

Anatomic Ligament Repair Restores Ankle and Syndesmotiic Rotational Stability as Much as Syndesmotiic Screw Fixation

Name Credentials

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Purpose: Unstable rotational ankle fractures with concomitant syndesmotiic disruption commonly occur. Currently, there is disagreement about the optimal treatment method for the syndesmotiic injury component in this patient population. While transsyndesmotiic screws are most often used, alternative syndesmotiic stabilization strategies such as syndesmotiic suture fixation and anatomic ligament repair have garnered increased interest. To date, no study has investigated the ability of an anatomic ligamentous repair strategy, consisting of posterior inferior tibiofibular ligament (PITFL) and deltoid ligament repair, to restore ankle and syndesmotiic stiffness. Our hypothesis was that the anatomic ligament repair strategy would provide equivalent ankle and syndesmotiic stiffness compared to a single 3.5-mm transsyndesmotiic screw.

Methods: Nondestructive external rotation stresses of 4 nM were applied to 8 cadaveric limbs using a hydraulic loading frame. Four conditions were tested using a repeated-measures design: intact and three repair conditions following a destabilizing ligamentous ankle injury with syndesmotiic disruption. The three repair conditions were tricortical transsyndesmotiic screw fixation, PITFL repair, and combined PITFL and deltoid ligament repair. External rotation of the ankle and syndesmosis were measured using a motion capture system and compared for each test condition. Repeated-measures one-way analysis of variance (ANOVA) statistical tests were performed to compare the ankle and syndesmotiic rotation findings between the three repair conditions and intact condition.

Results: Ankle stability was not fully restored by any of the three repair constructs. The intact ankle externally rotated approximately half as many degrees as the three repair conditions (Intact 10.9, Transsyndesmotiic screw 17.0, PITFL 21.4, and PITFL/deltoid 15.6). Direct comparison between the transsyndesmotiic screw and PITFL/deltoid repair specimens was not significantly different ($P = 0.84$). The intact condition demonstrated significantly fewer degrees of syndesmotiic rotation than the repair constructs (Intact 2.4, Transsyndesmotiic screw 5.2, PITFL 8.5, and PITFL/deltoid 6.9). Direct comparison between the transsyndesmotiic screw and PITFL/deltoid repair groups was not significantly different ($P = 0.21$). Each of the repair constructs resulted in an externally rotated fibula compared to the intact condition prior to external rotation testing. The soft-tissue repairs (PITFL 4.3°, PITFL/deltoid 3.9°) caused twice as much external rotation compared to the syndesmotiic screw (1.9°).

Conclusion: We found that combined repair of the PITFL and deltoid ligament restores an equivalent amount of ankle and syndesmotiic rotational stability as transsyndesmotiic screw fixation. Based on our findings, ligamentous repair can potentially be a viable treatment alternative in unstable ankle fracture patients with syndesmotiic disruption. However, the

clinical implications of a 2° greater fibular external malrotation with the ligamentous repair constructs compared to a single trans-syndesmotic screw is not known. Clinical outcome studies are needed to verify our biomechanical findings.

Table I. Summary and comparison of ankle and syndesmotic external rotation stability as well as resting fibular external rotation between intact and repair conditions

	Mean (n=8)	95% Confidence Intervals	P-value
Ankle Stability (degrees)			
Intact	10.9	6.4 – 15.4	-
Screw	17.0	11.4 – 22.6	0.008
PITFL	21.4	14.8 – 28.0	<0.001
PITFL+Deltoid	15.6	10.8 – 20.4	0.05
Syndesmotic Stability (degrees)			
Intact	2.4	1.6 – 3.2	-
Screw	5.2	3.6 – 6.8	0.013
PITFL	8.5	6.0 – 11.0	<0.001
PITFL+Deltoid	6.9	5.2 – 8.6	<0.001
Fibular External Rotation (degrees)			
Intact	-		-
Screw	1.9	1.0 – 2.9	0.016
PITFL	4.3	3.0 – 5.5	<0.001
PITFL+Deltoid	3.9	2.7 – 5.1	<0.001

The FDA has stated that it is the responsibility of the physician to determine the FDA clearance status of each drug or medical device he or she wishes to use in clinical practice.