Peripheral TFCC Tears Cause Ulnocarpal Instability: a Biomechanical Study

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INTRODUCTION:
Instability at the ulnocarpal joint can result from a number of etiologies, but the common thread among these causes for ulnocarpal instability (UCI) is often the presence of pathology in the triangular fibrocartilage complex (TFCC). To our knowledge, the clinical experience reported with UCI has not yet been substantiated by biomechanical evidence that peripheral TFCC tears result in UCI. The aim of the current study is to determine whether disruption of the peripheral TFCC causes ulnocarpal instability. Based on our clinical experience, we hypothesize that an arthroscopically-created tear of the articular disk of the TFCC will cause a significant decrease in ulnocarpal stability on mechanical testing.

METHODS:
24 fresh-frozen human upper extremities with intact elbow to the hand were held with the elbow in 90° flexion, the forearm in pronation and wrist in neutral on a servo-hydraulic loading machine. (Figure #1). A screw in the distal ulna was fixed to the loading actuator and a vertical load was sinusoidally cycled at ¼ Hz while the vertical movement was measured. From the load-displacement curve a range of dorsal-volar motion with minimal resistance (DVROM) was identified and the slope of the curve at the ends of the DVROM were measured and termed the ulnocarpal stiffness. A standardized 3 mm lesion of the ulnar-sided peripheral TFCC was then created via a single incision with a #15 scalpel blade under direct arthroscopic visualization. (Figure #2). The loading regimen was repeated after the TFCC lesion.

RESULTS:
The mean stiffness of the specimens was significantly decreased after creation of a peripheral tear in the articular disk of the TFC (p=0.003). The change in DVROM after the articular disk tear was not significantly different (p=0.08), see Table 1.

<table>
<thead>
<tr>
<th>N=20 specimens</th>
<th>Mean</th>
<th>SD</th>
<th>Paired t-Test</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stiffness: baseline</td>
<td>4.86 N/mm</td>
<td>3.68 N/mm</td>
<td></td>
</tr>
<tr>
<td>Stiffness: post tear</td>
<td>3.14 N/mm</td>
<td>2.47 N/mm</td>
<td>p=0.003*</td>
</tr>
<tr>
<td>DVROM: baseline</td>
<td>1.82 mm</td>
<td>0.63</td>
<td></td>
</tr>
<tr>
<td>DVROM post tear</td>
<td>2.33 mm</td>
<td>1.38</td>
<td>p=0.08**</td>
</tr>
</tbody>
</table>

Table 1: *Lower stiffness indicates more instability **Higher value of neutral zone indicates more instability

DISCUSSION cont.:
While the TFCC is composed of many contiguous structures, we chose only to transsect the peripheral attachment of the articular disk of the TFCC, representative of a Palmer 1B lesion. Further transection of the ulnocarpal ligaments and the other remaining components of the TFCC in the current experimental model would likely propagate the destabilization of both the UCJ and DRUJ, following the spectrum of instability proposed by Melone and Nathan. The current study is the first to establish biomechanical instability at the ulnocarpal joint after a TFCC lesion, particularly at the peripheral attachment of the articular disk. While these findings may seem intuitive in view of prior anatomic and clinical speculations, these results will help to define the pathomechanics underlying UCI. The presence of UCI after an isolated lesion of the articular disk suggests that both the distal radioulnar and ulnocarpal joints should be evaluated and accounted for in treatment strategies for patients with pathology of the TFCC, even if the ulnocarpal ligaments, ulnar collateral ligament, and ECU subsheath appear unaffected on advanced imaging modalities. Seeing that many surgeons are currently performing DRUJ reconstruction to treat patients with irreparable TFCC tears, one should evaluate the stability of the UCJ after reconstruction is complete and consider utilizing an adjunctive procedure to stabilize the UCJ if necessary.

SIGNIFICANCE:
A TFCC lesion at the peripheral attachment of the articular disk can cause ulnocarpal instability. Both ulnocarpal and radioulnar stability should be evaluated when assessing TFCC repair techniques and reconstructive procedures, even in the absence of pathology in the other soft tissue components of the TFCC.

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