# Supplementary Appendix

Supplement to:

# Baseline clinical and MRI risk factors for hamstring reinjury showing the value of performing baseline MRI and delaying return to play: a multicentre, prospective cohort of 330 acute hamstring injuries

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### **Table of contents**

Content	Page
Full authors affiliations	3
Hamstring Study Group (Qatari and Dutch)	5
Supplement Appendix 1: Method of included study	10
Supplement Appendix 2: Rehabilitation program of included study	11
Supplement Appendix 3: Data cleaning protocol	15
Supplement Appendix 4: Descriptive baseline statistics for total study population and separated into those who did or did not incur a reinjury within 2- and 12 months return to play.	17

References

22

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## Qatari and Dutch Hamstring Study Group

## GF Growth Factor study

- HIT Hamstring Injection Therapy Study
- HAR Rehabilitation of Acute Hamstring Injury Study
- HIR Diffusion Tensor Imaging (DTI) for hamstring injury Study

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## Supplement Appendix 1: Summary of methods of the included study.

	Growth Factor study <sup>1</sup>	Rehabilitation of Acute Hamstring Injury study <sup>2</sup>	Hamstring Injection Therapy study <sup>3</sup>	DTI for Hamstring Injury study (ongoing)
Study design	Randomized controlled trial	Randomized controlled trial	Randomized controlled trial	Prospective cohort
Country	Qatar (Aspetar), Doha	Qatar (Aspetar), Doha	Netherlands (Multicentre)	Netherlands (AUMC), Amsterdam
Intervention(s)	<ol> <li>Platelet Rich Plasma (PRP) + Rehab program.</li> <li>Platelet Poor Plasma (PPP) + Rehab program.</li> <li>No injection, only rehab. Program.</li> </ol>	<ol> <li>Rehab. Program + early introduction to lengthening exercises (day one of rehabilitation).</li> <li>Rehab. Program + late introduction to lengthening exercises (&gt;70% of self-rated max. speed).</li> </ol>	<ol> <li>Two Platelet-rich plasma (PRP) injections + rehab program.</li> <li>Two Isotonic saline injections (placebo) + rehab. Program.</li> </ol>	Voluntary physical therapy program
Study objectives	Efficacy of PRP in enhancing RTP following hamstring injury	Effect of early introduction of lengthening exercises on time to return to play and reinjury risk.	Efficacy of PRP in enhancing RTP following hamstring injury	Correlation between Diffusion Tensor imaging parameters return to play and reinjury.
Patient recruitment Eligibility criteria	<ul> <li>Recruitment through Qatar National Sports Medicine Program (NSMP) Inclusion: <ul> <li>Age: 18-50</li> <li>Gender: Male</li> <li>Available for follow-up</li> <li>Acute onset of posterior thigh pain</li> <li>Presenting an MRI ≤ 5 days from injury</li> <li>MRI confirmed grade I or II hamstring lesion</li> <li>Able to perform five sessions of physiotherapy a week at the study location Exclusion: <ul> <li>Contra-indication MRI</li> <li>Reinjury or chronic hamstring injury</li> <li>Concurrent other injuries inhibiting rehabilitation</li> <li>Unwilling to comply with follow-up</li> <li>Needle phobia</li> <li>Overlying skin infection</li> <li>Diabetes/immunocompromised state</li> <li>Medication with increased bleeding risk</li> <li>Medical contraindication to injection</li> </ul> </li> </ul></li></ul>	<ul> <li>Recruitment through Qatar National Sports Medicine Program (NSMP)</li> <li>Inclusion: <ul> <li>Age: 18-50</li> <li>Gender: Male</li> <li>Available for follow-up</li> <li>Acute onset of posterior thight pain when training or competing.</li> <li>Clinical diagnosis of an acute hamstring muscle injury is defined as (a) localised pain during palpation of hamstring muscle, (b) increased pain during isometric contraction (c) Localised pain when performing a passive straight leg test.</li> <li>Presenting an MRI ≤ 5 days from injury</li> <li>MRI confirmed grade I or II hamstring lesion</li> <li>Able to perform ≥ 3 sessions of physiotherapy a week at study location</li> </ul> </li> <li>Exclusion: <ul> <li>Verified or suspected previous hamstring injury in the same leg ≤ 6 months.</li> <li>Chronic hamstring complaints ≥2 months</li> <li>Grade III injury, including complete hamstring disruption of tendon avulsion (modified Peetrons)</li> <li>Contraindication to MRI</li> <li>No intention to return to full sport activity</li> <li>Refusal to receive one of the two therapies.</li> </ul> </li> </ul>	<ul> <li>Nationwide referral program for suspected hamstring injury</li> <li>Inclusion: <ul> <li>Age 18-50</li> <li>Clinical diagnosis of an acute hamstring injury: (a)</li> <li>History of acute onset of posterior thigh pain, (b) localized pain on palpation, (c) localized pain on palsive stretch of the hamstring, and (d) increased pain on isometric contraction</li> <li>Hamstring lesion on MRI (increased signal intensity on STIR and/or T2-weighted imaging)</li> </ul> </li> <li>Exclusion: <ul> <li>Not capable of doing an active exercise program</li> <li>Previous injection therapy for this injury</li> <li>No intention to return to full sports activity</li> <li>Contraindication to MRI</li> <li>Refusal to receive one of the two therapies.</li> <li>Injury caused by extrinsic trauma</li> <li>Chronic low back pain</li> <li>Chronic hamstring complaints ≥2 months</li> <li>Grade III lesion (total rupture) or avulsion on MRI.</li> </ul> </li> </ul>	<ul> <li>Online recruitment or referral by physician /physical therapist.</li> <li>Inclusion: <ul> <li>Age &gt;16</li> <li>Clinical diagnosis of an acute hamstring injury ≤7 days defined as:</li> <li>Anamnestic acute injury: Acute injury, localized pain posterior thigh, pain during palpation of hamstring muscle, -passive straight leg raise, - isometric contraction.</li> </ul> </li> <li>Exclusion: <ul> <li>Not capable of doing an active activity program</li> <li>No intention to return to full sports activity</li> <li>Injury caused by extrinsic trauma</li> <li>Contraindication to MRI</li> <li>Reinjury ≤2 months</li> <li>Complete proximal tendon avulsion (grade 3)</li> <li>Other concurrent injury inhibiting rehabilitation</li> </ul> </li> </ul>
Randomization & blinding	Three-block randomization. Blinded assessor Patient blinding for injection arm.	Computer-generated, random allocation sequence.	Computer-generated permuted block scheme. All were blinded except the coordinating physician (preparation of injection).	

## Supplement Appendix 2: Rehabilitation program of included study.

1. Hamstring Injection Therapy (Study Dutch Trial Register 2771)<sup>3</sup>

Physiotherapist	t supervised program (modified Heiderscheit,2010) <sup>4</sup>	
Phase	Content	Criteria to progress
Phase 1	<b>Goals:</b> Protection of scar formation, minimizing muscle atrophy and stimulating neuromuscular control.	Criteria for progression to the next phase
	<ul> <li>Protection. Exercises limited to pain-free range of motion (ROM). No excessive lengthening or resistance training of the hamstrings</li> <li>Ice application. For pain reduction, ice can be applied 2-3 times a day (maximal 3-5 minutes when using ice and maximal 15-20 minutes when using a cool pack).</li> </ul>	<ol> <li>Normal walking stride without pain</li> <li>very low-speed jog without pain</li> <li>Pain-free isometric contraction against sub-maximal (50- 70%) resistance during prone knee flexion manual strength test in 90° knee flexion.</li> </ol>
	<b>Exercises</b> . Ergometer cycling, low to moderate intensity stepping exercises (such as side, grapevine, and in place fast feet stepping), isometric exercises for lumbopelvic musculature, single limb balance exercises, and core-stability exercises (such as prone body bridge, side body bridge, and supine bent knee bridge). Exercises should be performed without pain.	
Phase 2	<b>Goals:</b> Regaining pain-free ROM and development of trunk and pelvis neuromuscular control with a progressive increase in movement speed.	Criteria for progression to the next phase
	<b>Protection.</b> No end-range lengthening of the hamstrings when muscle weakness is still present.	<ol> <li>Pain-free full strength (5/5) during prone knee flexion manual strength test in 90° knee flexion</li> <li>Pain-free forward and backward</li> </ol>
	<b>Ice application.</b> For pain reduction, ice can be applied after exercises (maximal 3-5 minutes when using ice and maximal 15-20 minutes when using a cool pack).	jogging at 50% of maximum speed.
	<b>Exercises.</b> Gradual increase in hamstring lengthening and intensity of exercises. Agility drills and core-stability exercises are performed with a progressive increase in speed and intensity. Based on the patient's tolerance, exercises are gradually increased in hamstring lengthening. Submaximal eccentric exercises are performed near mid-hamstring length. Start with anaerobic training and sport-specific skills, but take care to avoid end-range lengthening of the hamstrings or substantial eccentric work. Running should not be performed at a speed greater than 50% of the maximal speed.	
Phase 3	<b>Goals</b> : Symptom-free during all activities, normal concentric and eccentric hamstring strength through full ROM and full speeds, improvement neuromuscular control of trunk and pelvis, and improvement control in sport-specific movements.	Criteria for clearance to return to play Symptom-free (e.g., pain and stiffness) during: 1. full ROM;
	<b>Protection</b> . ROM is unrestricted. Sprinting and explosive acceleration should be avoided until full ROM and functional movement patterns (such as running, jumping, and cutting) can be performed pain-free.	<ol> <li>full-speed sprinting;</li> <li>sport-specific movements (such as jumping and cutting).</li> </ol>
	<b>Ice application</b> . For pain reduction, ice can be applied after exercises (maximal 3-5 minutes when using ice and maximal 15-20 minutes when using a cool pack).	
	<b>Exercises</b> . More challenging core-stability exercises by incorporating asymmetrical postures and motion exercises.	

	Eccentric exercises toward the end range of motion and	
	increasing resistance (e.g., lunche walk with trunk rotation,	
	supine single limb chair-bridge). Agility and sport-specific drills	
	involving quick direction changes and technique training.	
Home exercise	program (Progressive agility and trunk stabilisation from Sherry	and Best) <sup>5</sup>
Phase 1	Low- to moderate-intensity sidestepping, 3 × 1 minute;	Criteria for progression to the next
	<ul> <li>Low- to moderate-intensity grapevine stepping (lateral stepping with the trail leg going over the lead leg and</li> </ul>	phase:
	then under the lead leg), both directions, $3 \times 1$ minute;	1. Able to walk pain-free with normal
	<ul> <li>Low- to moderate-intensity steps forward and backward</li> </ul>	gait pattern (e.g., same stride
	over a tape line while moving sideways, $2 \times 1$ minute;	length and stance time on the
	<ul> <li>Single-leg stand progressing from eyes open to eyes</li> </ul>	injured leg and stance leg;
	closed, 4 × 20 seconds;	2. Able to do a pain-free high knee
	Prone abdominal body bridge (performed by using	march.
	abdominal and hip muscles to hold the body in a face-	
	down straight-plank position with the elbows and feet as	
	the only point of contact), $4 \times 20$ seconds;	
	Supine extension bridge (performed by using abdominal	
	and hip muscles to hold the body in a supine hook lying	
	position with the head, upper back, arms, and feet as the	
	points of contact), 4 × 20 seconds;	
	<ul> <li>Side bridge, 4 × 20 seconds on each side;</li> <li>Iso while sitting for 20 minutes.</li> </ul>	
Phase 2	<ul> <li>Ice while sitting for 20 minutes.</li> <li>Moderate- to high-intensity sidestepping, 3 × 1 minute;</li> </ul>	
Pliase Z	<ul> <li>Moderate- to high-intensity sidestepping, 5 × 1 minute,</li> <li>Moderate- to high-intensity grapevine stepping, 3 × 1</li> </ul>	
	minute;	
	<ul> <li>Moderate- to high-intensity steps forward and backward</li> </ul>	
	while moving sideways, 2 × 1 minute;	
	Single-leg stand windmill touches, 4 × 20 seconds of	
	repetitive alternate hand touches;	
	Push-up stabilization with trunk rotation (performed by	
	starting at the top of a full push-up, then maintaining this	
	position with one hand while rotating the chest toward	
	the side of the hand that is being lifted to point toward	
	the ceiling, pause and return to the starting position), $2 \times 10^{-1}$	
	15 repetitions on each side;	
	<ul> <li>Fast feet on the spot (performed by jogging in place with increasing uploating the fact only a family inches off</li> </ul>	
	increasing velocity, picking the foot only a few inches off	
	the ground), 4 × 20 seconds; Proprioceptive neuromuscular facilitation trunk pull-	
	downs with Thera-Band, 2 × 15 to the right and left;	
	<ul> <li>Symptom-free practice without high-speed maneuvers;</li> </ul>	
	<ul> <li>Ice for 20 min if any symptoms of local fatigue or</li> </ul>	
	discomfort are present.	
Notification	- The intensity of each exercise should be such that the	patient can perform the exercise
	pain-free;	- -
	<ul> <li>Low intensity: a velocity of movement that is less than</li> </ul>	n or near that of normal walking;
	<ul> <li>Moderate intensity, a velocity of movement greater the</li> </ul>	han normal walking but not as
	great assport;	
	<ul> <li>High intensity, a velocity of movement similar to sport</li> </ul>	t activity.

# 2. Growth Factor study (Clinicaltrials.gov NCT 01812564)<sup>1</sup>

Stage	Content	Criteria to progress
Stage 1	<ul> <li>All activities to be pain-free</li> <li>Two-leg squat, or if able, single-leg squat</li> <li>Maintain pelvis control, hip and knee alignment, squat to 45°, hold, return to start</li> <li>Supine Bridge—2 leg</li> <li>2 s up, 2 s down (4 s total per rep.) Begin at 45°. Must reach knee-hip-shoulder in alignment. 4×15</li> <li>supine isometric heel digs</li> <li>In supine, painlessly pull heel into bed through range. Can bias with tibial IR/ER when painless.</li> <li>Exercise bike</li> <li>Upright or recumbent can be substituted with the elliptical trainer</li> <li>Isometric manual-resisted hamstring</li> <li>The therapist applied resistance isometrically in varying angles in prone</li> <li>Soft tissue massage</li> <li>Proximal and distal to the injury site, lymphatic drainage.</li> <li>Active range of motion exercises</li> <li>Supine active knee flexion and extension, then Prone active</li> </ul>	Criteria to progress to stage 2: 1. Painless single-leg squat 2. Painless bike, 150W, 5 min 3. Full knee extension supine
Stage 2	flexion and extension Any exercise from stage 1 permitted, additionally:	Criteria to progress to stage 3: 1. Run ≥70% patient rated
	<ul> <li>Supine bridge—1 leg</li> <li>Same rate as for two legs, another knee in full extension, and thighs parallel throughout the exercise. 4×15</li> <li>Walk-Jog</li> <li>Walk 20 m corners, jog the 30 m straight, painless. Begin at 25% (self-rated) jog, progress to max70%.</li> <li>Triple extension walks</li> <li>100 m laps, every third step triple extension—i.e., alternating legs.</li> <li>'A' drill</li> <li>Walking late swing knee extension, painless. Alternating legs, 100 m lap.</li> <li>Soft tissue massage</li> <li>Can massage the injured area. Maximum allowed pain VAS: 4/10. The therapist uses caution with any report of discomfort, monitors symptoms, and adjusts accordingly.</li> <li>Stretching</li> <li>Hamstring (supine, 90° hip flexion, knee extension); SLR (supine to the onset of discomfort, add ankle DF)</li> <li>Initially active, patient-controlled, progress to passive, end range. SLR mobilisation if indicated.</li> <li>Resisted hamstring</li> <li>Note tibial rotation as indicated. 4×15 repetitions, aiming for fatigue</li> </ul>	<ol> <li>ROM hamstrings ≥75% uninvolved side</li> <li>ROM SLR ≥75% uninvolved side</li> </ol>
Stage 3	<ul> <li>Any exercises from stages 1 and 2, additionally:</li> <li>Single leg bridge</li> <li>1s repetition, 2s recovery. 4×8 repetitions.</li> <li>Single leg bridge, foot on the Swiss ball</li> <li>1s up, 2s down. 4×8 repetitions.</li> </ul>	Criteria to progress to stage 4 (sport-specific rehab): 1. 100% running speed 2. Painless high-speed direction changes

	<ul> <li>Interval running</li> <li>20 m jog, 30 m run. Begin running at 70% (patient-rated), progressing by 10% steps, painlessly. At 90%, progress by 5%. Monitor performance by hand timing.</li> <li>Modified T-Drill</li> <li>Direction changing running over T-Drill course. Begin at patient rated 70%, progress as able by 10% until 90%, then by 5%. Monitor performance by hand timing.</li> <li>Eccentric exercises</li> <li>Nordic Hamstrings, manual-resisted eccentric, prone catches, Arabesque (single leg stance, trunk flexion)</li> </ul>	
Stage 4	<ul> <li>Any exercises from stages 1–3, additionally on-field, football-specific drills:</li> <li>Direction change drills</li> <li>With and without the ball, 40 min</li> <li>Jumping drills</li> <li>10–15 min</li> </ul>	Criteria to progress to stage 5 (sport-specific rehab): 1. Painless completion of stage 4
Stage 5	Passes and run Long passes progression Crosses (static) Corner kicks Crosses (dynamic)	Criteria to progress to stage 6 (sport-specific rehab): 1. Painless completion of stage 5
Stage 6	Passes and run Shooting scenarios Competitive one versus-one drill Shooting scenarios Scoring scenarios	Criteria to progress to medical review for return to sport: 1. Painless completion of stage 6

DF, dorsiflexion; ER, external rotation; IR, internal rotation; Modified T-Drill, (always) forward running over the course of the Agility t test; ROM, range of motion; SLR, straight leg raise.

#### 3. Rehabilitation of Acute Hamstring Injury study (Clinicaltrials.gov NCT 02104258)<sup>2</sup>

Both groups received a similar 6-stage standard criteria-based rehabilitation program (see rehabilitation program for Growth Factor Study). The difference between the treatment groups was the introduction of the lengthening exercise (the extender, the diver, and the slider exercise)<sup>6,7</sup> at different time points. In the early lengthening group, the lengthening exercise was introduced on day 1 of rehabilitation. In the delayed lengthening group, the lengthening exercises were introduced after meeting the criteria of rehabilitation program stage 3 (able to run more than 70% of maximal speed).

4. DTI for Hamstring Injury study (Dutch Trial Register 2016\_033).

The patient was advised to be treated using a criteria-based rehabilitation program but on a voluntary basis. The optional rehabilitation protocol, such as Aspetar,<sup>8,9</sup> Sherry and Best,<sup>5</sup> Mendighucia<sup>10</sup> and Askling<sup>7</sup>). The rehabilitation protocols were available for patients and therapists on the study website.

Supplement Appendix 3: Data cleaning protocol.

## **Data cleaning Merged databases (Muscle Medics)**

Based on the Quality Manual of the Department of General Practice of the Erasmus MC)

#### Data to be cleaned

All variables merged from different databases:

- Qatar: Growth Factor Study (Clinical Trials.gov NCT 01812564)
- Qatar: Hamstring Acute Rehabilitation Study (Clinical Trials.gov NCT02104258)
- The Netherlands: Hamstring Injection Therapy Study (Dutch Trial Register 2771)
- The Netherlands: Diffuse Tensor Imaging (DTI) for Hamstring Injury Study (Dutch Trial Register 2016\_033)

#### **Data Cleaning Protocol**

- I. All new variables were derived/recoded from merged dataset (e.g., variables differentiating for injured and uninjured leg derived from the injured side and variables for right and left leg).
- II. Manually correcting all odd data points as found during the verifying process.
- III. Manually check the listed recoding formulas for coding errors.
- IV. Manually checking all measurements of a selection of 3% of all participants per database in a statistical software package to check for consistency with original measurements after performing pre-listed recoding formulas. When the percentage of disagreement exceeds 1.5% within a measurement of one-time point, the random selection is increased to 15% of all participants for that measurement. If the percentage of disagreement exceeds 1.5% within the measurements of one-time point in the extended selection of participants, all measurements from that time point will be digitally rescanned and data will be reprocessed. All indicated faults will be corrected in the original data sets.
- V. Checking data of each variable is within possible ranges and manually correcting data points based on the original study forms.
- VI. Checking data on logic inconsistencies, i.e., checking dates of time points of date of injury, first visit, time of MRI, RTP, and reinjury for chronologic consistency.

## Data cleaning log

- Randomly "baseline variables" data of \*\*\*
  - $\circ$   $\,$  3 out of 90 (3%) subjects of GF Study (subject GF18, GF32, GF66)  $\,$
  - $\circ$  3 out of 88 (3%) subjects of HAR study) (subject HAR 20, HAR49, HAR54)
  - 2 out of 80 (3%) subjects of HIT study (subjects HIT 125 and HIT 145)
  - o 4 out of 120 (3%) HIR study (subjects HIR 41, HIR 45, HIR 49, HIR 146)

are checked and compared to the source data (original database) by a colleague researcher, not involved in the process of data cleaning for the reviewed selection of data (Mokkenstorm & Zein).

Two data points were not consistent with the source data after scored twice of a total number of 4476 data points checked. This gives an error margin of 0.04%

Error margin: 2 / 4476 = 0,04%

#### Changes in raw data file after performing data cleaning protocol:

Subject	Variable	False outcome	Source data
GF18	All data is consistent with the original data files.		
GF32	All data is consistent with the original data files.		
GF66	All data is consistent with the original data files.		
HAR 20	All data is consistent with the original data files.		
HAR49	BaslinePainPalpationYN [Column GD]	2	1
HAR54	All data is consistent with the original data files.		
HIT 125	BaselineReceivedTreatment_Medication_Yes_No (col BX)	0	2
HIT 145	All data is consistent with the original data files.		
HIR 41	All data is consistent with the original data files.		
HIR 45	All data is consistent with the original data files.		
HIR 49	All data is consistent with the original data files.		
HIR 146	All data is consistent with the original data files.		

#### Supplement Appendix 4: Descriptive baseline statistics for total study population and separated into those who did or did not incur a reinjury within 2-

and 12 months return to play.

Demographics & History	Total population (n=	2 months reinjury (n= 330)		12 months reinjury (n= 308)	
	368)	No reinjury	Reinjury	No reinjury	Reinjury
		(n= 299; 91%)	(n= 31; 9%)	(n= 256; 83%)	(n= 52; 17%)
Age (year)*	26 (21-30)	26 (21-30)	28 (23-30)	26 (21-30)	26 (22-30)
Sex male; n (%)	357 (97.0%)	290 (97.0%)	29 (93.5%)	250 (97.7%)	49 (94.2%)
Side of hamstring injury; n (%)					
- Right	199 (54.0%)	156 (52.2%)	19 (61.3%)	134 (52.3%)	29 (55.8%)
- Left	169 (46.0%)	143 (47.8%)	12 (38.7%)	122 (47.7%)	23 (44.2%)
Days since injury (day)*	3 (2-4)	3 (2-4)	3 (2-5)	3 (2-4)	3 (2-4)
Height (meter)*	1.790 (1.740-1.840)	1.790 (1.740-1.840)	1.800 (1.710-1.860)	1.790 (1.740-1.840)	1.805 (1.725-1.852)
Weight (kg)*	75.500 (70-80)	75.700 (70.000 –	76.000 (71.200-82.050)	75.000 (70.000-83.250)	75.500 (70.000-82.025)
		84.000)			
Body Mass Index (kg/m <sup>2</sup> )*	23.525 (22.300-25.200)	23.500 (22.340-25.200)	23.800 (22.600-24.700)	23.500 (22.300-25.240)	23.720 (22.575-24.700)
Type of sports; n (%)					
- Football	255 (61%)	208 (70%)	25 (81%)	178 (70%)	42 (80.8%)
- Field hockey	26 (7%)	22 (7%)	1 (3%)	20 (8%)	3 (6%)
- Athletics	17 (5%)	13 (4%)	1 (3%)	11 (4%)	2 (4%)
- Futsal	14 (4%)	13 (4%)	-	11 (4%)	1 (2%)
- Handball	11 (3%)	7 (2%)	1 (3%)	6 (2%)	1 (2%)
- Basketball	10 (3%)	9 (3%)	1 (3%)	6 (2%)	1 (2%)
- Other / Unknown	31 (8%)	27 (9%)	2 (7%)	24 (10%)	2 (4%)
Level of Sports; n (%)					
- Professional	241 (65%)	194 (65%)	19 (61%)	168 (66%)	29 (56%)

17

- Absent	25 (6.8%)	45 (15.1%) 24 (8.0%)	-	23 (9%)	8 (15.4%) 1 (1.9%)
- Present	287 (78.0%) 56 (15.2%)	230 (76.9%) 45 (15.1%)	25 (80.6%) 6 (19.4%)	194 (75.8%) 39 (15.2%)	43 (82.7%) 8 (15.4%)
Discomfort restricted flexion 90 degrees; n (%)					
ength of painful area (cm)*	8.000 (5.000-12.000)	8.000(5.000-12.000)	8.000 (3.000-12.500)	8.000 (5.000-13.000)	8.000 (4.500-11.500)
		(n=299; 91%)	(n=31; 9%)	(n= 256; 83%)	(n= 52; 17%)
	(n= 368)	No reinjury	Reinjury	No reinjury	Reinjury
Physical examination	Total population	2 months reinjury (n=	330)	12 months reinjury (n= 308)	
- Unknown/missing	2 (0.5%)	2 (0.7%)	-	2 (0.8%)	-
- Non-sprinting	265 (72.0%)	215 (71.9%)	21 (67.7%)	187 (73%)	32 (61.5%)
- Sprinting	101 (27.4%)	82 (27.4%)	10 (32.3%)	67 (26.2%)	20 (38.5%)
njury Mechanism: Sprinting / non-sprinting; n (%)					
- Unknown/missing	10 (2.7%)	7 (2.3%)	1 (3.2%)	7 (2.7%)	1 (1.9%)
- Training	112 (30.4%)	94 (31.4%)	7 (22.6%)	80 (31.3%)	13 (25%)
- Match/Competition	246 (66.8%)	198 (66.2%)	23 (74.2%)	169 (66%)	38 (73.1%)
Aoment of injury; n (%)					
- Unknown/missing	38 (10.3%)	28 (9.4%)	5 (16.1%)	26 (10.2%)	6 (11.5%)
- No	198 (53.8%)	164 (54.8%)	12 (38.7%)	142 (55.5%)	20 (38.5%)
- Yes	132 (35.9%)	107 (35.8%)	14 (45.2%)	88 (34.4%)	26 (50%)
History of hamstring injury on ipsilateral leg; n (%)					
- Unknown/missing	38 (10.3%)	28 (9.4%)	5 (16.1%)	26 (10.2%)	6 (11.5%)
- No	149 (40.5%)	125 (41.8%)	9 (29.0%)	110 (43.0%)	14 (26.9%)
- Yes	181 (49.2%)	146 (48.8%)	17 (54.8%)	120 (46.9%)	32 (61.5%)
History of hamstring injury; n (%)					
- Unknown/missing	6 (2%)	3 (1%)	1 (3%)	3 (1%)	1 (2%)
<ul> <li>Non-professional (Recreational/competitive)</li> </ul>	121 (33%)	102 (34%)	11 (36%)	85 (33%)	22 (42%)

č	(n= 368)	No reinjury	Reinjury	No reinjury	Reinjury
MRI findings	Total population	2 months reinjury (n= 330)		12 months reinjury (n= 308)	
Time to RTP (days)*	31 (21-46)	31 (21-46)	32.50 (19.25-44.50)	31 (21-48.50)	31 (19-42)
		(n=299; 91%)	(n=31; 9%)	(n=256; 83%)	(n=52; 17%)
	(n= 368)	No reinjury	Reinjury	No reinjury	Reinjury
Return to play	Total population	2 months reinjury (n= 330)		12 months reinjury (n= 308)	
- force deficit (Newton)*					
<ul> <li>Uninjured leg (Newton)**</li> </ul>	37.300 (5.000-74.575)	37.300 (3.900-75.000)	20.800 (5.600-81.550)	37.300 (2.00-76.500)	23.500 (8.700-67.175)
- Injured leg (Newton)**	134.761 (±54.628) 176.611 (±48.563)	134.683(±54.986) 176.766 (±48.883)	147.314 (±54.330) 186.224 (±55.326)	133.036 (±54.337) 174.134 (±48.471)	147.306 (±49.814) 187.355 (±49.323)
Isometric knee flexion resistance force in 90° (Newton)	134.761 (±54.628)	124 692(+54 096)	147.314 (±54.330)	133.036 (±54.337)	147.306 (±49.814)
- force deficit (Newton)*			. ,		
- Uninjured leg (Newton)*	94.200 (42.325-150.100)	97.100 (43.000-149.550)	72.000 (40.950-183.400)	94.200 (41.100-149.100)	98.550 (44.425-179.475)
- Injured leg (Newton)**	243.000 (203.400-287.000)	247.100 (205.000-289.300)	227.500 (196.00276.300)	245.200 (202.700-287.150)	240.100 (199.000-278.60
Isometric knee flexion resistance force in 15° (Newton)*	145.254 (±79.507)	145.764 (±79.746)	138.686 (±79.660)	145.401 (±78.703)	141.202 (±82.233)
- Deficit in flexibility (degrees)*					
- Uninjured leg angle (degrees)*	10.000 (2.000-23.000)	13.927 (±15.778)	10.000 (2.000-20.000)	10.000 (2.000-23.500)	8.000 (0-20.000)
<ul> <li>Injured leg angle (degrees)**</li> </ul>	60.000 (50.000-72.000)	62.000 (50.000-72.000)	55.000 (50.000-61.000)	62.000 (50.000-72.000)	56.000 (50.000-68.000)
Passive straight leg raise test (degree)	70.473 (±17.256)	70.390 (±17.420)	63.621 (±17.885)	69.732 (±17.127)	65.082 (±17.560)
- Deficit in flexibility (degrees)					
- Uninjured leg angle (degrees)	15.000 (4.750-33.050)	15.000 (4.000-33.600)	11.000 (3.250-26.500)	13.000 (4.000-33.500)	18.500 (5.000-28.500)
- Injured leg angle (degrees)	84.600 (67.000-121.250)	85.000 (70.000-125.000)	90.000 (62.250-138.500)	85.000 (70.000-121.250)	90.000 (67.000-140.000)
Active knee extension test (degree)*	65.000 (40.000-105.500)	66.000 (41.650-109.000)	82.500 (41.000-128.000)	65.000 (40.000-102.000)	82.500 (42.500-128.00)
- Unknown					
- Absent	98 (26.6%)	86 (28.8%)	5 (16.1%)	81 (31.6%)	9 (17.3%)
- Present	147 (39.9%)	118 (39.5%)	9 (29.0%)	96 (37.5%)	17 (32.7%)
Discomfort during Active Knee Extension; n (%)	123 (33.4%)	95 (31.8%)	17 (54.8%)	79 (30.9%)	26 (50%)

		(n=299; 91%)	(n=31; 9%)	(n=256; 83%)	(n=52; 17%)
Involved muscles; n (%)					
- Biceps Femoris	284 (77.2%)	231 (77.3%)	24 (77.4%)	194 (75.8%)	44 (84.6%)
- Semimembranosus/Semitendinosus	84 (22.8%)	68 (22.7%)	7 (22.6%)	62 (24.2%)	8 (15.4%)
Modified Peetrons grading; n (%)					
- Grade 1	126 (34.2%)	100 (33.4%)	10 (32.3%)	85 (33.2%)	17 (32.7%)
- Grade 2	242 (65.8%)	199 (66.6%)	21 (67.7%)	171 (66.8%)	35 (67.3%)
Tendon involvement; n (%)					
- No tendon involvement	173 (47.0%)	140 (46.8%)	14 (45.2%)	118 (46.1%)	26 (50%)
- Tendon involvement	195 (53.0%)	159 (53.2%)	17 (54.8%)	138 (53.9%)	26 (50%)
MTJ involvement; n (%)					
- No MTJ involvement	104 (28.3%)	90 (30.1%)	4 (12.9%)	79 (30.9%)	8 (15.4%)
- MTJ involvement	264 (71.7%)	209 (69.9%)	27 (87.1%)	177 (69.1%)	44 (84.6%)
Extent of oedema (cm)*					
- Anteroposterior (cm)	1.900 (1.100-3.095)	1.800 (1.100-3.020)	2.000 (1.190-3.020)	1.800 (1.100-3.200)	1.500 (0.935-2.590)
- Transverse (cm)	2.200 (1.400-3.200)	2.200 (1.440-3.125)	1.855 (1.050-3.275)	2.230 (1.400-3.200)	2.020 (1.410-3.200)
- Craniocaudal (cm)	13.200 (8.300-20.000)	13.000 (8.238-19.925)	13.750 (7.275-16.800)	13.200 (8.200-19.900)	13.000 (7.700-16.225)
Extent of haematoma (cm)*					
- Anteroposterior	0.000 (0.000-0.700)	0.000 (0.00-0.700)	0.550 (0.000-0.900)	0.000 (0.000-0.600)	0.500 (0.000-0.800)
- Transverse	0.000 (0.000-0.820)	0.000 (0.00-0.810)	0.620 (0.000-0.955)	0.000 (0.000-0.810)	0.600 (0.000-0.900)
- Craniocaudal	0.000 (0.000-2.475)	0.000 (0.000-2.400)	1.700 (0.000-3.000)	0.000 (0.000-2.300)	1.500 (0.000-3.000)
IM tendon disruption; n (%)					
- No IM tendon disruption	183 (45.7%)	146 (48.2%)	13 (41.9%)	129 (50.4%)	23 (44.2%)
- IM tendon disruption	168 (49.7%)	146 (48.8%)	17 (54.8%)	124 (48.4%)	28 (53.8%)
- Unknown	17 (4.6%)	9 (3%)	1 (3.2%)	3 (1.2%)	1 (1.9%)
Complete IM tendon disruption; n (%)					
- No complete disruption	324 (88.0%)	266 (89%)	29 (93.5%)	232 (90.6%)	49 (94.2%)

20

- Complete disruption	27 (7.3%)	24 (8%)	1 (3.2%)	21 (8.2%)	2 (3.8%)
- Unknown	17 (4.6%)	9 (3%)	1 (3.2%)	3 (1.2%)	1 (1.9%)
Length of IM tendon disruption (cm)*	0.000 (0.000-5.600)	0.000 (0.000-5.600)	1.500 (0.000-5.225)	3.202 (0.000-5.650)	2.727 (0.000-4.800)
Presence of waviness; n (%)					
- No present	216 (58.7%)	178 (59.5%)	18 (58.1%)	152 (59.4%)	37 (71.2%)
- Present	131 (35.6%)	109 (36.5%)	11 (35.5%)	98 (38.3%)	13 (25%)
- Unknown	21 (5.7%)	12 (4%)	2 (6.5%)	6 (2.3%)	2 (3.8%)
Presence of fibrosis; n (%)					
- No present	322 (87.5%)	269 (90%)	26 (83.9%)	231 (90.2%)	45 (86.5%)
- Present	32 (8.7%)	23 (7.7%)	4 (12.9%)	20 (7.8%)	6 (11.5%)
- Unknown	14 (3.8%)	7 (2.3%)	1 (3.3%)	5 (2%)	1 (1.9%)

Abbreviations: IQR = Interquartile range; SD = Standard Deviation; RTP = return to play, MTJ = myotendinous junction, IM = Intramuscular.

\*Presented in median (IQR).

\*\*Presented in mean (SD).

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