

# C1. Eligibility Criteria

Reviewer ID: \_\_\_\_\_ Date: / /2009 Ref ID: \_\_\_\_\_

CRITERIA	Yes	No	Unclear
<b>1. PUBLICATION TYPE</b>			
a. Report of primary research	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
b. Published in 1990 or later	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
c. English language, except for nonoperative or postoperative rehabilitation	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<b>2. STUDY DESIGN</b>			
a. Enrolled ≥ 11 participants	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
b. One of the following designs (circle design):	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
i. RCT			
ii. CCT			
iii. Cohort			
iv. Case control			
v. Cross sectional			
vi. <i>Prospective</i> before-and-after (baseline data required)			
<b>3. POPULATION</b>			
a. >80% adult patients (≥18 years) [exclude pediatric, in vitro, cadaver]. Exclude professional athletes.	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
b. Partial- or full-thickness (including massive) RCT, confirmed using imaging (e.g. arthrography, ultrasound, MRI, etc) or intraoperative findings. [Exclude diagnosis based on physical exam/ history only]	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
c. Primary intention is treatment of RCT. Exclude if patients have RA or other inflammatory arthritis† (not OA), or are undergoing revision of failed RCT.	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<b>4. INTERVENTION (One of:)</b>			
a. Operative approaches: open, mini-open or arthroscopic repair, debridement or decompression [Exclude tendon transfers, arthroplasty, pain management]	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
b. Nonoperative intervention for treatment of RCT.			
c. Postoperative rehabilitation following RC repair.			
<b>5. OUTCOME</b>			
a. Numeric data reported on at least one of: quality of life, disability, time to return to work / activities, shoulder pain, range of motion, strength, adverse events.	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
b. Operative studies: Minimum 12 month follow-up for at least one outcome of interest [No restriction for nonoperative]	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Comments:

**REVIEWER'S DECISION :** Include  Exclude  Unsure

**FINAL DECISION:** Include  Exclude  Unsure

**NOTE:** To exclude must have said "NO" for at least one of 1-5.

**RELEVANT TO QUESTION(S):**

- 1. Does **early** surgical repair **compared to late** surgical repair (i.e., nonoperative intervention followed by surgery) lead to improved patient-important outcomes? [check only if study directly compares early vs. late]
- 2. What is the comparative effectiveness of **operative approaches**?
- 3. What is the comparative effectiveness of **nonoperative interventions**?
- 4. Does **operative repair vs. nonoperative treatment** lead to improved outcomes?
- 5. What are the associated **adverse effects** of operative and nonoperative therapies?
- 6. Which **prognostic factors** predict better outcomes? (specify)