

1 **Clinical outcome of latissimus dorsi tendon transfer and partial cuff repair**
2 **in irreparable postero-superior rotator cuff tear**

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6 **Abstract**

7 **Background and Purpose:** Irreparable Rotator Cuff Tears (RCTs) is a common cause of pain in adult
8 population, requiring in many cases a surgical treatment. Possible alternatives are debridement, partial repair,
9 muscle transfers and joint replacement. We evaluated two groups of patients with irreparable rotator cuff tear
10 treated surgically: one group received an arthroscopic assisted Latissimus Dorsi tendon transfer (LDTT), and
11 the other an arthroscopic rotator cuff partial repair. Aim of our study is to compare clinical results and
12 quality of life in two groups of patients with massive irreparable rotator cuff tear: one receiving an
13 arthroscopic Latissimus Dorsi tendon transfer and the other receiving an arthroscopic rotator cuff partial
14 repair.

15 **Methods:** 40 patients were assigned to two groups: 20 patients to group TT treated with latissimus dorsi
16 tendon transfer and 20 patients to group PR treated with a partial repair. The average follow-up duration was
17 2.8 years (1-5; SD: 3). Pre- and post-operative modified-UCLA shoulder score, ROM, measurement of the
18 strength and the RC-QOL were used to asses the outcome.

19 **Results:** Latissimus Dorsi Tendon Transfer showed significant improvements when compared to partial
20 repair in UCLA score results, strength and rc-qol (rotator cuff quality of life) questionnaire. No differences
21 were found between the groups in pain relief.

22 **Conclusion:** Both techniques are effective in reducing patients' symptoms. We believe that in younger,
23 high-demanding patients with no or mild osteoarthritis, the LDTT represents a valid treatment option with
24 better modified UCLA score improvement and strength at our follow-up.

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26 Conflict of interest: NO

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30 **Keywords:** rotator cuff tears; irreparable rotator cuff tears; partial repair; tendon transfer; latissimus dorsi
31 transfer;

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37 **Introduction**

38 Rotator cuff tears is a common cause of pain in adult population and often produce lasting symptoms as pain
39 and limitation of normal activities. Reduced acromion-humeral distance (<5 mm), fatty degeneration of the
40 muscle and huge tendinous tissue deficit are factors that suggest not to repair the lesion¹. Possible treatments
41 for irreparable rotator cuff tears are debridement associated to subacromial bursectomy and long head of the
42 biceps tenotomy, partial cuff repair, tendon transfers (latissimus dorsi, pectoralis major) and joint
43 replacement². Reverse total shoulder arthroplasty is often used in elderly patients where the rotator cuff
44 lesion coexists with degenerative gleno-humeral arthropathy. Latissimus Dorsi tendon transfer is advocated
45 in younger patients without gleno-humeral arthropathy, in which a postero-superior irreparable rotator cuff
46 tear causes pain and loss of function. Gervasi et al. proposed an arthroscopic LD transfer avoiding deltoid
47 sacrifice³. We didn't found in literature studies comparing the Latissimus Dorsi tendon transfer to other
48 techniques for the treatment of irreparable postero-superior rotator cuff tear. Aim of our study is to compare
49 clinical results and quality of life in two groups of patients with massive irreparable rotator cuff tear: one
50 receiving an arthroscopic Latissimus Dorsi tendon transfer and the other receiving an arthroscopic rotator
51 cuff partial repair.

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53 **Methods**

54 *Patient population*

55 Inclusion criteria were daily and nocturnal pain, previous conservative treatment (NSAIDs, intrarticular
56 injection of corticosteroids and physiotherapy) without results, strength loss and an intact or reparable
57 subscapularis tendon. Exclusion criteria were: shoulder instability, previous rotator cuff surgery, fracture of
58 the glenoid or smaller tuberosity, gleno-humeral osteoarthritis, prior surgery of the shoulder, cervical
59 radiculopathy, capsule-ligamentous lesions, inflammatory disease of the connective tissue; (6) other general
60 comorbidities (cardiovascular and cerebrovascular diseases, lower extremity ischaemia, neurological
61 diseases, and uncontrolled diabetes), or psychiatric illness. In the period between January 2007 and January
62 2011 we included in our study 40 patients respecting inclusion and exclusion criteria. These patients were
63 assigned to two groups: 20 patients to group TT (13 men and 7 women) treated with arthroscopic assisted
64 latissimus dorsi tendon transfer and 20 patients to group PR (11 men and 9 women) treated with a rotator
65 cuff partial repair. Patients were intraoperatively allocated to the two groups, according to the possibility to
66 first attempt a partial repair of the cuff. When the tissue's features allowed for partial repair, it was

67 performed, when they didn't allow, the tendon transfer was performed. The mean follow-up was 2.8 years
68 (1-5; SD: ± 3), demographic features are reported in Table 1. Patients evaluation was performed immediately
69 before the index operation, and postoperatively at a minimum 2-year follow-up.

70 *Surgical technique*

71 All surgical procedures were performed by the senior orthopaedic surgeon. All the patients received a pre-
72 operative interscalene block plus a general anesthesia. Patient were positioned in a lateral decubitus with the
73 shoulder and elbow flexed at 90° to allow both latissimus dorsi exposure and later arthroscopic transfer. The
74 same set-up was managed for group PR. Gravity joint irrigation was provided using 4 L saline bags hung at a
75 height of 8 feet. The extent of tear and the tendon retraction were measured intra-operatively in both the
76 coronal and sagittal planes according to the classification system described by Boileau et al⁴

77 Latissimus dorsi tendon transfer

78 The procedure was performed according to the technique described by Gervasi in 2007(Gervasi, Causero et
79 al. 2007)

80 Phase 1: diagnostic arthroscopy

81 Standard portals, including a posterior portal (P), an anterior inferior portal (A), a posterolateral portal (PL),
82 and an anterolateral portal (AL) are performed.

83 Phase 2: harvesting the tendon

84 The arm should be released from the traction and abducted and internally rotated. After probing the
85 latissimus dorsi tendon with a finger, a 6–7 cm long curved incision line is firstly marked and then made
86 along the muscle's profile at the axillary level. Using blunt dissection, the tendon can be isolated and
87 detached from its humeral insertion.

88 Phase 3: prepare the tendon

89 The two sides of the tendon are reinforced with suture stitches by differently colored high strength sutures.
90 Then the same is done for the end of the tendon, bridging the lateral- and medial-side stitches to strengthen
91 the tendon during its transfer through the subacromial space, preventing the tendon from splitting

92 Phase 4: tendon transfer

93 Once the limb traction is restored, we use one finger to isolate the fibers of the brachial triceps. At this point
94 it is identified through a PL portal vision the best way to pass a 30° curved grasper through the AL portal to
95 the armpit, between the teres minor and the posterior deltoid. Once the curved grasper has exited the axillary
96 incision, we use to pass two transparent suction tubes through the pathway to reduces the risk of rotating the

97 graft while shuttling it to the subacromial space. Finally a suture retriever is used to shuttle out of the AL
98 portal the lateral and the medial side tendon sutures through the lateral and the medial tube respectively.

99 Phase 5: tendon fixation (Fig.1)

100 To fix the tendon on the prepared site on the greater tuberosity, the medial and the lateral stitches are loaded
101 on 5.5 mm knotless anchors. The lateral one is placed as anteriorly as possible, aiming to the bicipital
102 groove.

103 Partial repair

104 After the footprint was identified at the greater tuberosity, through a shaver (Arthex, Naples, FL, USA) it
105 was prepared until a bleeding surface was achieved. We performed a partial repair of the irreparable lesion
106 according to the technique previously described by Burkhart et al⁵.

107 *Post-operative Management*

108 After the operation, the joints in the TT group were immobilised in a 45° abduction sling for 6 weeks. The
109 sling was then removed and patients were allowed for assisted passive mobilization on all planes and soft
110 active mobilization until the thirth postoperative month. The main target during this period was to achieve a
111 good neuromuscular control of the transfered Latissimus Dorsii tendon in its new role as a humeral head
112 stabilizer and external rotator. After 3 months trengthening excercises for the deltoid and the scapular
113 stabilizers were started.

114 In patient included in the partial repair group a sling was used for the first 4 postoperative weeks and were
115 allowed free flexion and internal rotation from the first postoperative day. At 4 weeks a progressive free
116 ROM in all directions was allowed. On the first day after surgery passive external rotation was started while
117 overhead stretching was allowed 4 weeks postoperatively to avoid damaging the repair. At 4 weeks, the sling
118 was removed, and overhead stretching with a rope and pulley was started. Strengthening of the deltoid and of
119 the scapular stabilizers were initiated at 8 weeks after the surgery.

120 Evaluation

121 *Imaging*

122 All patients received a standard pre-operative assessment using standard radiographs and MRI scans.
123 According to the classification of Hamada et al., the Acromio-Humeral Index (AHI) was preoperatively
124 assessed for each patient (Table 2). Fatty infiltration was evaluated using MRI scans and classified according
125 to Goutallier et al⁶

126 *Functional assessment*

127 A modified UCLA rating scale for pain, function, ROM, and patient satisfaction was used to evaluate each
128 patient pre-operatively and at follow-up. According to Ellman, an excellent UCLA score is 34 to 35 points; a
129 good score is 28 to 33 points; a fair score is 21 to 27 points and a poor score is 0 to 20 points⁷. Pre- and post-
130 operative measurement of the strength were performed through a handheld dynamometer (PowerTrack
131 MMT; JTech Medical Industries, Alpine, Utah, and Muscletester; Hoggan Health Industries, South Draper,
132 Utah).⁸ The ranges of motion in elevation, external rotation, internal rotation, and hand behind back lift-off
133 were assessed.

134 *Quality of life (RC-QOL)*

135 All patients completed a self-administered RC-QOL(rotator cuff quality of life) questionnaire. The RC-QOL
136 questionnaire is a simple disease-specific outcome measure that evaluate the impact of rotator cuff disease on
137 the general quality of life. The total score ranges from 0 (worst score) to 3400 (best score), results are given
138 as percentage (0-100%). This questionnaire has been translated and validated for the Italian language⁹.

139 *Statistical analysis*

140 We designed the investigation as a prospective case-control study; two independent populations (patients
141 undergoing arthroscopic latissimus dorsi tendon transfer, and patients receiving an arthroscopic rotator cuff
142 partial repair) were considered. The data used to design the study were the following: Alpha-value: 0.05,
143 Power: 0.8, Ratio between cases and control: 1, Probability of the event in cases: 0.3, Probability of the event
144 in controls: 0.3. According to the power analysis calculation, we needed a total of 18 patients in each group
145 to satisfy the above premises. We recruited 20 patients per group. The differences between preoperative and
146 postoperative active forward flexion, external rotation, internal rotation and UCLA shoulder score for both
147 the groups were assessed by an unpaired Student t test. The effects of tear size, tendon retraction, fatty
148 degeneration and AHI grade on outcome were also assessed by 1-way ANOVA. Statistical significance was
149 set at $P < 0.05$. Data are presented using mean, median or standard deviation, and range and data ranges as
150 appropriate. Statistical analysis was performed with the SPSS software package, version 11.0 (SPSS,
151 Chicago, IL).

152 *Source of Funding*

153 There was no external funding source for this study.

154 **Results**

155 *Associated procedures*

156 The associated procedures have been performed, are reported in Table 2.

157 *Range of motion*

158 ROM measures of both groups at the latest follow-up (post-operative forward elevation, internal rotation,
159 and external rotation) were significantly improved ($P<0.05$) compared to pre-operative values, with
160 significant intergroup differences (Table 3).

161 *Functional assessment* (Table 4)

162 Pain measures (Visual Analogue Scale, VAS) improved significantly from pre- to post-operative time for
163 both groups [Group TT: from a mean pre-operative value of 6.9 ± 1.7 to the final post-operative value of
164 1.3 ± 0.7 ($P<0.05$)]; [Group PR: mean preoperative value: 6.6 ± 1.8 ; final postoperative value:
165 1.5 ± 0.8 ($P<0.05$)]. Results from UCLA shoulder score showed a mean pre-operative value of 7.3 ± 2.5 for
166 group TT and 7.6 ± 3.9 ($P=n.s.$) for group B while the post-operative values at the latest follow-up showed a
167 statistically significant improvement in both groups [30.3 ± 4.2 for group TT and 20.1 ± 3.4 for group PR]
168 ($P<0.05$). Intergroup differences in functional and strength domains were statistically significant ($P<0.05$)
169 starting from the first post-operative month to the whole duration of the study. With regard to the strength,
170 there was a statistically significant improvement between pre-operative evaluation and the last follow-up for
171 both groups, but with significant intergroup differences ($P<0.05$). According to the UCLA rating system, in
172 group TT 12 patients (63%) had an excellent result (34–35 points), 5 (26%) a good result (28–33 points), and
173 2 (11%) a fair result (21–27 points), whereas in group PR 11 patients (55%) had an excellent result (34–35
174 points), 5 patients (25%) a good result (28–33 points), and 4 (20%) a fair result (21–27 points). There were
175 no poor results (0–20 points).

176 *Tendons' features*

177 The UCLA shoulder score demonstrated a statistically significant difference between patients of both groups
178 (TT and PR) with stage 2 degeneration and those with stage 3 or stage 4 fatty degeneration ($P < 0.0001$),
179 while no difference in outcome between those with stage 3 and those with stage 4 degeneration. The same
180 was noticed concerning the AHI. Patients with an AHI grade 1 achieved significant better UCLA outcomes
181 than those with an AHI grade 2 ($P < 0.0001$).

182 *Quality of Life (RC-QOL)*

183 The RC-QOL demonstrated a statistically significant difference between the groups [group TT: 81.8 ± 9.3 ;
184 group PR: 69.3 ± 8.7] ($P<0.05$) (Table 4).

185 *Ruptures*

186 Based on a clinical diagnosis, given a sudden loss of function, a case of LDT rupture was recorded, after 13
187 months from surgery. In this patient a reverse total shoulder arthroplasty was performed.

188 **Discussion**

189 Changing the insertion site, from the anatomical one to the great tuberosity, the Latissimus Dorsi muscle
190 become an external rotator^(10,11,12) and this is the biomechanical feature on which the LDT transfer lies. In
191 2007 Gervasi and colleagues(Gervasi, Causero et al. 2007) proposed an arthroscopically assisted LDT
192 transfer. Our technique proposes few changes compared to the Gervasi's one. We developed some tricks to
193 obtain the widest footprint coverage and to avoid the graft rolling and rotation while shuttling it to the
194 subacromial space: the particular pattern of tendon edge's stitches, the use of two separate sutures and of two
195 suction tubes and the use of an in-out positioned grasper to shuttle the tendon through the subacromial space.
196 A biomechanical study of Oh and colleagues¹³ demonstrated as the abnormally increased maximum internal
197 rotation occurring in massive rotator cuff tears was reversed after LDT transfer. But the authors also outlined
198 as an excessive muscle tension (as in the case of a LDT transfer with limited excursion) could cause,
199 paradoxically, lost of internal rotation. To avoid this troublesome scenario, we recommend an accurate
200 release of the muscle, allowing the tendon to reach the posterior rim of the acromion, thus ensuring sufficient
201 length once it is passed into the subacromial space. Our aim was to cover as more as possible the humeral
202 head. The wider is the coverage, the better will be the healing potential of the tendon and the higher will be
203 the depressive action on the humeral head. Moreover, we try to fix the tendon edge as anteriorly as possible
204 to the bicipital groove, to obtain the maximal tenodesis effect and the best balance between the subscapular
205 and the latissimus dorsi muscles. As suggested by Gervasi (Gervasi, Causero et al. 2007), when fixing the
206 tendon close to the articular cartilage and the long head of biceps groove, the fiber's distension generates an
207 elastic force pulling back distally and posteriorly along the LD bill axis, thus this force contributes maintain
208 the humeral head located at the rotation centre of the glenoid track. The main positive effects on external
209 rotation is achieved just by changing the biomechanical features of the LDT. At its natural insertion it acts as
210 an important restraint to external rotation, while the maximum moment-generating capacity is restored
211 significantly through each new insertion site¹⁴; for this reason we believe that it is more important to cover as
212 much surface as possible of the rotator cuff footprint. Results of our study show how partial repair leads to
213 pain relief and slightly improvement of shoulder function, conversely the tendon transfer allows for a greater
214 recovery of the shoulder active movements. In the post-operative period, patients' quality of life improved in
215 the overall cohort but better results were found in the tendon transfer group considering shoulder function
216 and strength. Our results, in agreement with the literature, demonstrated good to excellent recovery of
217 shoulder function.^(15,16,17,18) We attributed this successful outcomes to the careful postoperative rehabilitation
218 targeted to an extensive work to achieve the best neuromuscular control. Strengths of the study are a single
219 surgeon performing all the operations and the strictly inclusion and exclusion criteria. Limitations are given
220 by the short follow-up, the small study population and the lack for postoperative radiological controls. We
221 recorded 1 case of tendon rupture after 13 months from surgery and no clinical detectable failure of any
222 partial repair. The lack for radiological controls didn't allow us to record partial repair failures during the
223 follow-up period neither the possible progression of the AHI. Although it has been shown as latissimus dorsi
224 transfer is not able to avoid the risk for glenohumeral joint arthropathy¹⁹, our study shows its effectiveness in

225 younger patients leading to significative ROM, strength and pain reduction. An ideal candidate for a tendon
226 transfer has mild to moderate shoulder weakness associated with an irreparable posterior-superior rotator
227 cuff tear²⁰ .Also if both techniques are effective in reducing patients' symptoms we think that for high
228 demanding, younger patients, latissimus dorsi tendon transfer should be considered, since it restores an
229 higher shoulder strength compared to partial repair. We didn't found in literature other studies comparing the
230 LDTT to other techniques. It's not possible to demonstrate the superiority of one technique on the others but,
231 according to results of our study, we believe that in younger, high-demanding patients with no or mild
232 osteoarthritis, the LDTT represents a valid treatment option.

233 **Figures and tables**

234 **Figure 1:** picture showing the LDT fixed. Intrarticular arthroscopic view.

235 **Table 1:** patients's features and associated surgical treatments. Values are given as average with range in
236 brackets

237 **Table 2:** preoperative tendon features

238 **Table 3:** Range of movement. Values are given as average \pm standard deviation with range in brackets

239 **Table 4:** Pre-and post-operative values of UCLA, VAS and RC-QOL. Values are given as average \pm
240 standard deviation with range in brackets

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Figure1

Table1

	Group TT	Group PR
Sex	13 M; 7 W	11 M; 9 W
Age	62.5 years (range 45–77)	64.9 years (range 47-78)
Dominant arm involved	18	17
time from symptoms to surgery	7 months (range 7-23)	8 months (5–13)
Associated treatment		
Acromioplasty	4	6
LHB tenotomy	14	13
Subscapular repair	2	3

Table 1: patients's features and associated surgical treatments. Values are given as average with range in brackets

Table2

	Group TT	Group PR
Size		
Large (3-5 cm)	4	7
Massive (> 5 cm))	16	13
Tendon retraction		
Stage III	6	4
Stage IV	14	16
Location		
SSP	4	8
SSP+ISP	16	12
Stage of Fatty infiltration		
0	0	0
1	0	0
2	8	10
3	9	9
4	3	1
AHI grade		
1	3	4
2	17	16
3	0	0
4	0	0
5	0	0

Table 2: preoperative tendon features

Table3

ROM	Group TT		p Value	Group PR	
	Baseline	Last follow-up		Baseline	Last follow-up
Passive External rotation, degree	22.6±13.5 (15-55)	59.1±10.2 (53-74)	<0.05	18.3±11.7 (17-33)	57.4±9.1 (35-62)
Active, External rotation, degree,	14.5±11.3 (9-26)	41.2±8.7 (31-52)	<0.05	15.8±9.2 (11-31)	38.4±12.0 (33-54)
Internal rotation, degree	a level between L3-S1	11 pts T8; 5 pts T9; 4 pts T10		a level between L3-S1	14 pts T8; 4 pts T9; 2 pts T10
Passive Forward flexion, degree	119.8±13.0 (105-130)	171.2±9.7 (148-178)	<0.05	129.2±18.2 (90-160)	169±10.9 (145-180)
Active Forward flexion, degree	83.5±11.0 (72-98)	131±9.0 (117-145)	<0.05	86.3±8.2 (68-89)	110±12.7 (98-132)

Table 3: Range of movement. Values are given as average ± standard deviation with range in brackets

Table4

	Group TT		p Value	Group PR	
	Baseline	Last F-U		Baseline	Last F-U
UCLA	7.3±2.5 (4-9)	30.3±4.2 (29-34)	<0.05	7.6±3.9 (4-10)	20.1±3.4 (18-25)
VAS	6.9±1.7 (6-9)	1.3±0.7 (1-3)	<0.05	6.6±1.8 (6-9)	1.5±0.8 (1-3)
RC-QOL	n.a.	81.8±9.3 (78-92)		n.a.	69.3±8.7 (63-77)

Table 4: Pre-and post-operative values of UCLA, VAS and RC-QOL. Values are given as average ± standard deviation with range in brackets.