantly improved ( $P \leq .003$ ) from preoperatively to 1-year follow-up. However, despite this initial improvement, the number of patients reporting that they were dissatisfied with the procedure increased from 1-year follow-up (6%) to >2-year follow-up (32%). Additionally, the VAS score was significantly worse at >2-year follow-up (3.16) compared with 1-year follow-up (2.13) (P = .039).

Three studies evaluated range of motion in patients undergoing partial repair of MIRCTs (Table 1).<sup>12,16,22</sup> Duralde and Bair<sup>16</sup> found an average increase in both forward flexion (114° to 154°) and external rotation (44° to 54°) over an average follow-up of 43 months. Cuff et al<sup>12</sup> found no significant changes in forward flexion  $(-14^{\circ}; P = .07)$ , external rotation  $(+1^{\circ}; P > .99)$ , or internal rotation (-4%; P > .99) from preoperatively to postoperatively. However, only patients with preoperative forward flexion  $>120^{\circ}$  were included in this study. Galasso et al<sup>22</sup> compared forward flexion, abduction, external rotation, and internal rotation of patients' affected shoulder with those of the contralateral shoulder postoperatively. No significant difference was found between the 2 shoulders with regard to forward flexion or external rotation, although patients did have an average  $10^{\circ}$  less in abduction (P <.001) and significantly less internal rotation (P < .001) in the affected shoulder compared with the contralateral shoulder at an average follow-up of 7 years.

Several studies have attempted to examine preoperative factors and their effects on clinical outcomes after partial repair of MIRCTs.<sup>16,22,59</sup> Shon et al<sup>59</sup> investigated a variety of factors such as patient demographics, tear size, and fatty infiltration of the rotator cuff tendons. The authors found that fatty infiltration of the teres minor was the only preoperative factor associated with poor outcomes. Galasso et al<sup>22</sup> found male patients to have significantly greater postoperative strength in abduction, external rotation, and internal rotation (all P < .001) compared with female patients, while younger patients displayed greater postoperative range of motion in abduction (P = .019) and external rotation (P < .03) compared with older patients. However, Duralde and Bair<sup>16</sup> also examined similar preoperative

factors and found no correlation between clinical outcomes and sex, age, or preoperative duration of symptoms. Additionally, Chen et al<sup>11</sup> found only a lower preoperative ASES score, a higher preoperative VAS score, and night pain to be associated with a greater degree of functional improvement, while age, sex, diabetes status, smoking status, acromiohumeral distance, and preoperative duration of symptoms had no effect on clinical outcomes.

Although partial repair of MIRCTs has shown promising clinical outcomes, studies have revealed a relatively high failure rate after this procedure.<sup>11,12,16,22,59</sup> Chen et al<sup>11</sup> found the rate of repair failure to be 41.6%. Failed procedures are often revised to subsequent partial repair<sup>16</sup> or rTSA.<sup>12,22</sup> The rate of complications other than failure or the need for revision is low with partial repair of MIRCTs and is reported to be 4%.<sup>16</sup>

## SUPERIOR CAPSULAR RECONSTRUCTION

MIRCTs are often associated with superior migration of the humeral head in relation to the glenoid.<sup>6,40,49</sup> SCR is a technique first described by Mihata et al<sup>40</sup> to reconstruct the superior glenohumeral joint capsule and to prohibit superior migration of the humerus. This is achieved by arthroscopically attaching a fascia lata autograft medially to the superior glenoid and laterally to the greater tuberosity while simultaneously adding sutures between the graft and residual torn rotator cuff tendons (Figure 1).<sup>39</sup> This in turn prevents superior migration of the humeral head, optimizing the force coupling necessary for arm elevation. In addition to a fascia lata autograft, recent studies have described the use of an acellular dermal allograft for this procedure.<sup>6,13,39</sup> Indications for this procedure include failed conservative management in a patient with an irreparable rotator cuff tear, lack of significant osteoarthritis, superior migration of the humeral head, and subjective shoulder dysfunction.<sup>39</sup>

Studies<sup>13,37,39,48</sup> have reported on SCR in human participants and compared clinical outcomes preoperatively and postoperatively (Table 1). Mihata et al<sup>39</sup> analyzed a group of 24 shoulders in 23 consecutive patients undergoing SCR with a fascia lata autograft. Significant improvements were demonstrated in the mean Japanese Orthopaedic Association score (48.3 to 92.6; P < .00001), ASES score (23.5 to 92.9; P < .00001), and UCLA score (9.9 to 32.4; P < .00001) at an average follow-up of 34.1 months. Additionally, average active elevation and external rotation both increased significantly by  $64^{\circ}$  (P < .001) and  $14^{\circ}$  (P < .01), respectively. The acromiohumeral distance also increased significantly from 4.6 mm to 8.7 mm (P < .001) over the study period, and no progression of osteoarthritis or rotator cuff muscle atrophy occurred in any patient.

Denard et al<sup>13</sup> evaluated a group of 59 patients over a minimum 1-year follow-up who underwent SCR with a human dermal allograft for the treatment of MIRCTs. Compared with preoperatively, the ASES score improved from 43.6 to 77.5 (P < .001), the VAS score decreased from 5.8 to 1.7 (P < .001), and the Subjective Shoulder Value (SSV) improved from 35.0 to 76.3 (P < .001). This same study also



Figure 1. Arthroscopic images of superior capsular reconstruction (SCR). (A) Massive rotator cuff tear. (B) Anchor placement. (C) Graft passage and coupling the graft to the posterior rotator cuff. (D) Coupling sutures tied. (E) Completed SCR.

analyzed range of motion and found significant improvements in forward flexion (130° to  $158^\circ$ ; P < .001), external rotation (36° to  $45^{\circ}$ ; P = .008), and internal rotation (L3 to L1; P < .001). The acromiohumeral distance also improved from a mean of 6.6 mm to 7.6 mm after just 2 weeks postoperatively. However, this outcome was not maintained at >1-vear follow-up, and the acromiohumeral distance actually decreased to a mean of 6.7 mm (P = .89). Postoperative magnetic resonance imaging results revealed only 45% of the grafts to have healed at final follow-up. Graft healing did correlate with better outcomes, as 100% of the patients with complete graft healing had successful outcomes as well as a significantly better mean ASES score (P = .027)and mean VAS score (P = .038) when compared with the nonhealed group. There was also a significantly greater prevalence of preoperative subscapularis atrophy (P =.006) in the nonhealed group.

An additional study<sup>48</sup> analyzed a group of 88 consecutive shoulders undergoing arthroscopic SCR using an acellular dermal allograft for MIRCTs with a minimum 12-month follow-up. The mean VAS and ASES scores improved from 4.0 to 1.5 (P = .005) and from 52 to 82 (P = .005), respectively. Mean range of motion values at a minimum 1-year follow-up also improved with regard to active forward flexion (120° to 160°; P = .007) and active abduction (103° to 159°; P = .02). Radiographic analysis from this study showed that the acromiohumeral interval improved from a mean of 7.1 mm preoperatively to 9.7 mm at 1-year follow-up (P = .049). Additionally, the superior capsular distance improved from a mean of 52.9 mm preoperatively to 46.2 mm at 1-year follow-up (P = .011). Lee and Min<sup>37</sup> investigated similar clinical outcomes after SCR for MIRCTs but also included in their analysis predictive factors for a retear. Although the authors found promising results in a group of 36 shoulders at a mean follow-up of 24.8 months (ASES: 50.3 to 84.0 [P < .01]; CM: 56.3 to 82.8 [P = .02]), they also found poor posterior remnant tissue and inadequate acromiohumeral interval improvement in the immediate postoperative phase to be predictive of failure and graft retears.

Failure rates after SCR for MIRCTs are moderate. Mihata et al<sup>39</sup> reported that 16.7% of patients undergoing SCR sustained a retear of either the graft or the repaired rotator cuff tendons. However, these patients either had severe fatty degeneration of the infraspinatus tendon or a history of rotator cuff surgery. Lee and Min<sup>37</sup> reported that 36.1% of patients sustained a graft retear. Denard et al<sup>13</sup> reported 25.4% of cases to be failures, defined as a final ASES score of <50, a <17-point improvement in the ASES score over the study period, or revision to repeat SCR or rTSA. The revision rate after SCR has been reported as 18.6%, with the majority of revisions being conversion to rTSA.<sup>13</sup> Intraoperative and postoperative complications other than failure or the need for revision occurred at a rate of 6.8% in 1 study and included an infection requiring debridement and placement of an antibiotic spacer and persistent biceps pain requiring biceps tenodesis.<sup>13</sup>

# **GRAFT INTERPOSITION**

The inability of the torn rotator cuff tendon to reach the anatomic footprint on the proximal humerus is a key characteristic of an MIRCT. In 1985, Post<sup>52</sup> described using a graft to bridge the gap between the torn rotator cuff tendon and the anatomic footprint of the tendon. More recently, this has been accomplished by first arthroscopically mobilizing the torn and retracted native rotator cuff tendon,

followed by a mini-open approach in which the graft is passed into the shoulder, sutured to the native rotator cuff tendon, and then anchored to the anatomic footprint on the greater tuberosity.<sup>30,45</sup> The purpose of this is to re-create the biomechanical action of the torn rotator cuff by filling the void between the retracted tendon and its insertion. Graft options for this procedure vary, and studies have reported on the use of synthetic grafts, allografts, and xenografts.<sup>30,45,52</sup> Graft interposition is indicated in patients with symptomatic massive and irreparable tears of the supraspinatus and infraspinatus who have failed conservative treatment and lack glenohumeral arthritis.<sup>45</sup> Some authors<sup>45</sup> have suggested that patients with severe atrophy and fatty infiltration of the torn tendon should not be considered for graft interposition, as these conditions are irreversible and reconstructing the tendon would likely not restore strength and function.

Neumann et al<sup>45</sup> reported on the clinical outcomes of 60 patients who underwent graft interposition with a porcine acellular dermal matrix xenograft for the treatment of MIRCTs (Table 1). The mean VAS score decreased from 4.0 preoperatively to 1.0 at a mean follow-up of 50.3 months (P < .001). The mean modified ASES score at final follow-up was 87.8, although the authors did not state whether this was a significant change from preoperatively. Mean range of motion measurements improved significantly over the study period in terms of active forward flexion  $(140.7^{\circ} \text{ to}$ 160.4°; P < .001), active external rotation at 0° of abduction  $(55.6^{\circ} \text{ to } 70.1^{\circ}; P = .001)$ , and active internal rotation at  $90^{\circ}$ of abduction (52.0° to 76.2°; P = .001). Strength was defined on a 10-point scale,<sup>45</sup> and supraspinatus strength improved from 7.7 preoperatively to 8.8 postoperatively (P < .001), while infraspinatus strength improved from 7.7 preoperatively to 9.3 postoperatively (P < .001). Postoperative ultrasonography revealed 91.8% of grafts to be intact, indicating a failure rate of 8.2%.

Two studies<sup>30,61</sup> have described the use of a human dermal allograft for graft interposition in the setting of MIRCTs (Table 1). Gupta et al<sup>30</sup> observed a group of 24 patients over a mean 36-month follow-up period and found significant improvements in both the ASES score (66.6 to 88.7; P = .0003) and 12-Item Short Form Health Survey (SF-12) score (48.8 to 56.8; P = .03). Additionally, the VAS score significantly decreased from 5.4 preoperatively to 0.9 at follow-up (P = .0002). Mean active forward flexion  $(111.7^{\circ} \text{ to } 157.3^{\circ}; P = .0002)$ , external rotation  $(46.2^{\circ} \text{ to }$ 65.1°; P = .001), and abduction (105.0° to 151.7°; P =.002) all significantly improved as well. Postoperative ultrasonography showed 76% of repairs to be intact, while all other repairs were found to be at least partially intact. Venouziou et al<sup>61</sup> demonstrated similar results in a group of 14 patients with a mean follow-up of 30.2 months. The mean VAS score improved from 7.4 preoperatively to 1.7 at 18-month follow-up (P = .001) and was maintained until final follow-up. The mean ASES score also improved significantly from 23.8 to 72.3 postoperatively (P = .001). Range of motion improved in terms of forward flexion  $(73.6^{\circ} \text{ to}$  $129.3^{\circ}$ ; P = .002), abduction (67.5° to  $117.9^{\circ}$ ; P = .002), and external rotation (7.9° to  $43.2^{\circ}$ ; P = .001). The authors found that a smaller gap size between the retracted tendon

and greater tuberosity correlated with a significantly improved postoperative VAS score, ASES score, and range of motion. However, there was no significant correlation between postoperative outcomes and age, sex, duration of symptoms, type of acromion, presence of acromioclavicular joint arthritis, muscle atrophy, or fatty infiltration.

Synthetic grafts have also been used for graft interposition in the treatment of MIRCTs.<sup>2,54</sup> Among a cohort of 41 patients, Audenart et al<sup>2</sup> found a significant improvement in the CM score from 25.7 to 72.1 (P < .001) at a mean follow-up of 43 months. Ranebo et al<sup>54</sup> conducted a longterm study on 13 consecutive patients treated with graft interposition for MIRCTs using a synthetic graft made from Dacron (Table 1). Ten patients were reached at a mean follow-up of 18 years, with a mean CM score of 46 and a mean Western Ontario Rotator Cuff Index (WORC) score of 59. Ultrasonography at follow-up revealed that 7 of the 10 (70%) patients' grafts were not intact and that 9 of 10 (90\%) patients also had full- or partial-thickness tears of the subscapularis. Therefore, the authors of this study concluded that graft interposition with a synthetic graft could not preserve rotator cuff integrity or prevent rotator cuff tear arthropathy.

# BALLOON SPACER ARTHROPLASTY

Balloon spacer arthroplasty is a procedure that was first described by Savarese and  ${\rm Romeo}^{56}$  in 2012. As mentioned previously, because of the severe disruption to the rotator cuff musculature in patients with MIRCTs, the humeral head is prone to superior migration, which can disrupt shoulder function. Balloon spacer arthroplasty was designed to prevent this superior migration by inserting a biodegradable saline-filled balloon spacer between the humerus and acromion.<sup>56</sup> This is accomplished arthroscopically using a cylindrical insertion device to introduce the spacer to the subacromial space and then inflating the spacer to its maximum volume with saline solution (Figure 2). The patient is then subjected to a period of rehabilitation, and the balloon dissolves over a period of 12 to 18 months. It is hypothesized that this restores shoulder biomechanics by permitting smooth, frictionless gliding within the joint as well as by allowing for effective action of the deltoid muscle.<sup>26,56</sup> Gervasi et al<sup>26</sup> recently described a fluoroscopy-guided technique for inserting the balloon under local anesthesia. Balloon spacer arthroplasty is contraindicated in patients with a known allergy to the device material, patients having an active or latent infectious process, or patients with signs of tissue necrosis in the subacromial space but otherwise is indicated in patients who continue to have pain or dysfunction due to MIRCTs after failure of conservative management.<sup>56</sup> However, balloon spacer arthroplasty is only available in Europe or the United States through a currently ongoing US Food and Drug Administration trial.

Senekovic et al<sup>58</sup> first reported on the clinical outcomes of arthroscopic balloon spacer arthroplasty for MIRCTs in a group of 20 consecutive patients (see Table 1). The authors found a significant increase in the CM score as early as 6



**Figure 2.** Arthroscopic images of balloon spacer arthroplasty. (A) Cylindrical insertion device entering the subacromial space. (B) Deflated spacer within the subacromial space. (C) Spacer inflating with saline solution.

weeks postoperatively (42.8) compared with preoperatively (33.4) (P = .010), with an average CM score increase of 31.5 points (P < .0001) at a final follow-up of 3 years. Senekovic et al<sup>57</sup> performed a similar study in 24 patients and found that the CM score significantly increased from 34.2 preoperatively to 67.4 at 5-year follow-up (P < .0001) (Table 1). In a group of 44 consecutive patients undergoing arthroscopic balloon spacer arthroplasty, Piekaar et al<sup>51</sup> found the mean Oxford Shoulder Score and CM score to increase from 21.8 to  $32.4 \ (P < .001)$  and from 37.1 to  $60.2 \ (P < .001)$ , respectively, at 12-month follow-up (Table 1). This same study demonstrated a significant decrease in pain on a simple 1to-10 numeric scale (6.84 to 3.27; P < .001). Gervasi et al<sup>27</sup> found significant increases in the CM and ASES scores of 37.9 points (P < .0001) and 51.5 points (P < .0001), respectively, at 1-year follow-up (Table 1).

Deranlot et al<sup>14</sup> evaluated a group of 37 consecutive patients (39 shoulders) undergoing arthroscopic balloon spacer arthroplasty for MIRCTs (Table 1). At a mean follow-up of 32.8 months, the average adjusted CM score increased significantly from 44.8 preoperatively to 76 at final follow-up (P < .001). Additionally, the mean adjusted CM score at final follow-up was significantly greater than at 1-year follow-up (P = .02). This study found significantly increased range of motion at final follow-up compared with preoperatively in terms of forward flexion (130° to 160°; P =.02), abduction (100° to 160°; P = .01), and external rotation  $(30^{\circ} \text{ to } 45^{\circ}; P = .0001)$ . Despite these promising clinical results, radiographic evidence from this study showed that the mean acromiohumeral distance actually decreased from 8.2 mm preoperatively to 6.2 mm at final follow-up (P = .002). Similarly, Prat et al<sup>53</sup> found no significant difference in the acromiohumeral distance at a mean followup of 14.4 months in a group of 22 patients treated with arthroscopic balloon spacer arthroplasty for MIRCTs. The same study found a significant increase in the mean UCLA score at follow-up (10.9 to 15.9; P = .001) but no significant differences in preoperative and postoperative range of motion values for active forward flexion (90° to 106.5°; P= .17), active external rotation (34.1° to 37.5°; P = .48), and active internal rotation (L5 to L4; P = .37). However, there was a moderate-strong correlation (r = 0.64) between preoperative range of motion and subjective general satisfaction after the procedure.

The failure rate after balloon spacer arthroplasty for MIRCTs is reported to be 3% to 8.3%.<sup>14,27,53,57,58</sup> One study<sup>14</sup> reported a single case of failure due to migration of the spacer anteriorly, which was later revised to a new spacer with satisfactory clinical outcomes. Most other cases of failure in the literature were revised to rTSA.<sup>27,57,58</sup> The complication rate is also low after balloon spacer arthroplasty but was reported to be 16.7% in 1 study.<sup>53</sup> Synovitis was reported to occur in 10% of patients in 1 study,<sup>57</sup> while another study reported the incidence of transient neural damage and both superficial and deep wound infections.<sup>53</sup>

## TRAPEZIUS TRANSFER

Tendon transfer is an available treatment option for patients presenting with MIRCTs. Lower trapezius transfer is one such procedure, in which the lower trapezius tendon is transferred to the humeral head to take the place of an irreparable posterior-superior rotator cuff tear.<sup>18,19</sup> With this type of tear, the humeral rotational position, joint reaction forces, and kinematics of the shoulder are altered compared with an intact shoulder.<sup>46</sup> Repair with lower trapezius transfer restores these biomechanical measures to the level of an intact shoulder.<sup>46</sup> Although first reported as an open procedure, Elhassan et al<sup>19</sup> described an arthroscopic approach in which the lower trapezius muscle is dissected and its tendinous insertion at the medial aspect of the scapular spine is detached. This is then augmented with an Achilles tendon allograft, which is attached to the supraspinatus footprint on the greater tuberosity. Lower trapezius transfer is typically indicated in younger and active patients with a posterior-superior irreparable rotator cuff tear who have minimal to mild glenohumeral arthritis.<sup>18</sup> Another study<sup>28</sup> described and attempted to determine the efficacy of superior trapezius transfer for the treatment of an irreparable subscapularis tear. However, this study was met with poor clinical outcomes, and the authors do not recommend it as a viable treatment option for MIRCTs.

Only 1 study has reported on clinical outcomes after lower trapezius transfer for the treatment of MIRCTs (see Table 1).<sup>19</sup> At a mean follow-up of 47 months, the mean SSV significantly increased from 54% preoperatively to 78% postoperatively (P < .01). The mean DASH score

significantly improved from 52 to 18 (P < .01). Preoperative range of motion included  $70^{\circ}$  of forward flexion,  $40^{\circ}$  of abduction, and 20° of external rotation, which all significantly increased (all P < .01) to mean values of  $120^{\circ}$ ,  $90^{\circ}$ , and 50°, respectively, while mean internal rotation was maintained at the L3 spinous process level. Interestingly, patients with  $>60^{\circ}$  of forward flexion and abduction preoperatively demonstrated a significantly greater improvement in range of motion compared with patients with less preoperative range of motion. The radiographic evaluation from this study demonstrated that 26 of 33 (79%) patients had proximal migration of the humeral head preoperatively, with a mean acromiohumeral distance of 2.3 mm, which increased to 8 mm at final follow-up. Failure occurred in 1 of 33 patients (3%) because of a postoperative infection that required debridement and later revision to shoulder fusion. Seroma formation was reported in 4 of 33 patients (12%), but no other complications were reported in this study.

#### LATISSIMUS DORSI TRANSFER

Latissimus dorsi transfer was first described by Gerber et al<sup>25</sup> in 1988 as a surgical technique used for patients with MIRCTs, specifically of the supraspinatus and infraspinatus tendons. The original technique involves harvesting the latissimus dorsi tendon from its insertion on the floor of the intertubercular groove of the humerus and fixing it to the superolateral humeral head. This allows the latissimus dorsi to function as an external rotator as well as a direct countering force to superior migration of the humeral head on attempted flexion and abduction of the shoulder. The original technique was performed as an open procedure, although an arthroscopically assisted technique has been described in multiple studies.  $^{8,10,29,33-35,42,50}$  Indications for this procedure include younger patients who are suffering from severe functional disabilities caused by an irreparable posterior-superior rotator cuff tear and patients with minimal to no glenohumeral arthritis.<sup>1,34,43</sup>

Numerous studies have discussed functional and clinical outcomes after latissimus dorsi transfer for MIRCTs (see Table 1).<sup>§</sup> One study<sup>24</sup> evaluated the functional outcome scores of 46 cases in which open transfer was performed. At a minimum 10-year follow-up, the mean SSV increased from 29.0% preoperatively to 70.1%, and the mean CM score improved from 47.3 to 63.5 (both P < .0001). Additionally, mean forward flexion increased from 118.0° to 132.4° (P = .029), mean abduction increased from 112.1° to 122.6° (P = .089), and mean external rotation increased from 17.9° to 32.5° (P = .0001).

Another study<sup>17</sup> evaluated 108 patients undergoing open latissimus dorsi transfer for MIRCTs. At a mean follow-up of 9.3 years, the CM score improved to 62.0 from 36.1 preoperatively (P < .0001), excluding the 10% of patients in whom the procedure failed. The mean ASES score improved from 30.1 to 70.2 (P < .0001), and the mean VAS score decreased from 7.8 to 2.4 (P < .0001). However, an increase in rotator cuff arthropathy and a decrease in the acromiohumeral distance (5.9 mm to 4.9 mm; P < .0001) were also noted.

Eight studies have examined clinical outcomes after arthroscopically assisted latissimus dorsi transfer.  $^{8,10,17,24,29,34,42,50}$  Grimberg et al  $^{29}$  evaluated 55 cases clinically and radiographically at a mean follow-up of 29 months. The authors noted that the CM score improved significantly from 37.0 preoperatively to 65.4 postoperatively (P < .001). Additionally, the SSV improved from a mean of 26% preoperatively to 71.1% postoperatively (P < .001). Active forward flexion increased from  $134^{\circ}$  to 157° (P < .001), mean active abduction increased from 67° to  $92.5^{\circ}$  (P < .001), and mean active external rotation increased from  $29^{\circ}$  to  $41.5^{\circ}$  (P < .001). Mean abduction strength increased from 1.4 kg to 4.8 kg, and the mean CM pain subscore improved from 1.7 preoperatively to 12.6 postoperatively. The authors noted that there was no statistical difference between preoperative and follow-up acromiohumeral distance and that there was also no increase in osteoarthritic stage with this procedure in contrast to the open technique.<sup>17,24</sup> Kanatli et al<sup>34</sup> evaluated patients undergoing the same technique after a mean of 26.4 months and noted similar improvements in functional outcome scores. They also found that active forward flexion improved from 58° to 130° (P < .001), active abduction increased from 51° to 129.7° (P < .001), and active external rotation increased from  $13.3^{\circ}$  to  $32^{\circ}$  (P < .001). The mean acromiohumeral distance significantly improved from 3.13 mm preoperatively to 5.67 mm postoperatively (P < .001) in this study.

In another study,<sup>33</sup> 9 patients who underwent arthroscopic-assisted latissimus dorsi transfer were analyzed to determine changes in maximum shoulder flexion/ extension, abduction/adduction, and internal/external rotation. The authors found a significant increase in shoulder range of motion in all movements after 6 months compared with preoperatively (P < .001). Similarly, Castricini et al<sup>10</sup> noted a significant improvement in external rotation at a mean follow-up of 27 months (P < .05). The authors also noted a significant improvement in the mean CM score and pain score (P < .05) and no significant osteoarthritis progression or proximal migration of the humeral head after surgery. Petricciolo et al<sup>50</sup> also noted significant improvements in shoulder range of motion after latissimus dorsi transfer. Of the 33 patients included in their retrospective study, it was noted that forward flexion improved from an average of 138° preoperatively to 168° (P < .05) and that active external rotation increased from an average of  $7^{\circ}$  to  $34^{\circ}$  (P < .05) at an average follow-up of 35.7 months.

One study<sup>42</sup> evaluated a group of 24 patients undergoing arthroscopic latissimus dorsi transfer for irreparable subscapularis tendon tears. At a follow-up of >1 year, the mean CM score improved from 46 to 69 (P < .001), the ASES score improved from 40 to 70 (P < .001), and the VAS score improved from 6 to 2 (P = .006). This study also found significant improvements in active forward flexion (135° to 166°; P = .016) and internal rotation (L5 to L1; P =.010), while active external rotation trended toward a

<sup>§</sup>References 8, 10, 17, 24, 29, 33–35, 42, 50.

significant improvement (51° to 68°; P = .062). Postoperative magnetic resonance imaging studies of these patients showed adequate healing of the transferred latissimus dorsi tendon to the humeral head in all patients.

Several studies have investigated preoperative factors that affect the clinical outcomes of latissimus dorsi transfer for MIRCTs.<sup>1,8,17,24,29,50</sup> Anastasopoulos et al<sup>1</sup> suggested that patients with irreparable posterior-superior rotator cuff tears with associated atrophy or fatty degeneration of the subscapularis and deltoid, as well as a large critical shoulder angle, are more likely to have poor clinical outcomes. Successful outcomes were associated with a preoperative critical shoulder angle of <36°. El-Azab et al<sup>17</sup> noted that damage to the deltoid had an adverse effect on revision latissimus dorsi transfer. Gerber et al<sup>24</sup> noted that patients had a higher risk of being dissatisfied if they had teres minor atrophy, preoperative forward flexion <90°, superior fixation of latissimus dorsi transfer, and workers' compensation status. Grimberg et al<sup>29</sup> suggested that patients had a higher chance of being satisfied if they were men, were younger than 65 years, had no history of shoulder surgery, and underwent antegrade-type fixation consisting of a round button applied on the anterior humeral cortex at the distal end of the humeral tunnel using 2 knotted sutures for fixation on the button.

Latissimus dorsi transfer has shown promising clinical results in select patients, with a moderate rate of failure.<sup>17,24,29,34,35,43</sup> The highest reported failure rate for this procedure is  $38\%^{35}$  within the first 2 years postoperatively. However, other studies<sup>17,24,34,43</sup> have reported much lower rates of failure, with 1 study reporting an 86% satisfaction rate at 10-year follow-up.<sup>17</sup> Kany et al<sup>35</sup> found that patients with failure of the repair had significantly worse clinical outcomes than patients with adequate tendon healing. Complications include stiffness, traumatic failure of the transfer, hematoma, subscapularis retears, resolving nerve dysesthesia, and deltoid reattachment failure, although appropriate patient selection is of paramount importance, and the procedure requires a high level of surgical skill and experience.<sup>24,35</sup>

## LIMITATIONS

The limitations of this review should be noted. There is a lack of randomized controlled trials comparing these procedures with each other or to rTSA. Many of the studies included in this review present a short-term follow-up. In contrast, Gerber et al<sup>23</sup> have reported on the 15-year followup of rTSA with a relatively high failure rate, which may be related to older prosthetic designs and a long duration of follow-up, making these techniques difficult to compare. Several of the studies included in this review lack postoperative imaging results to confirm the integrity of repairs or grafts. Many of these studies also have an inconsistent definition of a successful outcome. For example, 1 study<sup>34</sup> reporting on the outcomes of arthroscopic latissimus dorsi transfer considered an average postoperative CM score of 59.73 to be a successful result.

#### CONCLUSION

Reverse total shoulder arthroplasty has gained favor as a surgical treatment option for patients presenting with MIRCTs. However, complications after rTSA occur at a rate of 20% to 50%, and there are concerns regarding the longevity of these implants, as revision is necessary in 10% to 33% of cases. Arthroscopic debridement provides pain relief and improved clinical outcomes but does not provide functional benefits for patients with MIRCTs. Partial repair is an alternative treatment option available for patients with an irreparable supraspinatus tendon and reparable infraspinatus and subscapularis tendons; this technique partially restores shoulder biomechanics and has shown promising clinical outcomes but high failure rates as well. SCR reduces superior translation of the glenohumeral joint for patients with MIRCTs, and while preliminary outcomes are promising, additional investigation is necessary to confirm these results. Graft interposition has shown success with dermal allografts and xenografts in short-term followup studies. However, synthetic grafts, especially at longterm follow-up, have demonstrated a high risk of retears. Balloon spacer arthroplasty is a simple and minimally invasive technique that improves clinical outcomes but does not appear to have an effect on the acromiohumeral distance. Both latissimus dorsi and trapezius transfer provide significant functional improvement and are typically indicated in younger, active patients who present with MIRCTs. Future studies involving randomized controlled trials are necessary for confirming the efficacy of these procedures as well as to determine when each is indicated based on specific patient and anatomic factors.

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