Supplementary file 3. GRADE

Table S1. Diagnos	stics										
Test	Test result	Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Number of participants	Diagnostic accuracy	Certainty
Subacromial imp	ingement						-				
Composit test (combination of Hawkins- Kennedy, Neer, Painful arc,	Positive Negative	1	Prospective cohort study	Not serious	Not assessed	Serious ^a	Serious ^b	None	55	LR+=2.93 LR-=0.34	Low Low
Empty can/Jobe, external rotation against resistance) ¹											
Internal posteros	uperior imp	pingement					-				
Posterior impingement test ²	Positive Negative	1	Prospective cohort study	N/I	Not assessed	Not assessed	Not assessed	Not assessed	69	LR+=5.0 LR-=0.29	N/A ^c
Anterior instabili	ity			•			•	•		•	
Apprehension ¹	Positive Negative	2	Retrospective cohort studies	Not serious	Not serious	Serious ^a	Serious ^b	Large effect ^d	409	LR+=17.21 LR-=0.39	Moderate Low
Relocation ¹	Positive Negative	3	Cohort studies	Not serious	Serious ^e	Serious ^a	Serious ^f	None	509	LR+=5.48 LR-=0.55	Very low Very low
Surprise ¹	Positive Negative	2	Cohort studies	Not serious	Serious ^e	Serious ^a	Serious ^f	None	128	LR+=5.42 LR-=0.25	Very low Very low
Apprehension + relocation ¹	Positive Negative	1	Prospective cohort study	Not serious	Not serious	Serious ^a	Serious ^b	Large effect ^d	46	LR+=39.68 LR-=0.19	Moderate Moderate
SLAP				•			•			•	
Biceps load II ^{3c}	Positive Negative	1	Prospective cohort study	Not serious	Not assessed	Serious ^a	Serious ^b	Large effect ^g	127	LR+=26.38/PPV=92.1 LR-=0.11/ NPV=95.5	Moderate Moderate
Biceps-Labrum c	omplex inju	ıries									
O'Brien's active compression; Inside ^{4 d}	Positive Negative	1	Prospective cohort study	Not serious	Not assessed	Serious ^a	Serious ^b	None	116	LR+=1.62/PPV=63.2 LR-=0.27/NPV=77.8	Low Low
O'Brien's active compression; Junctional ^{4 d}	Positive Negative	1	Prospective cohort study	Not serious	Not assessed	Serious ^a	Serious ^b	None	116	LR+=2.48/PPV=82.4 LR-=0.15/NPV=77.8	Low Low
O'Brien's active compression;	Positive Negative	1	Prospective cohort study	Not serious	Not assessed	Serious ^a	Serious ^b	Large effect ^h	116	LR+=2.00/PPV=65.7 LR-=0.08/NPV=92.6	Low Moderate

Bicipital tunnel ⁴											
1											
Throwing test; Inside ^{4 d}	Positive Negative	1	Prospective cohort study	Not serious	Not assessed	Serious ^a	Serious ^b	None	116	LR+=2.32/PPV=71.2 LR-=0.36/NPV=72.1	Low Low
Throwing test;	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	None	116	LR+=3.42/PPV=86.5	Moderate
Junctional ^{4 d}	Negative		cohort study	serious					-	LR-=0.35/NPV=60.5	Low
Throwing test;	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	None	116	LR+=2.09/PPV=66.7	Low
Bicipital tunnel ⁴	Negative		cohort study	serious						LR-=0.40/NPV=72.1	Low
Bicipital tunnel	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	None	116	LR+=1.92/PPV=67.2	Low
palpation; Inside ^{4 d}	Negative		cohort study	serious						LR-=0.16/NPV=85.7	Moderate
Bicipital tunnel	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	Large effecth	116	LR+=3.43/PPV=86.6	Moderate
palpation; Junctional ^{4 d}	Negative		cohort study	serious						LR-=0.09/NPV=85.7	Moderate
Bicipital tunnel	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	Large effect ^h	116	LR+=2.24/PPV=68.2	Low
palpation; Bicipital tunnel ⁴	Negative		cohort study	serious						LR-=0.04/NPV=96.4	Moderate
Yergasons test;	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	None	116	LR+=2.13	Low
Inside ^{4 d}	Negative		cohort study	serious						LR-=0.76	Low
Yergasons test;	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	None	116	LR+=6.57	Low
Junctional ^{4 d}	Negative	1	cohort study	serious		a :	G i h	N	116	LR-=0.83	Low
Yergasons test; Bicipital tunnel ⁴	Positive Negative	1	Prospective cohort study	Not serious	Not assessed	Serious ^a	Serious ^b	None	116	LR+=12.43 LR-=0.75	Moderate Low
Rotator cuff inju						1					
Painful Arc ⁵	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	None	104	LR+=3.70	Low
<u>a 1 7 % 66</u>	Negative		cohort study	serious	a i i	a · · ·	a i h			LR-=0.36	Low
Gerber/Lift-off test ⁵	Positive Negative	2	Prospective cohort studies	Not serious	Serious ⁱ	Serious ^a	Serious ^b	None	233	LR+=1.40-1.50 LR-=0.63-0.85	Low Low
External rotation	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	None	203	LR-=0.03-0.83 LR+=2.60	Low
against	Negative	1	cohort study	serious	Not assessed	Serious	Serious	None	203	LR-=0.49	Low
resistance ⁵	_		_								
Full can ⁵	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	None	104	LR+=2.40	Low
	Negative		cohort study	serious						LR-=0.37	Low
Empty can/Jobe ⁵	Positive	3	Prospective	Not	Serious ^e	Serious ^a	Serious ^f	None	337	LR+=1.30	Very low
F II	Negative		cohort studies	serious						LR-=0.64	Very low
Full rotator cuff		1		NT (G · · · ·	0 · h	N	27	1.0. 7.20	T
External rotation	Positive Negative		Prospective	Not	Not assessed	Serious ^a	Serious ^b	None	37	LR+=7.20 LR-=0.57	Low Low
lag ⁵	negative	1	cohort study	serious	1					LK-=0.3/	LOW

Internal rotation	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b	Large effect ^h	37	LR+=5.60	Low
lag ⁵	Negative		cohort study	serious						LR-=0.04	Moderate
Drop sign ⁵	Positive	1	Prospective	Not	Not assessed	Serious ^a	Serious ^b		37	LR+=3.20	Low
	Negative		cohort study	serious						LR-=0.35	Low
Explanations:											
^a Downgraded one	level due to	the popula	tion being non-a	thletes							
^b Downgraded one	e level due to	a limited i	number of includ	ed studies							
^c The article by M	eister et al. v	vas not obt	ained in full text	hindering r	isk of bias assess	sment					
^d Positive test resu	lt upgraded	one level d	ue to high diagno	stic accura	cy						
e Downgraded one	level due to	significan	t heterogeneity in	the pooled	lestimate						
f Downgraded one	level due to	wide 95%	confidence inter	vals in poo	led estimates						
g Positive and neg	ative test res	ults were u	pgraded one leve	l due to hig	gh diagnostic acc	curacy					
h Negative test res	ult was upgr	aded one le	evel due to high d	iagnostic a	ccuracy						
in 11	1 1 1 7	1 .	1		1.						

ⁱ Downgraded one level due to large variability in point estimates between studies

Table S2. Prevention	1				1		1	1	1	1
Outcome/intervention	Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Number of participants	Effect (95% CI)	Certainty
Risk of shoulder prob	lems (all sl	noulder problems)								
Oslo Sports Trauma Research Center Shoulder Injury Prevention program vs. usual care ⁶	1	Randomized controlled trial	Not serious	Not assessed	Not serious	Serious ^a	None	660	OR= 0.72 [0.52 to 0.98]	Moderate
Risk of shoulder prob	lems (subs	tantial shoulder nr	oblems)						I	
Oslo Sports Trauma Research Center Shoulder Injury Prevention program vs. usual care ⁶	1	Randomized controlled trial	Not serious	Not assessed	Not serious	Serious ^a	None	660	OR= 0.78 [0.53 to 1.16]	Moderate
Risk of shoulder injur	·y		•		•	•	•		·	
Shoulder Control program vs. usual care ⁷	1	Randomized controlled trial	Not seripus	Not assessed	Not serious	Serious ^a	None	464	HRR= 0.44 [0.29 to 0.68]	Moderate
Throwing injury prevention program vs. usual care ⁸	1	Randomized controlled trial	Not serious	Not assessed	Not serious	Serious ^a	None	237	HR= 0.48 [0.21 to 1.08]	Moderate
FIFA 11+ shoulder prevention program vs. usual care ⁹	1	Randomized controlled trial	Not serious	Not assessed	Not serious	Serious ^a	None	726	IRR= 0.28 [0.13 to 0.60]	Moderate
Intervention including Sleeper's stretch vs. usual care ¹⁰	1	Prospective cohort study	Serious ^b	Not assessed	Not serious	Serious ^a	None	46	HR=0.35 [0.13 to 0.94]	Very low
Intervention including Sleeper's stretch and prone shoulder external rotation exercise vs. usual care ¹⁰	1	Prospective cohort study	Serious ^b	Not assessed	Not serious	Serious ^a	None	60	HR=0.47 [0.20 to 1.10]	Very low
Pas et al. ¹¹	1	Randomized controlled trial	Serious ^a	Not assessed	Not serious	Serious ^a	None	579	OR=0.96 (p=0.93)	Low
Achenbach et al. ¹²	1	Randomized controlled trial	Serious ^a	Not assessed	Not serious	Serious ^a	None	579	Absolute risk reduction= - 2.5% [-10.3 to 5.4]	Low

Intervention including functional exercises using resistance bands or dumbbells vs. usual care ¹³	1	Randomized controlled trial	Serious ^c	Not assessed	Not serious	Serious ^a	None	26	Between-group difference in number of shoulder pain episodes= 2.8 [CI not reported], p=0.02	Low
Patient-reported shou	ılder pain									
Intervention including shoulder strengthening exercises vs. usual care ¹⁴	1	Randomized controlled trial	Not serious	Not assessed	Not serious	Serious ^a	None	206	Between group difference in VAS= 0.1 [CI not reported], p=0.746	Moderate
Explanations: ^a Downgraded one leve ^b Downgraded one leve ^c Downgraded one leve OR, odds ratio; HR, ha	el due to cri el due to hig	tical risk of bias in R gh risk of bias in ROI	OBINS-I 3-2	analog scale						

Table S3. Treatment										
Intervention/outcome	Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Number of participants	Effect (95% CI or SD)	Certainty
Subacromial impinge	ment					-		-		
Intervention including shoulder specific warm-up and exercises vs. no intervention on pain (NRS) ¹⁵	1	Randomized controlled trial	Not serious	Not assessed	None	Serious ^a	None	30	Within-in group baseline and follow-up scores [SD] Intervention 7.88 [1.02] to 3.56 [1.31] Control 7.71 [0.83] to 8.00 [0.88]	Moderate
Intervention including strengthening exercises (no comparator group) on pain and function (SPADII) ¹⁶	1	Prospective cohort study	Not serious	Not assessed	None	Serious ^a	None	47	Within-in group baseline and follow-up scores [SD] 29.86 [17.03] to 11.7 [13.78]	Very low
Supraspinatus tendin	opathy		-	•			1			
Interventions including hyperthermia or ultrasound vs. passive stretches on pain (VAS) ¹⁷	1	Randomized controlled trial	Not serious	Not assessed	None	Serious ^a	None	37	Within-in group baseline and follow-up scores [SD] Hyperthermia 5.96 [0.83] to 1.2 [0.63] Ultrasound 6.3 [0.86] to 5.15 [0.87] Passive stretches 6.1 [0.89] to 4.9 [0.88]	Moderate
Interventions including hyperthermia or ultrasound vs. passive stretches on physical function (Constant Murley Score) ¹⁷	1	Randomized controlled trial	Not serious	Not assessed	None	Serious ^a	None	37	Within-in group baseline and follow-up scores [SD] Hyperthermia 58.6 [3.9] to 82.0 [5.7] Ultrasound 58.9 [2.8] to 61.8 [4.2]	Moderate

									Passive stretches 59.5 [2.7] to 63.3 [5.6]	
Shoulder pain									59.5 [2.7] to 05.5 [5.0]	
Intervention including anteroposterior mobilisation of the shoulder joint vs. manual treatment vs. attention on pain (VAS) ¹⁸	1	Randomized controlled trial	Not serious	Not assessed	None	Serious ^a	None	31	Within-group changes [95% CI] Mobilization 0.6 [0.1 to 1.1] Manual treatment 0.0 [0.0 to 0.3] Attention	Moderate
Intervention including anteroposterior mobilisation of the shoulder joint vs. manual treatment vs. attention on physical function (DASH) ¹⁸	1	Randomized controlled trial	Not serious	Not assessed	None	Serious ^a	None	31	0.2 [-0.2 to 0.7] Within-group changes [95% CI] Mobilization 0.3 [-2.7 to 3.4] Manual treatment 0.5 [-0.3 to 1.3] Attention 0.7 [-0.6 to 2.0]	Moderate
Intervention including posture correcting exercises vs. no intervention on physical function and pain (ASES) ¹⁹	1	Randomized controlled trial	Serious ^b	Not assessed	None	Serious ^a	None	28	Within-in group baseline and follow-up scores [SD] Intervention Right shoulder: 89.1 [11.2] to 89.3 [14.6] Left shoulder: 89.9 [11.4] to 91.1 [10.6] Control Right shoulder: 90.8 [11.7] to 86.4 [17.9] Left shoulder: 90.7 [12.4] to 86.9 [15.5]	Low
Intervention includes strengthening	1	Prospective cohort study	Serious ^c	Not assessed	None	Serious ^a	None	29	Within-in group baseline and follow-up scores [SD]	Very low

exercises (no comparator group) on pain (VAS) ²⁰									3-months follow-up: 7.5 [2.3] to 3.4 [1.8] 6 months follow-up: 7.5 [2.3] to 2.9 [2.1]	
Intervention including scapula- focused stretching and strengthening exercises (no comparator group) on pain (VAS) ²¹	1	Prospective cohort study	Serious ^c	Not assessed	None	Serious ^a	None	31	7.5 [2.5] to 2.9 [2.1] Within-in group baseline and follow-up scores [SD] 3-months follow-up: 7.2 [1.3] to 2.4 [1.8] 6 months follow-up: 7.2 [1.3] to 2.6 [1.4]	Very low
Explanations: ^a Downgraded one leve ^b Downgraded one leve ^c Downgraded one leve	l due to hig	gh risk of bias in ROI	B-2							

NRS, numeric rating scale; SPADI, Shoulder Pain and Disability Index; DASH, Disabilities of the Arm, Shoulder, and Hand; ASES, The American Shoulder and Elbow Surgeons Shoulder Score.

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