



Letter to the Editor regarding Young et al: “Reverse shoulder arthroplasty with and without latissimus and teres major transfer for patients with combined loss of elevation and external rotation: a prospective, randomized investigation”



To the Editor:

We read with great interest the article entitled “Reverse shoulder arthroplasty with and without latissimus and teres major transfer for patients with combined loss of elevation and external rotation: a prospective, randomized investigation” by Young et al.²² We would like to commend the authors for their investigation of an important research question: Do “CLEER” patients (combined loss of elevation and external rotation) who undergo reverse shoulder arthroplasty (RSA) with latissimus dorsi and teres major transfer (+LD/TM) have better functional results than those treated with RSA alone? They performed a randomized controlled trial (RCT) and found no significant differences at 2 to 3.5 years of follow-up. At least 3 important methodological weaknesses limit the strength of this RCT and deserve discussion: patient selection, statistics, and interpretation of results.

Patient selection

CLEER is a well-defined clinical and anatomic entity.^{3,4,6} Patients present with shoulder pseudoparalysis and complete loss of active external rotation. Imaging studies show massive irreparable posterosuperior cuff tears with severe fatty infiltration (Goutallier grades 3 and 4) of infraspinatus and atrophy/absent teres minor (Tm). Unfortunately, some of the recruited patients in the study of Young et al did not meet CLEER criteria. As shown in Table II, some patients had *no pseudoparalysis* (active forward elevation $\geq 100^\circ$) and the authors included 5 patients who had an *intact Tm* (Table I) or grade 2 fatty infiltration of Tm. Furthermore, as shown in Figure 3, some patients had no active external rotation deficit with a *preoperative* Activities of Daily Living requiring active External Rotation (ADLER) score near 30 points (maximum) and thus should not have been

labeled as CLEER patients nor included in the study. As acknowledged by the authors themselves and shown in Table III, the median *preoperative* ADLER score of their cohort is far higher (17/30 points in the RSA group and 16/30 points in the RSA+LD/TM group) than in the usual population affected by CLEER ($4 \pm 3/30$).^{3,4,6,7,9,16,17,20}

Statistics

The authors calculated a minimum sample size of 20 total patients. The authors quote 2 publications^{3,6} for their power calculation, but chose the least restrictive ($26^\circ \pm 4^\circ$)³ and used a unilateral statistical test. The choice of the other referenced results ($25^\circ \pm 5^\circ$),⁶ a bilateral test (as it is customary), and a classical hypothesis of 20% lost to follow-up would have led to a minimum of 40 patients required to reach statistical power (20 patients *per* group). Furthermore, the number of randomized patients ($n = 31$) dropped to only 22 reviewed at a minimum of 2 years for patient-reported outcomes, and only 12 assessed for active ROM. With the methodology employed, the final 2-year follow-up was underpowered to detect differences between groups, as acknowledged by the authors themselves: “The relatively small sample size of this study certainly leads to potential study frailty.”²²

Interpretation of the results

First, despite the fact that the purpose of this study was to analyze restoration of active ER after RSA \pm LD/TM, the authors could not compare pre- and postoperative range of motion as they failed to report *preoperative* values of active ER. Secondly, the authors suggest that the spontaneous resolution of the preoperative hornblower sign, observed in 58.3% of patients in the control group (RSA alone), could be attributed to increased posterior deltoid recruitment coupled with the use of a constrained prosthesis and a slightly lateralized center of rotation (2–3 mm of

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lateralization).¹ To prove this interpretation, the authors should have stratified results according to the *presence or absence of Tm on preoperative imaging studies*. Unfortunately, they fail to do it. Greiner et al¹¹ performed this type of stratification in their RCT study comparing the rotational outcomes of lateralized RSA vs. medialized (Grammont type) RSA. They observed that lateralized RSA (10 mm BIO-RSA) significantly improved active ER in patients with an intact Tm muscle, but not when this muscle was absent/atrophied. In other words, regardless of the type of reverse prosthesis used (lateralized or medialized), a functional Tm muscle is needed to improve postoperative *active ER*. As already shown in many studies,^{3,4,9,17,21} a lateralized RSA is able to improve *passive ER* (by delaying posterior impingement), but a viable Tm muscle is needed to improve *active ER*.

The primary purpose of our letter is to re-emphasize the indications for RSA+LD/TM. This combined procedure is indicated *only* in CLEER patients with a positive hornblower sign and atrophy/absence of the Tm muscle on imaging.^{3,4,6} The absence/atrophy of Tm is a risk factor for poorer functional results, *whatever* the surgical procedure: biceps tenotomy/tenodesis,^{2,19} cuff repair,^{12,14} LD tendon transfer,^{8,10,15} and of course, RSA.^{4,7,9,17} The Tm contributes up to 45% of ER power, mainly in abduction, which is important to control hand positioning in space.^{2,5,13,14,18,21} Even when not active, the tendon transfer provides a tenodesis effect and allows patients to control their hand position in space and perform ADLs. In patients with persistently deficient active ER and handicap for ADLs after isolated RSA, Puskas et al¹⁶ have even shown that secondary LD transfer significantly improved active mobility and subjective results.

In summary, in light of the methodological weaknesses noted, we find it unsurprising that this study reported equivalent functional results in patients treated with RSA alone and with RSA+LD/TM. We encourage the authors to perform further studies, better controlling for these biases, and hopefully providing a “CLEER answer” to this difficult clinical problem. In the meantime, we will continue performing RSA+LD/TM in CLEER patients with no Tm, given the data from the aforementioned references.

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Pascal Boileau, MD, PhD
Mikaël Chelli, MD, MSc
ICR (Institute for Sports & Reconstructive Bone & Joint Surgery)
Nice, France
E-mail: boileau.p@wanadoo.fr

Tyler R. Johnston, MD
UC Irvine Medical Center
Orange, CA, USA

Gabriel Cardenas, MD
Instituto de Seguridad del Trabajo
Santiago, Chile

Marc-Olivier Gauci, MD, PhD
Institut Universitaire Locomoteur et du Sport
University Hospital of Nice
Nice, France

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