

# The Clinical and Biomechanical Performance of All-Suture Anchors: A Systematic Review



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**Purpose:** This systematic review aimed to clarify the relative strengths and weaknesses of the all-suture anchors (ASAs) in both clinical and experimental studies. Our hypothesis was that there would be similar clinical and experimental data for ASAs regarding the biomechanical properties, clinical outcomes and complication rates. **Methods:** A systematic review of MEDLINE and Embase databases was performed. The inclusion criteria for clinical studies were both retrospective or prospective study design and minimum 1-year follow-up; for biomechanical studies, the inclusion criteria were performance on either cadaver and animal bones or synthetic surfaces. Studies were excluded if the studies were not in English or if they were review articles, commentaries, letters, case reports, or technical notes. The risk of bias assessment was done using the Methodological Index for Non-randomized Studies (MINORS) tool. **Results:** We included 13 experimental and 3 clinical studies. The least displacement under cyclic loading was recorded with Q-Fix. Failure mode was mostly by suture breaking for the Q-Fix, whereas anchor pullout was the most common for the others. Cadaver humerus' greater tuberosity seemed to be less durable for the ASAs. Tests on cadaver glenoid showed similar biomechanical properties when compared to a control anchor. Studies investigating clinical and radiologic findings were very few, and only 3 case series were included in this review. Clinical findings of patients treated with ASAs for instability and rotator cuff repair showed satisfactory results and little increase in the complication rate (retear or revision surgery because of loose anchor). **Conclusions:** ASAs have similar or better biomechanical properties compared to regular anchors. Low-profile design seems to be an important advantage. Case series can not distinguish between the possible clinical benefits and/or risks. **Clinical Relevance:** ASAs have similar biomechanical properties when compared with other types of anchors. Their strength and performance vary with anatomic location, which may influence clinical success.

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For the purpose of reattachment of tendons and other soft tissues to bone, suture anchors are widely used with minimally invasive techniques, especially in the upper extremity. These devices have been improved over time, with the ability to accommodate multiple sutures, the use of biodegradable and biocomposite materials and

knotless designs. Suture anchors made completely of suture material were developed in the past decade. These all-suture anchors (ASAs) are based on ultrahigh molecular-weight polyethylene-containing sutures. The anchor portion of the device typically consists of a sleeve or tape, also made from suture material, through which the ultrahigh molecular-weight polyethylene-containing suture is woven. When the ASA is inserted into bone and the suture limbs are pulled, the sleeve or tape is cinched up to compress against the overlying cortical bone creating a "ball," which serves as the anchor. Potential benefits include decreased bone damage. They are radiolucent and nondegradable. ASAs used for the glenoid are smaller and usually have 1 or, at most, 2 sutures, whereas those used for rotator cuff tendon repairs are larger and are either double- or triple-loaded.<sup>1,2</sup>

Biomechanical testing reveals the failure mode and strength of ASAs and shows that they have sufficient strength for soft tissue-to-bone healing in the majority of clinical settings.<sup>3</sup> However, most biomechanical studies are in vitro and done at time 0, and only a few in vivo animal studies, which may not reflect the

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human clinical response, are available.<sup>3-15</sup> A literature search of clinical outcomes and complications related to ASAs discovered relatively few studies.<sup>16-18</sup>

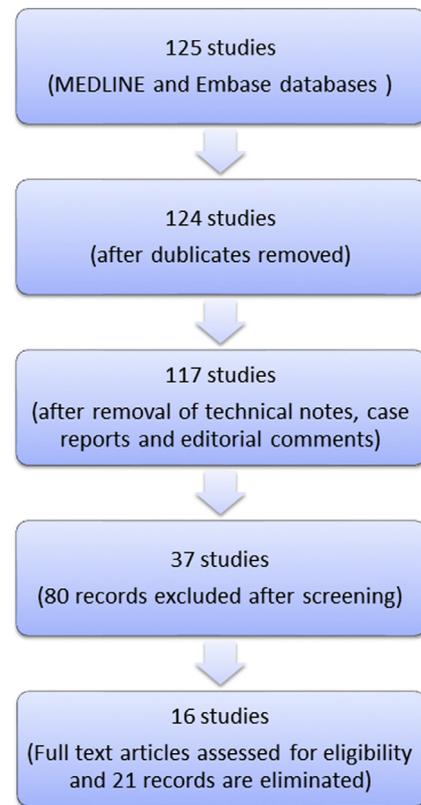
This study sought to address the following questions: (1) Because it is not feasible to obtain ASA biomechanical data clinically, is that high failure strength consistent with the clinical experience?; (2) Is tunnel widening or cyst formation previously reported in animal studies reported clinically?<sup>7</sup>

This systematic review focused on biomechanical test results and clinical outcomes of different sizes and models of ASAs used for soft tissue-to-bone fixation. Our hypothesis was that clinical outcomes and complication rates of ASAs are correlated with their biomechanical test results.

## Methods

This study was performed according to the Preferred Reporting Items for Systematic reviews and Meta-analyses (PRISMA).<sup>19</sup> MEDLINE and Embase databases were searched for published randomized or nonrandomized controlled studies and prospective or retrospective case series presenting the clinical results of soft-tissue repairs using ASAs and also experimental studies testing the biomechanical properties of ASAs in cadavers and animals and on synthetic surfaces. We independently searched the databases for the key phrases “all-suture anchor” OR “ASA.” All articles that were identified using these search terms on February 15, 2019, were then reviewed and discussed among the authors, and a decision whether to include or exclude them was made according to inclusion and exclusion criteria. The inclusion criteria for clinical studies consisted of both retrospective and prospective study design and any ligament or tendon-to-bone repairs by ASAs with a minimum 1-year follow-up period. Evaluations were expected to be done by clinical-outcome measuring, radiologic findings and complication rates. The inclusion criteria for biomechanical studies consisted of all studies testing the biomechanical performance of ASAs on either cadaver and animal bones or synthetic surfaces. Test protocols might include soft tissue-to-bone repair models or the general performance of these anchors.

Exclusion criteria for this review were non-English manuscripts, review articles, commentaries, letters, case reports, and technical notes. For studies in which a kin relationship was identified, manuscripts were joined to make a single record so as to avoid repeating the same findings. One investigator selected studies for inclusion in the review. Articles were then independently assessed by a second investigator, who confirmed eligibility. The following variables were recorded in a predesigned database: general manuscript characteristics (author, year and journal), study design, testing protocols and findings for biomechanical studies,



**Fig 1.** PRISMA flow diagram. From the initial 125 records, 37 full-text articles were reviewed for eligibility. Ultimately, 16 studies (13 experimental, 3 clinical and radiologic studies) were included.

imaging findings (tunnel widening, cyst formation), and clinical outcomes and complications (failure modes) for the clinical studies.

Biomechanical study data were reported quantitatively for cyclic loading, ultimate load to failure, displacement at failure, and stiffness-test results. Failure mode was qualitatively reported as anchor pullout, suture breaking, anchor breaking, and tendon tearing.

The quality of all included studies was independently evaluated using the Methodological Index for Non-randomized Studies (MINORS) tool<sup>20</sup> (Appendix Table 1).

## Results

The initial database search retrieved 125 titles and abstracts. After screening records for duplicates and inclusion criteria, 37 full-text articles were reviewed for eligibility. Ultimately, 16 studies (13 experimental studies, 3 clinical and radiologic studies) were included. The levels of evidence for the all 3 clinical studies were Level IV. The study’s flow diagram is shown in Fig 1.

The mean quality rating of all biomechanical studies using the Methodological Index for Non-randomized Studies MINORS tool was  $20.5 \pm 1.2$  points (of a maximum of 24 points, 85.2%; range 19-22), mean

**Table 1.** Characteristics of the Included Biomechanical Studies

	Author	Purpose of the study	Study design	Sample size	Test protocol
Studies of glenoid	Barber FA 2013 <sup>3</sup>	General performance of suture anchors.	Biomechanical performance on porcine distal femur metaphysis a. Cortex intact b. Decorticated bone	n = 10-14	1. Cyclic loading 10-100N (100 and 200 cycles) 2. Ultimate load to failure 3. Displacement at failure 4. Stiffness 5. Mode of failure
	Barber FA 2017 <sup>4</sup>	General performance of ASAs.	Biomechanical performance on porcine cortical bone and biphasic PU foam.	n = 10-20	1. Cyclic loading 10-100N (100 and 200 cycles) 2. Ultimate load to failure 3. Displacement at failure 4. Stiffness 5. Mode of failure
	Dwyer 2014 <sup>5</sup>	Pretensioning the ASA for better performance	Biomechanical performance on bovine proximal tibia and human cadaver glenoid (control: bioabsorbable screw anchor: Bio Mini Revo 3.1 mm)	n = 8	1. Displacement at 50N 2. Ultimate load to failure 3. Stiffness 4. Mode of failure
	Pfeiffer 2014 <sup>6</sup>	To examine histologic characteristics during the healing period and to compare immediate biomechanical characteristics of ASA and biocomposite anchor in human cadaveric glenoid	Histologic response, in vivo study in dogs Biomechanical performance on cadaver glenoid (control 2.4-mm BioComposite SutureTak)	n = 6 (dogs, shoulder) n = 8 (human cadaver glenoid)	1. Cyclic loading 5-25N (100 cycles) 2. Ultimate load to failure 3. Displacement at failure 4. Mode of failure
	Douglass 2017 <sup>7</sup>	Performance of ASAs on simulated acetabular and glenoid bone	Biomechanical performance on monophasic PU foam (30 pcf for acetabular, 20 pcf for glenoidal bone) (Control BioRaptor 2.3 PEEK)	n = 7-11	1. Cyclic loading 10-50 N ( 200 cycles), 10-100 N (additional 200 cycles) 2. Ultimate load to failure 3. Displacement at failure 4. Stiffness 5. Mode of failure
	Erickson 2017 <sup>8</sup>	Performance on glenoid bone	Biomechanical tests on cadaver glenoid (control PEEK anchor 2.3 mm Bioraptor)	n >30	1. Load to 2 mm displacement 2. Ultimate load to failure
	Ruder 2018 <sup>9</sup>	Effect of drill length (21 mm/17 mm/13 mm) on biomechanical performance of ASA (JuggerKnot 1.5)	Biomechanical performance on cadaver glenoid	n = 32	1. Cyclic loading 10-60 N ( 200 cycles), 2. Ultimate load to failure 3. Mode of failure (catastrophic/clinical)
Studies of biceps tenodesis	Chiang 2016 <sup>10</sup>	Performance of ASA (Y knot) on biceps tenodesis	Biomechanical performance on subpectoral biceps tenodesis. cadaver humerus (control group: interference screw)	n = 8	1. Cyclic loading 5-70 N (500 cycles) 2. Ultimate load to failure 3. Displacement at failure 4. Stiffness 5. Mode of failure
	Hong 2018 <sup>11</sup>	Performance of ASA (Y knot) on biceps tenodesis	Biomechanical performance on suprapectoral biceps tenodesis. cadaver humerus (control group: interference screw)	n = 8	1. Cyclic loading 5-70 N (500 cycles) 2. Ultimate load to failure 3. Displacement at failure 4. Stiffness 5. Mode of failure

(continued)

Table 1. Continued

Author	Purpose of the study	Study design	Sample size	Test protocol
Galland 2013 <sup>12</sup>	Performance of ASAs on rotator cuff repair (2 Juggerknot 1.4mm vs One 5.5 mm screw anchor)	Biomechanical performance on bovine humerus greater tubercule	n = 15	1. Ultimate load to failure 2. Displacement at failure 3. Mode of failure
Goschka 2015 <sup>13</sup>	Performance of ASAs on double row rotator cuff repair	Biomechanical performance on human cadaver shoulder	n = 9	1. Cyclic loading 10-100 N (500 cycles) 2. Load to 5 mm displacement 3. Ultimate load to failure 4. Stiffness 5. Mode of failure
Nagra 2017 <sup>14</sup>	Performance of ASAs on rotator cuff repair	Biomechanical performance cadaver human shoulder (control group Twinfix PEEK)	n = 5	1. Cyclic loading 10-180N (200 cycles) 2. Ultimate load to failure 3. Displacement at failure 4. Mode of failure
Oh 2018 <sup>15</sup>	Effect of insertion and traction angle on performance of ASAs for rotator cuff repair.	Biomechanical performance on biphasic PU and cadaver humerus.	n = 5 (for PU) n = 9 (for cadaver)	1. Stiffness 2. Ultimate load to failure 3. Yield load

ASA, all-suture anchor; PU, polyurethane.

quality rating of all clinical studies was  $13 \pm 1$  (of a maximum of 16 points, 81.2%; range 12-14) (Appendix Table 2). The mean quality rating of all 16 studies was 84.5%. The most common reason for point deductions was item 5, unbiased assessment of the study's endpoint. All except 2 studies (Pfeiffer et al.,<sup>6</sup> Willemot et al.<sup>16</sup>) received 0 points for this item because the observers were not blinded to the study's endpoints.

Studies to measure biomechanical performance of ASAs are performed on various bone models: porcine cortical bone and distal femur metaphysis,<sup>3,4</sup> bovine proximal tibia,<sup>5</sup> bovine humerus greater tubercle,<sup>12</sup> monophasic polyurethane (PU) foam,<sup>4,7</sup> biphasic PU foam,<sup>15</sup> cadaver glenoid,<sup>8</sup> cadaver greater tubercle,<sup>13</sup> and cadaver humerus to test biceps tenodesis.<sup>10,11</sup> It is obvious that bone models and test protocols are not standardized, and both show high variability. Biomechanical test protocols measured the displacement under cyclic loading, displacement at failure, ultimate load to failure, stiffness, mode of failure, and load to 2 mm or 5 mm displacement (Table 1). Most of the biomechanical studies had a control suture anchor (screw anchor or polyetheretherketone [PEEK] anchor), and the results were compared.

Clinical studies were reviewed, and outcome data were reported according to clinical scores acquired from the Visual Analog Scale, the Western Ontario Shoulder Instability Index, the Disabilities of the Arm, Shoulder and Hand Score, and the Constant and Murley scoring systems. Imaging findings and complication rates were also reported in those clinical studies (Table 2).

ASA models used in the included studies were Juggerknot 1.4, 1.5 and 2.9 mm (Biomet); Iconix 1, 2 and 3 (Stryker); Y-Knot 1.3, 1.8 and 2.9 mm (Conmed); Draw tight 1.8 and 3.2 mm (Parcus); Q-fix 1.8 and 2.8 mm, Suture Fix Ultra 1.7 mm (Smith & Nephew); Omega Knot (ARC, Korea) (Table 2).

Cyclic loading tests showed that the least displacement was recorded with Q-Fix anchors. Also, failure mode was mostly by suture breaking for the Q-Fix, whereas anchor pullout was the most common for the others (Table 3). Four biomechanical studies on cadaver glenoid bone compared the ASAs with a control bioabsorbable, biocomposite or PEEK anchor. In 2 of these studies, control anchors showed better results than ASAs (Y-Knot and Juggerknot),<sup>5,6</sup> but in the remaining 2 studies, ASAs (Suture-Fix Ultra and Q-Fix) showed better results than the control PEEK anchors despite their larger diameters than ASAs.<sup>7,8</sup> Two studies simulating biceps tenodesis on cadavers both showed superior biomechanical results for control screw anchors when compared to ASAs (Y-Knot).<sup>10,11</sup> Simulating the rotator cuff repair, 3 biomechanical studies compared ASAs with a control

**Table 2.** Characteristics of the Included Clinical Studies

Study	Purpose of the study	Study design and Level of Evidence	Sample size	Method
Willemot 2016 <sup>16</sup>	Radiologic and clinical outcomes arthroscopic labral repairs with ASAs.	Postoperative MRI scan DASH, Constant and WOSI Level of Evidence: IV	n = 20 patients 58 anchors	Radiologic appearance of bone at the anchor site was judged by the presence of cyst formation, tunnel widening (> 2 mm) or bone edema. Clinical scores
Byrd 2017 <sup>17</sup>	Incidence of intraoperative pull-out of ASAs for acetabular labral repairs.	Retrospective review of intraoperative anchor failure incidence Level of Evidence: IV	434 patient 2007 anchors	(1) The age and gender of all cases; (2) the number of cases in which labral repair was performed; (3) the number of anchors used; (4) the number of cases in which intraoperative anchor failure occurred; (5) the number of anchors that failed; and (6) the age and gender of those cases in which anchor failure occurred
Van Der Bracht 2018 <sup>18</sup>	Clinical and radiologic (at least 1 year later) study to investigate the feasibility and safety of ASAs in arthroscopic rotator cuff repair	Prospective cohort; VAS score (for pain and satisfaction), constant Murley score, strength of SS muscle by dynamometer and radiologic findings Level of evidence: IV	n = 20 patients 48 anchors	Integrity of the cuff repair, cyst formation around anchor, ingrowth of the bone into the anchor, and integrity of the bone tunnel border were evaluated. Clinical scores

ASA, all-suture anchor; WOSI, Western Ontario Shoulder Instability Index; DASH, Disabilities of the Arm, Shoulder and Hand score; VAS, Visual Analog Scale.

**Table 3.** Test Results of the Biomechanical Studies

Studies of	Author	Anchor	Cyclic loading	Ultimate load to failure	Displacement at failure	Stiffness N/mm	Mode of failure	Result
Glenoid	Barber FA 2013 and 2017 <sup>3,4</sup>	JuggerKnot 1.4 – 1.5 – 2.9	Porcine bone: 1.5: 1.39 mm 2.9: 1.44 mm	Porcine bone: 239N – 290N – 519N	Porcine bone: 0.22 mm – 0.22 mm – 0.22 mm	198 – 57 – 76	1.4: Anchor pullout and Suture break 1.5 – 2.9: Suture break only	Ultimate load at failure was correlated directly with the number of sutures. Y-Knot demonstrated greater displacement than the JuggerKnot and Q-Fix
		Iconix 1- 2 - 3	Porcine bone: 1.87 mm – 1.55 mm – 1.44 mm	Porcine bone: 209N – 469 N – 570N Biphasic PU foam: 235N – 520N	Porcine bone: 0.31 mm – 0.23 mm – 0.20 mm Biphasic PU foam: 0.23 mm – 0.43 mm	65 – 83 – 89	1: Anchor and suture break 2: Mostly suture break 3: Mostly anchor break	Both JuggerKnot (81%) and Q-Fix (97%) anchors failed predominantly by the suture breakage; however, Y knot had high anchor pull out rate.
		Y knot 1.3 – 1.8 – 2.9	Porcine bone: 2.4mm – 2.0mm – 3.52mm	Porcine bone: 250N – 477N – 603N Biphasic PU foam: 152N – 531N – 657N	Porcine bone: 0.45 mm – 0.33 mm – 0.55 mm Biphasic PU foam: * - 0.23 mm – 0.19 mm	65 – 74 – 84	All: High anchor pullout rate	None of the all-suture anchors reached a clinically significant 5 mm displacement during cyclic loading.
		Draw Tight 1.8 – 3.2	Porcine bone: 2.12 mm – 2.62 mm	Porcine bone: 290N – 418N Biphasic PU foam: 263N – 191N	Porcine bone: 0.30 mm – 0.30 mm Biphasic PU foam: 0.24 mm – 0.20 mm	41 – 49	All: High anchor pullout rate	
		Q Fix 1.8 – 2.8	Porcine bone: 1.22 mm – 1.58 mm	Porcine bone: 346N – 495N Biphasic PU foam: 292N – 495N	Porcine bone: 0.19 mm – 0.23 mm Biphasic PU foam: 0.11mm – 0.16mm	55 – 57	All: Suture break (1 anchor pullout)	
Dwyer 2014 <sup>5</sup>		YKnot 1.3 mm (Handset) (bovine tibia/ human glenoid)	-	140N / 91 N	Displacement at 50N: 4.6mm / 7.5mm	8.7 / 4.3	The primary mode of failure in all-suture anchors was anchor pullout	Pretensioning the YKnot to 60 N ensures that the anchor is well fixed, consistently eliminating laxity and displacement in both high-density bovine and lower-density cadaveric bone
		YKnot (60 N Pretensioned) (bovine tibia/ human glenoid)	-	135N / 145 N	Displacement at 50N: 1.9mm / 1.9mm	21 / 21.7	The primary mode of failure in all-suture anchors was anchor pullout	Some anchor pullout was seen below the 60 N set as the pretensioning force
		Bio Mini Revo 3.1 mm (control)	-	206N/107 N	Displacement at 50N: 3.5mm / 2.7mm	12.8 / 14.4	6 anchor pullout, 2 eyelet failure	
Pfeiffer 2014 <sup>6</sup>		JuggerKnot 1.4 mm	2.9 mm	141 N	13.7mm	-	Anchor pullout (8)	Consistent cavity formation, significant
		Control 2.4-mm BioComposite SutureTak	1.3 mm	136.7 N	3.2mm	-	Anchor pullout (5) and breakage of the top portion of the anchor (3)	expansion of the drill tunnel, was associated with the JuggerKnot anchors in the canine glenoid. All sutures incite foreign body reactions to varying degrees wherever they are placed in the body JuggerKnot anchor was slipping

(continued)

Table 3. Continued

Author	Anchor	Cyclic loading	Ultimate load to failure	Displacement at failure	Stiffness N/mm	Mode of failure	Result
Douglass 2017 <sup>7</sup>	JuggerKnot 1.4 – 1.5 – 2.9	-	20 pcf: JuggerKnot 2.9: 194N 30pcf: JuggerKnot2.9: 301N	-	-	20 pcf:Anchor pullout 30 pcf: Anchor pullout	within the prepared hole before failure. Clinically, such slipping leads to a loss of reduction and can hinder healing. Based on the biomechanical findings in human bone and histologic findings in canine subjects, ASAs may be at risk for clinical failure ASAs exhibited less displacement and greater maximum loads in higher density (30pcf) bone substitute.
	Iconix 1- 2 – 25 - 3	-	20 pcf: Iconix 2: 163N Iconix 25: 196N (highest) Iconix 3: 180N 30 pcf: Iconix 25: 307N Iconix 3: 276N	-	-	<b>20 pcf:</b> Anchor pullout <b>30 pcf:</b> Anchor pullout	The cyclic displacement and maximum load of ASAs vary widely depending on anchor design and bone density Q-Fix 1.8, however, performed better than all other anchors in displacement and had maximum failure loads comparable with the highest values of the other anchors tested.
	Y-Knot 1.3 – 1.8	-	20 pcf: YKnot 1.8: 176N	-	-	<b>20 pcf:</b> Anchor pullout <b>30 pcf:</b> Anchor pull out	Failure mode is mostly anchor pullout, probably due to monophasic structure of the PU substitute. But this situation changed with Q fix anchor on 30pcf substitute (suture break)
	Suture Fix Ultra 1.7	-	-	-	-	<b>20 pcf:</b> Anchor pullout <b>30 pcf:</b> Anchor pull out	
	Q Fix 1.8	Least	30 pcf: 291N	Best result (least displacement) in both 20pcf and 30pcf	-	<b>20 pcf:</b> Anchor pullout <b>30 pcf:</b> Suture break	
	Bioraptor PEEK 2.3mm (Control)	0.3 mm (20 pcf) 0.4 mm (30 pcf) (200 cycles)	-	-	-	-	-
Erickson 2017 <sup>8</sup>	JuggerKnot 1.4	-	171.5 N	Load to 2 mm displacement: 36 N	-	-	A second-generation all-soft suture anchor (Suture Fix Ultra) showed greater loads to 2 mm of displacement than a first-generation all-soft suture anchor (JuggerKnot).
	Suture Fix Ultra 1.7	-	182.5 N	Load to 2 mm displacement: 42 N	-	-	Both all-soft suture anchors had higher load to failure than PEEK anchor (Bioraptor 2.3 PK).
	Bioraptor PEEK 2.3 mm (control)	-	132N	Load to 2 mm displacement: 39 N	-	-	
Ruder 2018 <sup>9</sup>	JuggerKnot 1.5	21 mm: 2.50 17 mm: 1.70 13 mm: 1.13	21 mm: 194 N 17 mm: 190 N 13 mm: 138 N	-	-	High clinical failure (3mm and 5mm) with 21mm depth. No clinical failure with 13mm.	Inserting the anchor at a depth of 17mm reduced the displacement after cyclic loading without reducing the ultimate load to failure.

(continued)

Table 3. Continued

	Author	Anchor	Cyclic loading	Ultimate load to failure	Displacement at failure	Stiffness N/mm	Mode of failure	Result
<b>Studies on Biceps tenodesis</b>	Chiang 2016 <sup>10</sup>	Y-Knot 1.3mm / Milagro Bioreplaceable Screw 8 × 23 mm	8.1 mm / (Control 3.4 mm)	239N/(control 254.4 N)	20.3 mm/ (control 13.3 mm)	26/(control 27.7)	ASA: Anchor pullout (n: 4), Tendon tear (n: 4) Control: (screw pullout (2), tendon tear (6))	ASA technique displayed values of ultimate failure load and stiffness comparable to that of the interference screw technique. However, the cyclic and failure displacement values of the interference screw trials were significantly less than that of the ASA
	Hong 2018 <sup>11</sup>	Y-Knot 1.3 mm / Milagro Bioreplaceable Screw 8 × 23mm	6.4mm/(control: 3.7 mm)	186.6 N/(control 203.8 N)	16.3 mm / (control 13.3 mm)	26.1 / (control: 27.1)	ASA: Anchor pullout (n:6), Tendon tear (n:2) Control: (screw pullout (0), Tendon tear (8))	ASA technique displayed values of ultimate failure load, failure displacement and stiffness comparable to that of the interference screw technique. However, the cyclic displacement values of the interference screw trials were significantly less than that of the ASA
<b>Studies of rotator cuff repair</b>	Galland 2013 <sup>12</sup>	JuggerKnot 1.4 mm X2 / (Control Screw Anchor 5.5 mm)	-	ASA (X2): 265N /control: 325N (P: 0.09)	ASA (X2): 23 mm /control: 21 mm (P:0.46)	-	ASA: 12 anchor pullout, 1 thread fracture/control screw anchor: 8 anchor pullout, 5 eyelet fractures	There was no statistically significant difference between pullout strength and displacement of a double-fixed bone ASA and a single-fixed control SA
	Goschka 2015 <sup>13</sup>	Medial row/ lateral row Iconix2/Iconix2	-	313.2N	-	-	Anchor pullout: 5 Suture tear: 3 Knot slippage:1	The biomechanical performance of anchor configurations using the ICONIX2 would be comparable to that of the configuration of solid-body anchors, as no significant differences were found between groups for any metric tested.
		Iconix2 / ReelX 3.9	-	457.9N	-	-	Anchor pullout: 6 Suture tear: 2 Suture pull out of anchor: 1	(No significant change in between groups for all tests)
		Iconix2/ReelX 4.5	Max gap formation in anterior anchors Min gap formation in posterior anchors	420.2N	-	-	Anchor pullout: 6 Suture tear: 1 Suture pullout of anchor: 1 Eyelet break:1	
	Control: CorkScrew4.5/ SwiveLock 4.75	Max gap formation in posterior anchors Min. gap formation in anterior anchors	430.9 N	-	-	Anchor pullout: 4 Suture tear: 1 Suture pullout of anchor: 2 Muscle tear: 2		
		100 cycles		137.8 N	33.7 mm			

(continued)

Table 3. Continued

Author	Anchor	Cyclic loading	Ultimate load to failure	Displacement at failure	Stiffness N/mm	Mode of failure	Result
Nagra 2017 <sup>14</sup>	JuggerKnot	22.6 mm				Anchor pullout (4)	Mean load to failure values were significantly higher for the traditional anchor (181.0 N) compared with the ASAs (mean 133.1 N). The JuggerKnot anchor had greatest displacement at 50, 100 and 150 cycles, and at failure, reaching statistical significance over the control at 100 and 150 cycles (22.6 mm, and 29.6 mm). All ASAs showed substantial (> 5 mm) displacement between 50 and 100 cycles (6.2 to 14.3). ASAs predominantly failed due to anchor pullout (95% vs 25% of traditional anchors), whereas a higher proportion of traditional anchors failed secondary to suture breakage. Pullout strength was higher at the 45° than at the 90° traction angle (all $P < .05$ ) consistent with the deadman theory. Pullout strength was significantly higher at the 90° and 75° than at the 45° insertion angle in both high-density saw bones and cadaveric humeri (all $P < .05$ ). Pullout strength was not significantly different by ASA type (all $P > .05$ ). Ultimate load to failure and yield load at the posterior insertion point of the greater tubercle were significantly lower than at the anterior and middle insertion. Stiffness was also significantly lower at the posterior point than at the middle point.
	2.9mm					Suture failure (1)	
	Iconix 3	17.9 mm	103.9 N	22.7 mm		Anchor pull out (5)	
	Y-Knot 2.8mm	15.1 mm	145.8 N	23.6 mm		Anchor pull out (5)	
	Q Fix 2.8mm	11.8 mm	144.9 N	20.3 mm		Anchor pull out (5)	
Oh 2018 <sup>15</sup>	Control PEEK	12.5 mm	181 N	19.7 mm		Anchor pull out (1)	
	Twinix Ultra					Suture failure (2)	
	Omega Knot 2.9 mm		Maximum load on nonosteoporotic synthetic model: 45° traction, 75° insertion: 467.2N			Eyelet fracture (1)	
	Y-Knot 2.8 mm		45° traction, 90° insertion: 467.1N				

ASA, All-Suture Anchor.

**Table 4.** Test Results of the Clinical Studies

Study	Anchor/Fixation	Clinical outcome	Radiologic findings	Complications	Result
Willemot 2016 <sup>16</sup>	JuggerKnot 1.4 mm Labral repair	Satisfactory clinical results: WOSI: 70.6, DASH: 18.9 Constant: 89.3 No recurrence of instability.	None of the patients displayed large cyst formation. Small cysts (grade 3) were found in 2 patients (2 anchors). Tunnel widening (grade 2) was apparent in 3 patients (3 anchors) with an average widening of 3.3 mm (range 3-4 mm). Bone edema (grade 1) at the anchor-site was seen in 6 patients (8 anchors). The remaining 9 patients (45 anchors) did not display reactive bone changes.	No complications	Promising early radiographic and clinical outcome after arthroscopic glenohumeral labral repair using all-suture anchors. In this cohort of 20 patients with 58 anchors and a mean follow-up of 19 months, bone reactions were few and low grade on the postoperative MRIs, independent of anchor position. Clinical scores demonstrate satisfactory functional outcomes without recurrence of subluxation or dislocation.
Byrd 2017 <sup>17</sup>	Q Fix 1.8 mm Acetabular labral repair	—	—	A total of 33 anchors pulled out among 30 patients, representing a 1.6% incidence among all anchors. No statistically significant difference compared with the patient population in which no anchor pulled out. Most common reason for failure was to have the anchor securely imbedded in bone.	An overall failure rate of 1.6% seems quite acceptable.
Van Der Bracht 2018 <sup>18</sup>	JuggerKnot 2.9 mm Double row rotator cuff repair (Both ASA)	VAS for pain: 6.88 to 2.12 VAS for satisfaction: 9.18 Post op constant: 79.05 No difference in SS muscle strength between operated and nonoperated sides	1 retear, 5 small tears at musculotendinous junction Local fluid collection (not encapsulated) 10.4% (89.6% no fluid around anchor) No cyst formation (fluid diameter twice the anchor diameter) Thin, uninterrupted tunnel wall	1 deep wound infection One repeat arthroscopy 61 days later (1 loose lateral anchor)	No fluid could be detected between the anchors and the edge of the bony tunnel for 90% of the anchors. Full rotator cuff integrity was seen in 19 patients, with only 1 patient sustaining a retear. Clinical results are comparable with an arthroscopic rotator cuff repair using classic anchors. Study shows promising early radiographic and clinical results after arthroscopic rotator cuff repair using ASAs.

ASA, All-Suture Anchor; DASH, Disabilities of the Arm, Shoulder and Hand score; VAS, Visual Analog Scale; WOSI, Western Ontario Shoulder Instability Index.

screw or PEEK anchor. Although 2 of these studies<sup>12,13</sup> did not find any significant difference between groups, 1 study<sup>14</sup> found better biomechanical results for control PEEK anchor when compared to ASAs (Q-Fix, JuggerKnot, Iconix, and Y-Knot). Test results, especially the failure mode, relied on the bone model. The cadaver humerus greater tubercle seemed to be less durable with the ASAs than other bone models, and failure mode is more likely to be by anchor pull out (Table 3).

Studies investigating clinical and radiologic findings were very few, and only 3 case series (2 prospective, 1 retrospective) were included in this review (Table 2 and Table 4).<sup>16-18</sup> Each study had a different surgical repair modality for differing anatomic locations (gleno-humeral labral repair,<sup>16</sup> acetabular labral repair<sup>17</sup> and rotator cuff tendon repair<sup>18</sup>). Clinical failure was found to occur seldomly, anchor pullout was seen only in a rotator cuff repair study group in 1 patient in 1 anchor (n = 48). According to radiologic findings, cyst formation around the anchor was seen only in the gleno-humeral labral repair study group and included only 2 anchors (n = 58).

## Discussion

This systematic review found that ASAs have similar biomechanical properties when compared with metal screw, bioabsorbable, biocomposite, or PEEK anchors. However, only 3 level IV clinical studies were identified; consequently, insufficient data are currently available to draw conclusions about clinical outcomes.

With this systematic review, it has been demonstrated that the biomechanical performance of ASAs is affected primarily by the bone model. Synthetic PU foam is a commonly used model for ASA testing. Douglass et al. used 2 different monophasic PU foam models to test ASAs and found better biomechanical results with increasing density.<sup>7</sup> The Q-Fix, especially, showed less displacement with cyclic loading and changed the mode of failure from anchor pullout to suture breaking by increasing the PU density from 20 pound-force per cubic foot (pcf) to 30 pcf. Barber et al. advocated the usage of this biphasic artificial PU model to mimic cancellous (12 pcf PU foam) and cortical bone (fiber-filled epoxy coating).<sup>4</sup> They tested and compared different types of ASAs in the biphasic PU as well as in porcine femur cortical bone. They found similar test results for each type of ASA in both models. These results supported the use of biphasic PU models for experimental studies.

On the other hand, differing mechanical results were also reported in cadaver studies. Even in the same cadaver region, different insertion sites for the ASAs would end up with very different results. These differences can be explained by the bone density at the

insertion site. Cortical bone density seems to be the main determinant of the initial fixation strength of an ASA. Placing an ASA in an area with good cortical density such as the glenoid rim results in failure through suture breaking or soft-tissue failure, whereas placing an ASA in a weak cortical area such as the greater tuberosity of the humeral head may result in anchor pullout.

Although ASAs showed results comparable to those of controls on load-to-failure tests, results of cyclic loading tests should be evaluated with caution. Test protocols and displacement cut-off values in those studies are inconsistent (Tables 1 and 2). Lack of a standard test protocol seems to be an important drawback for comparing anchors. Authors used differing ranges of cyclic tests varying between 100 and 500 times.<sup>3,4,6,7,10,11,13,14</sup> As the number of cycles increases up to 5×, this can significantly affect the outcomes. Cut-off values for evaluating the amount of displacement is another area of debate. Some authors used 2 mm as a level for significant displacement during cyclic loading, whereas others used 5 mm. In a real-world scenario, a displacement more than 2 mm in a labral fixation would interfere with healing.

Few authors studied the insertion angle, the pre-tensioning or the drilling depth of ASA anchors. Oh et al. showed that ASAs demonstrated better mechanical properties when inserted at a more vertical angle (90 and 75 degrees) than 45 degrees.<sup>15</sup> The stability of an ASA depends mainly on the cortical bone, so insertion of an all-suture anchor using a vertical vector seems to be logical. Dwyer et al. found that 60 Newton pre-tensioning an ASA ensures that the anchor is well fixed, consistently eliminating laxity and displacement.<sup>4</sup> Because of the possibility of a handset ASA to become loose and displace prior to reaching high tension forces, pre-tensioning the ASA by a predetermined force seems to be useful. Ruder et al. reported less displacement by decreasing the preset drill depth of a specific ASA.<sup>9</sup> Cortical thickness at the insertion point of an ASA seems to be the main determinant of drilling depth. As the cortical thickness increases, drill depth should be increased to ensure a proper anchor deployment. An improper drill depth may cause 2 major problems. A thick cortex associated with a short drill depth may prevent adequate suture deployment. However, longer drill depths will position the anchor far from the cortex, which may leave a small amount of spongy bone between the anchor and the near cortex. This may cause a pistoning motion and interfere with the stability of the anchor. A zone-specific and/or personalized insertion depth for ASAs may help surgeons get the most out of ASAs.

Clinical findings of patients treated with ASAs for arthroscopic shoulder instability and rotator cuff repair

showed satisfactory results and little increase in the complication rate (retear, revision surgery because of loose anchor) (Table 4).<sup>18</sup> These clinical study results seem correlated with the biomechanical studies. Patients treated with ASAs for the purposes of either labral or rotator cuff repair were radiologically investigated in postoperative period and showed very little bone reaction or cyst formation. These findings counter the study of Pfeiffer et al.,<sup>6</sup> who showed consistent cavity formation and significant expansion of the drill tunnel in an in vivo study in the canine glenoid. However, there are not enough clinical data to define the risks for cyst formation using the ASAs.

Current data retrieved by this review lead us to the following observations:

- The weakest link in an all-suture anchor repair construct seems to be the cortical structure; therefore, the surgical decision should be based on the bone quality.
- Various ASAs have differing application techniques and preset drill depths. Surgeons should be familiar with these anchors to get the most out of them.
- Standardized mechanical protocols to test anchors are still lacking. Any direct comparison among various mechanical studies should be made with great caution.
- A perfect experimental model for bone is still missing. Biphasic polyurethane foam seems to be the best current option.

The principal limitation of the study is that a relatively small number of studies was identified, and they used widely diverse biomechanical measurement methodologies. Second, investigations were done in various soft tissue structures, such as ligamentous (labral) or tendinous (rotator cuff and biceps tendon) tissues.

## Conclusions

In conclusion, all suture anchors have similar or better mechanical properties than regular anchors. Low-profile design seems to be an important advantage that would preserve bone tissue during application. Radiologically investigated case series showed very little bone reaction or cyst formation in patients treated with ASAs. Clinical findings in these patients who underwent arthroscopic shoulder and acetabular labral repair and rotator cuff repair showed satisfactory results and had low complication rates. However, these case series cannot distinguish between the possible clinical benefits and/or risks of ASAs.

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