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1 **Is knee neuromuscular activity related to anterior cruciate ligament injury risk? A pilot**
2 **study**

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24 **ABSTRACT**

25 *Background:* Limited evidence exists on neuromuscular risk factors for anterior cruciate
26 ligament injuries, with most work mainly focussing on the strength of hamstrings and
27 quadriceps muscles. In this prospective pilot study, we explored if neuromuscular activation
28 patterns of the quadriceps and hamstrings during a drop vertical jump influence anterior cruciate
29 ligament injury risk.

30 *Methods:* Forty-six female athletes performed a drop vertical jump at baseline. Injuries were
31 monitored throughout a one year follow-up. Neuromuscular activation patterns of the vastus
32 medialis, vastus lateralis, hamstrings medialis and hamstrings lateralis, and selected landing
33 kinematic and kinetic profiles (knee flexion, knee abduction and hip flexion angles, and knee
34 abduction moments), were compared between athletes who sustained a non-contact anterior
35 cruciate ligament injury and those who remained injury-free. EMG vector fields were created
36 to represent neuromuscular activation patterns of muscle pairs around the knee joint rather than
37 only considering individual muscle activations, and compared using Statistical Parametric
38 Mapping.

39 *Results:* Four athletes sustained an anterior cruciate ligament injury. Significantly greater
40 {hamstrings medialis, hamstrings lateralis}, {vastus lateralis, hamstrings lateralis} and
41 {hamstrings lateralis, vastus medialis} activations, mainly due to greater hamstrings lateralis
42 activation, were found in the injured group around peak loading and just before take off ($P <$
43 0.001). No group differences were found in knee flexion, knee abduction and hip flexion angles,
44 or knee abduction moments.

45 *Conclusions:* This pilot study revealed initial evidence that athletes already showed altered
46 neuromuscular activation patterns prior to sustaining an anterior cruciate ligament injury,
47 namely increased lateral and posterior muscle activations.

48 **Key words:** ACL injury, neuromuscular activation, risk factor, drop vertical jump, injury
49 prevention, electromyography

50

51 **1. INTRODUCTION**

52 Anterior cruciate ligament (ACL) injuries are common during dynamic sports activities in the
53 young, active population and often have important short and long-term physical, psychological
54 and professional consequences [1], resulting in a lengthy absence from sports and high
55 economic cost for society [2]. Therefore, establishing neuromuscular and biomechanical risk
56 factors for ACL injury can assist the development of effective prevention programmes [3] as
57 both neuromuscular and biomechanical risk factors are potentially modifiable through
58 interventions [4,5]. To establish the risk factors for a specific injury, prospective studies with
59 injury as primary outcome are needed. So far, many studies assessed neuromuscular and/or
60 biomechanical changes after ACL injury or after intervention programs but very few
61 prospective studies assessed ACL injury risk with ACL injury as primary outcome [6,7]. To
62 date, only 4 prospective studies assessed neuromuscular risk factors for ACL injuries [8–11],
63 and only 4 prospective studies assessed biomechanical risk factors [12–15], delivering
64 contradictory outcomes.

65 Of the four prospective studies investigating the relationship between neuromuscular factors
66 and ACL injury risk, three studies focussed on muscle strength (isokinetic strength of the
67 quadriceps and hamstrings, and reciprocal muscle strength ratios) [8,9,11]. Myer et al. [11]
68 found that 22 female athletes who went on to sustain an ACL injury demonstrated decreased
69 hamstrings strength and unchanged quadriceps strength during concentric isokinetic testing
70 (300°/s) compared to 88 female controls. Similarly, Söderman et al. [9] found that 5 ACL
71 injured female athletes out of 146 players showed a reduced hamstrings to quadriceps ratio

72 (H/Q ratio) during concentric isokinetic strength tests (90°/s) prior to injury. However, on the
73 contrary, in the study of Uhorchak [8] no differences were found in H/Q ratios during isokinetic
74 strength tests at 60°/s (concentric action of quadriceps, eccentric action of hamstrings) between
75 24 ACL injured adolescents and 895 non-injured adolescents (males and females). Only one of
76 the four studies focused on neuromuscular activation patterns [10]. Zebis et al. [10] investigated
77 neuromuscular activation patterns during cutting maneuvers in 55 elite female athletes
78 (handball and soccer). The 5 athletes that sustained an ACL injury during the follow-up period
79 (2 match seasons) showed reduced activity of the semitendinosus and increased activity of the
80 vastus lateralis during the preparatory phase (10ms before initial contact) [10].

81 Based on the limited evidence, we hypothesize that imbalances in neuromuscular activation do
82 play a role in ACL injury risk. Furthermore, as neuromuscular activation patterns can be
83 modified by prevention programs [16], more prospective studies on neuromuscular risk factors
84 for ACL injury are needed in order to optimize prevention programmes. [17]. However,
85 neuromuscular risk factors first need to be identified, and if possible independently confirmed
86 by further prospective studies. This is very time consuming and often expensive as one has to
87 screen a large number of participants and subsequently follow them over an extensive period
88 of time to ensure that an adequate number of injurious events take place. The search for
89 neuromuscular factors could therefore benefit from pilot work in which a comprehensive
90 exploratory analysis precedes hypothesis testing [18,19]. Therefore, the aim of this pilot study
91 was to help justify future large cohort studies by exploring whether female athletes who went
92 on to sustain a non-contact ACL injury already showed meaningful differences in their knee
93 neuromuscular activation patterns during the landing phase of a bilateral drop vertical jump
94 (DVJ) prior to injury when compared to uninjured controls.

95

96 **2. METHODS**

97 **2.1. Participants**

98 Forty-six female athletes (21 soccer, 9 handball and 16 volleyball) aged between 16 – 28 years
99 participated in this study. All athletes, who were members of a an elite level team (first national
100 division), were tested at the beginning of their respective playing season and injuries were
101 monitored for one year. Participants who had a previous ACL or posterior cruciate ligament
102 injury, a previous lower extremity injury within the three months prior to testing, or another
103 lower extremity injury within the one-year follow-up period, had been excluded. All
104 participants provided informed consent, and the study was approved by the local ethics
105 committee.

106 **2.2. Injury registration**

107 The medical staff of each involved team registered all lower extremity time loss injuries during
108 follow-up. A time loss injury was defined as having occurred during sports participation and
109 resulting in being unable to take a full part in future training or match play [20]. We adapted
110 this definition to include “unable to take a full part in future training or match play for at least
111 two consecutive weeks”.

112 **2.3. Test protocol**

113 Each test session started with a standardized warm-up (two series of eight bilateral squats and
114 eight bilateral jumps) [21–23]. Body mass and height were measured before the test session
115 using scales (SECA, Hamburg, Germany) and a portable stadiometer (SECA, Hamburg,
116 Germany). Standardized indoor footwear (Indoor Copa, Kelme, Elche, Spain) was worn and
117 where necessary, long hair was tied up to avoid marker occlusion.

118 Subsequently, all athletes were asked to perform bilateral DVJ's. Bilateral DVJ's are commonly
119 used for screening in clinical settings to assess injury risk [13,14]. The protocol is briefly

120 summarized below. Participants were instructed to drop off a 0.3 m high box with their feet
121 initially positioned 0.2 m apart on the box, and upon landing to immediately perform a
122 maximum vertical jump. Participants were also instructed to reach upwards with both hands as
123 high as possible, as if performing a block in volleyball [24]. Participants were allowed to
124 familiarize themselves with the tasks by performing three practice repetitions before the start
125 of the tests. Subsequently, a minimum of three valid trials were completed. A trial was excluded
126 if subjects jumped off the box instead of dropping, if both feet did not land on the force plates,
127 if subjects reached upwards with only one hand, or if subjects clearly lost balance upon landing
128 [14]. A short rest period between consecutive trials was permitted to avoid fatigue [21].

129 **2.4. Data collection**

130 A wireless EMG system (Zerowire, Aurion, Milan, Italy) was used to record muscle activity at
131 1000Hz of the vastus lateralis (VL), vastus medialis (VM), biceps femoris (referred to as
132 hamstring lateralis, HL) and semitendinosus (referred to as hamstring medialis, HM) using
133 surface electrodes positioned according to the SENIAM guidelines [25]. All electrode locations
134 were shaved and gently cleaned with 70% isopropyl alcohol to reduce skin impedance. Silver-
135 silver chloride, pre-gelled bipolar surface EMG electrodes (Ambu Blue Sensor, Ballerup,
136 Denmark) were placed over the muscle belly and aligned with the expected muscle fibre
137 orientations, with 2 cm inter-electrode distance. Fauth et al. [26] observed that surface
138 electromyography (EMG) was reliable for measuring mean muscle activation of HL, HM, VL
139 and VM during the performance of a DVJ (ICC 0.83-0.97). As we were interested in the muscle
140 activation during the entire landing phase (from initial contact until take off), rather than the
141 mean muscle activation, we additionally assessed the reliability of the EMG data in 4 uninjured
142 subjects. These analyses showed both high intra- and inter-session reliability (see Appendix).

143 Secondary to the neuromuscular activation patterns, we also measured knee and hip flexion
144 angles, knee abduction angles and knee abduction moments as these have previously been

145 identified as potential biomechanical risk factors [12–14]. Therefore, we recorded three-
146 dimensional kinematic data using six MX-T20 optoelectronic cameras (VICON, Oxford, UK)
147 sampling at 100 Hz, synchronized with data recorded from two 0.8 x 0.3 m² force plates (AMTI,
148 Watertown, USA) sampling at 1000 Hz.. Each participant had 44 spherical reflective markers
149 positioned according to the eight segment ‘Liverpool John Moores University’ model including
150 feet, upper and lower legs, pelvis and trunk. This model was previously described in detail and
151 shown to be reliable for measuring kinematics and kinetics during DVJ [21].

152 **2.5. Data analysis**

153 All modelling and data processing were undertaken in Visual 3D (v.4.83, C-Motion, Kingston,
154 ON, Canada). Only the first landing (first contact) within each DVJ trial was used for analysis
155 [21]. Raw EMG signals were high pass filtered using a digital filter at a cut-off frequency of
156 10 Hz, full wave rectified, and low-pass filtered with a 4th order zero-lag Butterworth filter at a
157 cut-off frequency of 6 Hz. The EMG signal amplitudes were subsequently normalized to the
158 maximum root mean square amplitude (over a period of 100 ms) of 3 isometric maximum
159 voluntary contractions. Marker trajectories and forces were filtered using a 4th order low pass
160 Butterworth filter with a cut-off frequency of 18 Hz [27]. Initial contact and take off events
161 were created when the vertical force crossed a 20 N threshold. Knee and hip flexion angles,
162 knee abduction angles and moments were calculated using inverse dynamics. External joint
163 moments are described in this study; i.e. an external knee abduction moment will abduct the
164 knee (move the distal end of the tibia away from the midline of the participant’s body).

165 Kinetic, kinematic and EMG data were time normalized to 101 data points starting at 100 ms
166 before initial contact until take off (Fig. 1). The short time period prior to initial contact was
167 included based on Zebis et al. [10] who showed that neuromuscular pre-activity might be a risk
168 factor for ACL injury albeit during cutting. Therefore 0% of normalised time corresponds to
169 100 ms prior to initial contact and 100% corresponds to take off, whereas initial contact is

170 situated around 17% and peak loading around 50%. For all included variables, the average of
171 three trials was calculated for each participant and then means were calculated across the
172 groups.

173 **2.6. Statistical analyses**

174 Participants' baseline characteristics were compared between the control and the ACL-injured
175 group using independent sample Student's t-tests (Table 1).

176 To compare neuromuscular activation patterns of muscle pairs around the knee joint rather than
177 only considering individual muscle activation patterns as independent observations, time-
178 varying EMG vector fields were created. A major advantage of this technique is that it accounts
179 for inter-muscle covariance. The following vector fields were created to represent the vector
180 magnitude of anatomically relevant muscle groupings: an overall EMG {VM,VL,HM,HL}
181 (time) vector field, an anterior {VM,VL} (time) vector field, a lateral {VL,HL} (time) vector
182 field, a posterior {HM,HL} (time) vector field, a medial {HM,VM} (time) vector field and 2
183 diagonal vector fields: a {HM, VL} (time) vector field, and a {HL, VM} (time) vector field
184 [28]. To statistically compare these vector fields between groups, seven Hotelling's T^2 tests
185 were conducted (the vector field equivalent of an independent samples Student's t-test) using
186 Statistical Parametric Mapping (SPM) [28,29] for which Alpha was set at 0.05. SPM has
187 become well established in biomechanical research. Basically, first a test statistic is calculated
188 for each time node (e.g., t-values). Second, the threshold problem of repeated comparisons over
189 time is handled by modeling the behavior of random time-varying signals as a random field
190 based on signal smoothness [18,30]. Given the differences in group sizes, equality of variance
191 between groups was not assumed. Post-hoc SPM independent t-tests were used to estimate the
192 contribution of individual muscles to vector field based group differences. Detailed examples,
193 theoretical background and interpretations of vector field and SPM statistics are available
194 elsewhere [28,29]. In summary, if at certain times of the landing the T^2 or t -statistic exceeded

195 the critical threshold then a significant difference between groups was assumed of which the
196 probability (P -value) depends on the magnitude and duration of the threshold crossing.

197 Finally, external knee abduction moments, knee abduction angles, knee flexion and hip flexion
198 angles were compared between the ACL-injured and the control group across the landing phase
199 (excluding the 100ms pre-landing phase), again using SPM independent sample t-tests.

200

201 3. RESULTS

202 During the one year follow-up, four participants sustained a non-contact ACL injury. All
203 injuries occurred during competitive match play (1 in soccer, 2 in handball, 1 in volleyball).
204 These ACL injuries were diagnosed with magnetic resonance imaging (MRI) and all underwent
205 ACL reconstruction. Three out of four ACL injuries affected the non-dominant leg. The
206 dominant leg was defined as the preferred leg to kick a ball [31].

207 Of the remaining non-ACL-injured participants, five participants sustained an ankle inversion
208 trauma, and two other participants sustained an overuse knee injury during the one year follow-
209 up period (one pes anserinus tendinopathy and one degenerative lateral meniscus lesion with
210 associated cyst). As the purpose of this study was to assess ACL injury risk, the latter seven
211 participants who got injured during the follow-up were excluded, resulting in a control group
212 of 35 participants in total (Fig. 2). No significant differences were found for age, weight, height
213 and body mass index between the ACL-injured and the control group (Table 1).

214 3.1. Within group analysis

215 Both in the control group and the ACL-injured group, no significant differences were found
216 between limbs for the amplitude of the overall EMG {VM,VL,HM,HL} (time) vector, for knee
217 abduction moments, knee abduction angles, or for knee or hip flexion angles ($P > 0.05$). As our
218 study was not concerned with leg preference and since there were no significant leg differences
219 we pooled our data across legs resulting in a control sample consisting of 70 legs and an ACL-
220 injured sample consisting of 8 legs.

221 3.2. Between group analysis

222 To investigate whether there was any difference in neuromuscular activation between the ACL-
223 injured athletes and the non-injured athletes, the overall multi-muscle EMG vector
224 {VM,VL,HM,HL} was compared between both groups. A significant difference in the overall

225 multi-muscle EMG vector {VM,VL,HM,HL} was found between the ACL-injured group and
226 the control group from peak loading until the last part of the acceleration phase just before take
227 off (34-100% time; $P < 0.001$).

228 Subsequently, 6 Hotelling's T^2 tests were performed to identify which muscle pairs were
229 significantly different between both groups and were thus responsible for the significant
230 differences in the overall multi-muscle EMG vector. These Hotelling's T^2 tests showed
231 significantly greater amplitudes of the {HL, HM} (t) vector, the {HL, VL} (t) vector and the
232 {HL,VM} (t) vector in the ACL-injured group compared to the control group during the peak
233 loading phase (43-80% time, $P < 0.001$; 55-71% time, $P < 0.001$ and 58-79% time, $P < 0.001$
234 respectively) and during the last part of the acceleration phase just before take off (93-100%
235 time; $P < 0.001$, 84-100% time; $P < 0.001$ and 90-100% time; $P = 0.002$ respectively) (Fig. 3
236 and 4). Furthermore significantly greater amplitudes in {HM, VL} (t) vector were found in the
237 ACL-injured group the last part of acceleration phase just before take off (91-100% time, $P <$
238 0.035).

239 Finally, post-hoc SPM independent t-tests showed significantly greater HL amplitudes in the
240 ACL-injured group compared to the control group during the peak loading phase (51-78%; P
241 < 0.001) and during the last part of the acceleration phase just before take off (92-100% time;
242 $P = 0.006$) (Figure 3). No differences were found between groups for the individual muscle
243 activation amplitudes of VM, VL and HM (Fig. 3 and 5).

244 Furthermore, no significant differences in external knee abduction moments, knee abduction
245 angles, knee flexion, and hip flexion angles were found between the ACL injured group and the
246 control group (Fig. 6). However, a clear tendency was observed for the ACL-injured group to
247 show larger knee abduction angles and smaller knee flexion angles, seen by the SPM{t}-curve
248 being consistently respectively below and above 0.

249 4. DISCUSSION

250 The purpose of this pilot study was to prospectively explore the presence of neuromuscular
251 differences in athletes who would go on to suffer an ACL injury compared to uninjured controls.
252 Distinct muscle activation patterns were found in the ACL injured group compared to controls,
253 primarily showing greater activation levels during peak loading and just before take-off, mostly
254 due to increased lateral hamstring activations.

255 It is commonly accepted that quadriceps and hamstrings co-contractions are necessary to ensure
256 dynamic knee joint stability during dynamic activities [32]. Balanced hamstrings to quadriceps
257 co-contraction might protect against anterior tibial translation [33,34] and appropriate medial
258 to lateral co-contraction of the hamstrings and quadriceps should protect the knee joint from
259 high knee abduction loads [35]. However, when the lateral hamstrings are favored in this co-
260 contraction, then this may have detrimental consequences leading to increased risk of injury.
261 For example, Serpell et al. [36] have shown that during a step-up task, selectively increasing
262 the lateral hamstrings-quadriceps co-activation results in more ACL elongation, while stronger
263 medial hamstring-quadriceps co-activation reduces ACL elongation and knee joint rotation,
264 abduction, translation and distraction. Furthermore, the prospective study of Zebis et al. [10]
265 demonstrated that a neuromuscular imbalance between lateral and medial activation during a
266 side cutting manoeuvre was associated with increased ACL injury risk. The authors suggested
267 that sufficient HM activity against VL activity is important to compress the medial knee joint
268 and thereby limit the risk of excessive abduction of the knee and minimizing the strain on the
269 ACL [10]. This is in agreement with Palmieri-Smith et al. [37], who showed that increased
270 activation of the vastus lateralis and hamstrings lateralis during the preparatory phase of a 1m
271 forward hop was associated with higher peak knee valgus angles in female athletes.

272 However, comparing our results with other studies should nonetheless be done with some
273 apprehension, as muscle activations were assessed in a variety of dynamic tasks. For example,
274 Husted et al. [38] showed that neuromuscular pre-activity of the HM, HL and VL was only low
275 to moderately correlated between side cutting manoeuvres and the DVJ. Therefore, we recently
276 investigated the relationship between muscle activation of quadriceps and hamstrings and the
277 kinematics of the knee and hip joint during the performance of a DVJ in female athletes [22].
278 We found that those athletes who had a more erect landing pattern (less knee and hip flexion),
279 showed an increased {VL, HL} activation.

280 Thus, we expected that the altered muscle activation patterns in the ACL-injured group would
281 go together with significant differences in the selected landing kinematics or kinetics (knee
282 abduction moment, knee abduction angle, knee flexion angle and hip flexion angle). In this
283 pilot study we did not find significant differences between groups but a clear tendency was
284 observed for the ACL-injured group to show larger knee abduction angles and smaller knee
285 flexion angles. The fact that landing kinematics and kinetics were not significantly different
286 between the ACL-injured group and the control group suggests that neuromuscular activation
287 patterns might be more sensitive for predicting injury risk in well-trained female athletes.
288 Probably, subtle alterations in muscle activation only influence the stability and
289 arthrokinematics of the knee joint and therefore may not necessarily result in significant
290 changes in joint angles and moments.

291 The risk of ACL injuries is multifactorial and therefore the results of this paper should neither
292 be interpreted in isolation nor prematurely. The aim of this study was not to find an ‘ultimate
293 predictive factor’ for ACL injury risk, or to criticize existing factors, but was to explore whether
294 neuromuscular activation patterns are worthwhile consideration in the multifactorial approach
295 to prevent ACL injuries. Recently, Bittencourt et al. [39] published a conceptual paper to
296 propose a new framework on sports injury prevention, in which injury prediction is based on

297 risk pattern recognition. They proposed a ‘web of determinants’ that visualizes the complex
298 interactions between the different risk factors. Based on the results of our pilot study we suggest
299 that also neuromuscular activation patterns may play a meaningful role in this ‘web of
300 determinants for ACL injury risk’. In the past, muscle activation could only be measured with
301 laboratory techniques and therefore it was not feasible to implement it in injury prevention
302 programmes. But recent technological developments (e.g. sport clothes with embedded textile
303 electrodes) made it possible to also measure muscle activation in field settings

304 To our knowledge, this prospective pilot study is the first study that comprehensively
305 investigated the relationship between neuromuscular activation patterns and the incidence of
306 ACL injuries, yet it comes with some limitations. First, we would like to reiterate that the
307 sample size of this pilot study was small and therefore we should be extremely cautious not to
308 generalize the observations as facts. Second, other muscles such as the gastrocnemius and glutei
309 can influence dynamic knee joint stability but these were not measured [40,41]. Third, as we
310 did not measure tibio-femoral contact forces, ACL strain, or actual joint kinematics for example
311 through the use of video fluoroscopy, we cannot describe the underlying mechanics that explain
312 whether a specific neuromuscular activation pattern would result directly in an increased ACL
313 injury strain. The calculation of medial and lateral tibio-femoral contact forces might reveal for
314 example an unloading phenomenon of the medial knee compartment compared to high contact
315 forces at the lateral knee compartment, but this would need to be confirmed.

316

317 **5. CONCLUSIONS**

318 This pilot study has prospectively demonstrated altered neuromuscular activation patterns in
319 ACL injured elite female team sports athletes. Participants who sustained an ACL injury
320 showed increased lateral hamstring activation during peak loading and the push off phase of a

321 DVJ. Previously suggested kinematic or kinetic indicators of increased risk of injury were not
322 confirmed, providing a preliminary rationale for including neuromuscular activations in future
323 large scale prospective studies.

324

325 **CONFLICT OF INTEREST**

326 This research did not receive any specific grant from funding agencies in the public,
327 commercial, or not-for-profit sectors.

328

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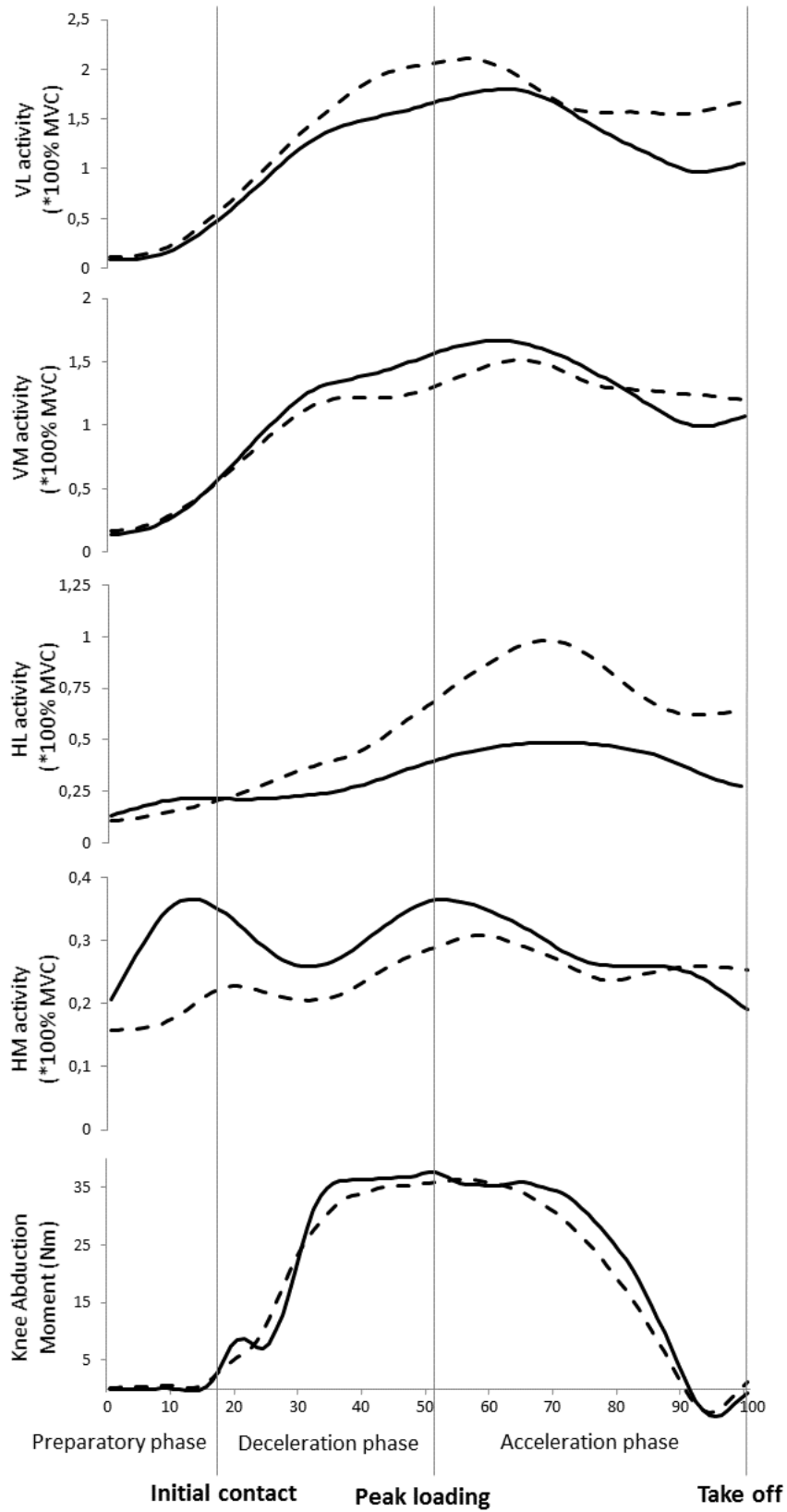
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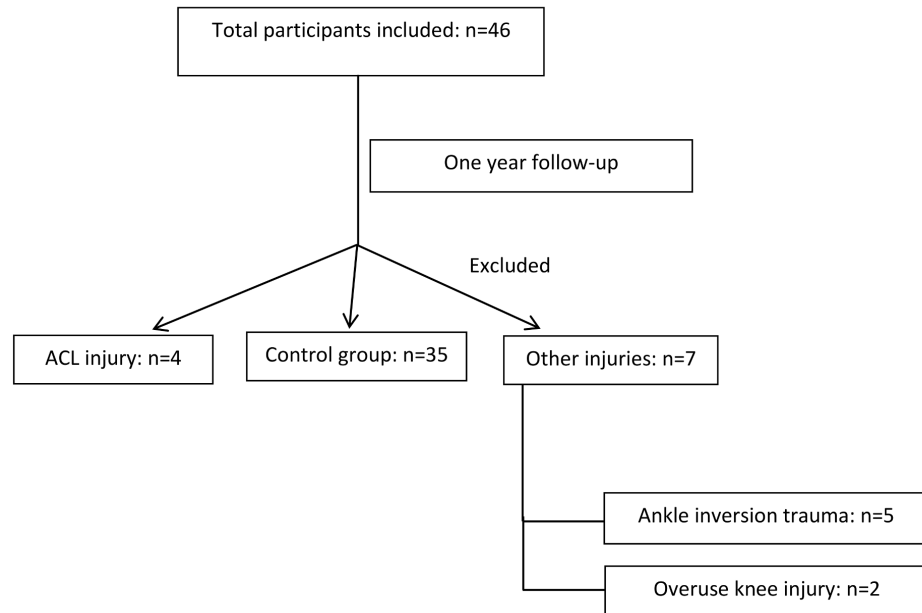
481

482 **Fig. 1**

483 Mean data of the ACL-injured group (dotted line) versus the control group (bold line) to
484 illustrate the different time periods during DVJ from 100ms before initial contact until take off.

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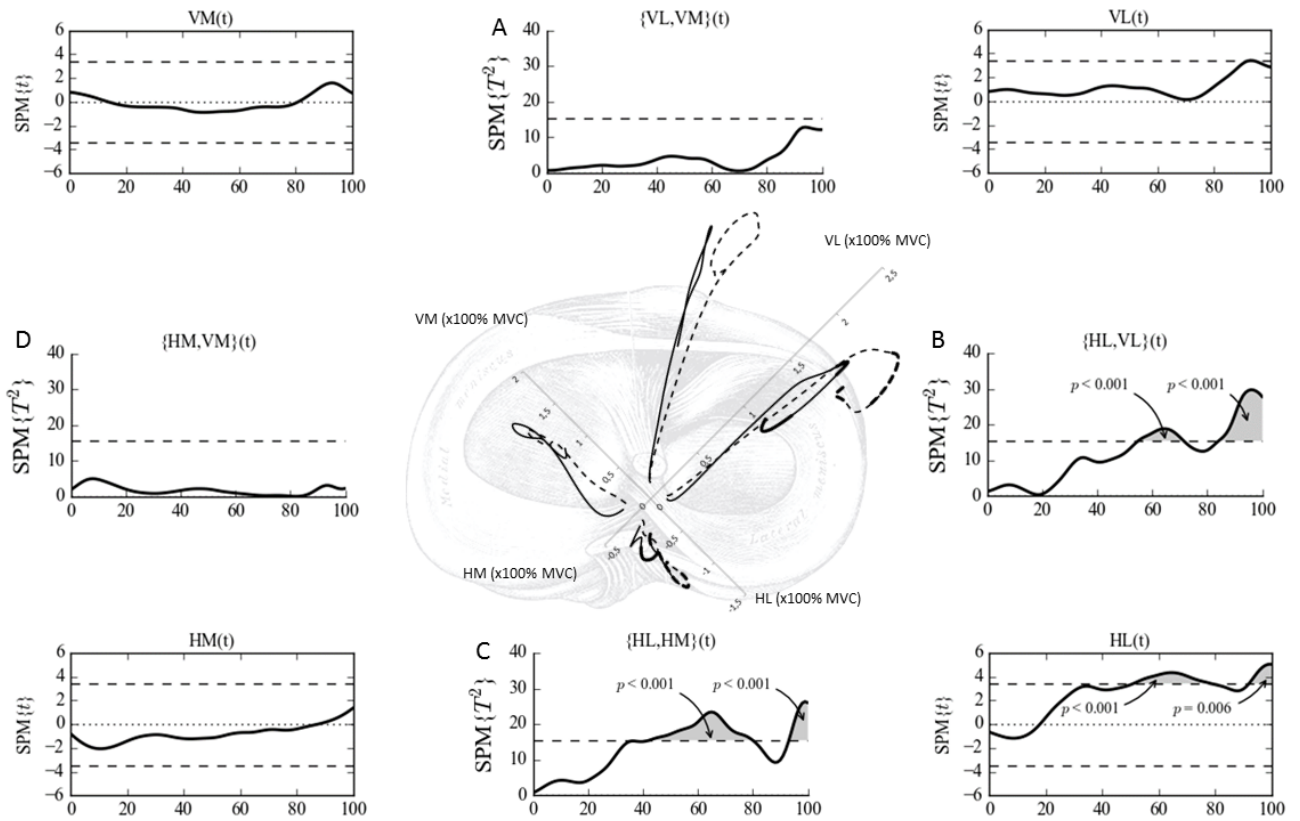


487

488 **Fig. 2**

489 Flowchart of the participants

490



491

492 **Fig. 3**

493 Top view representation of group differences.

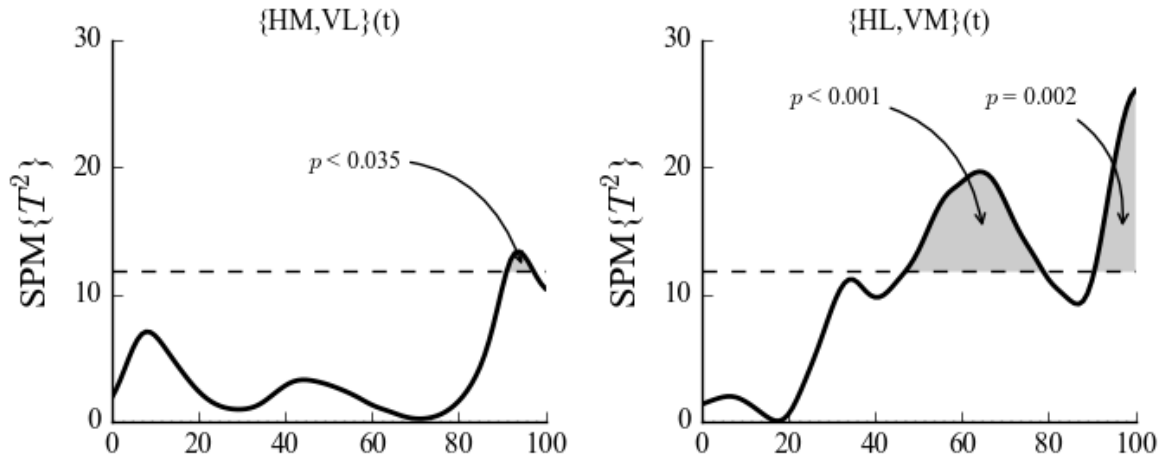
494 *Central figure:* Differences in two-muscle activation patterns between the ACL-injured (dotted
 495 line) and the control group (solid line) are visualized. The bold parts represent the time periods
 496 in which significant differences between groups were found.

497 *A,B,C,D:* Trajectory level SPM analyses show the differences between groups for the anterior
 498 {VM,VL}, lateral {HL,VL}, posterior {HM,HL} and medial {HM,VM} EMG vector. The
 499 horizontal dashed line represents the critical threshold ($P < 0.05$).

500 *Corner figures:* Trajectory level SPM analyses show differences between groups for the
 501 individual amplitudes of VM, VL, HM and HL. The horizontal dashed line represents the
 502 critical threshold ($P < 0.05$).

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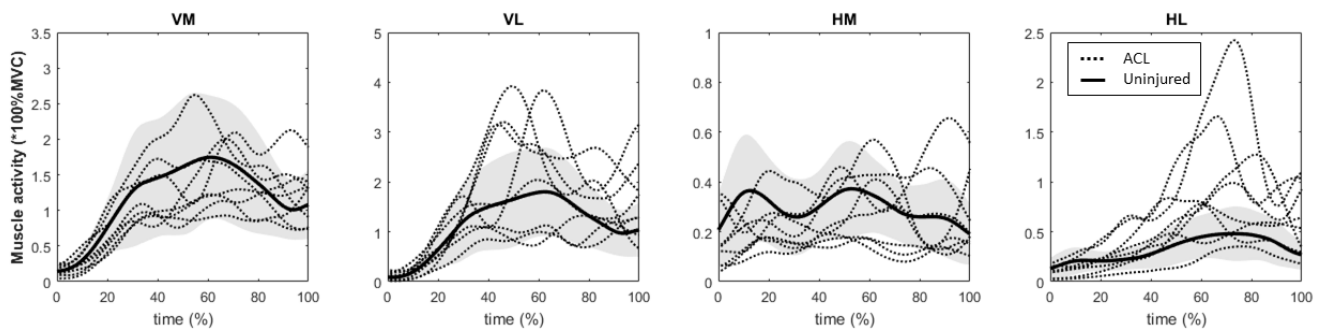
505

506 **Fig. 4**

507 Trajectory level SPM analyses show the differences between groups for the {HM, VL} and the
508 {HL, VM} EMG vector. The horizontal dashed line represents the critical threshold ($P < 0.05$).

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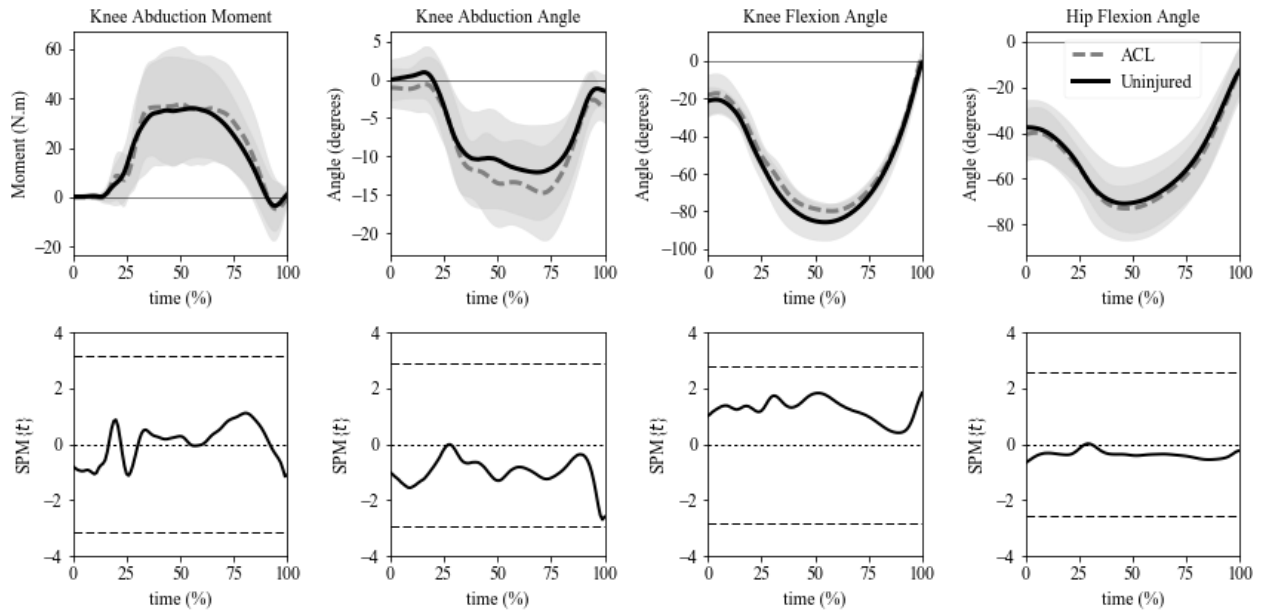
510



511 **Fig. 5**

512 This figure visualizes the activation of the HL, HM, VL and VM muscles. The 8 dotted lines
513 represents the data of the ACL injured athletes. The solid line represents the average activation
514 of the respective muscles in the control legs and the shaded zone represents the standard
515 deviation.

516



517

518 **Fig. 6**

519 The upper figures illustrate the mean data for the respective parameters. The dotted line
 520 represents the ACL-injured group and the bold line represents the control group. Standard
 521 deviation clouds are represented by the shaded areas. The lower figures illustrate the SPM
 522 output. No significant differences were found between groups for the external knee abduction
 523 moment, knee abduction angle, knee flexion angle and the hip flexion angle over the entire time
 524 period during DVJ. However, a clear tendency is observed for the ACL-injured group to show
 525 larger knee abduction angles and smaller knee flexion angles, seen by the being consistently
 526 respectively below and above 0.

527

Table 1: Participants characteristics

	Control group	ACL-injured group	P-value
Participants (n)	35	4	
Age, yrs (Mean ± SD)	20.69 ± 3.19	21.02 ± 2.96	0.85
Body height, m (Mean ± SD)	1.72 ± 0.09	1.72 ± 0.11	0.88
Body weight, kg (Mean ± SD)	64.94 ± 7.45	62.58 ± 6.94	0.44
BMI, kg/m ² (Mean ± SD)	22.05 ± 1.61	21.21 ± 1.02	0.31

n: number of participants; SD: standard deviation; BMI: body mass index

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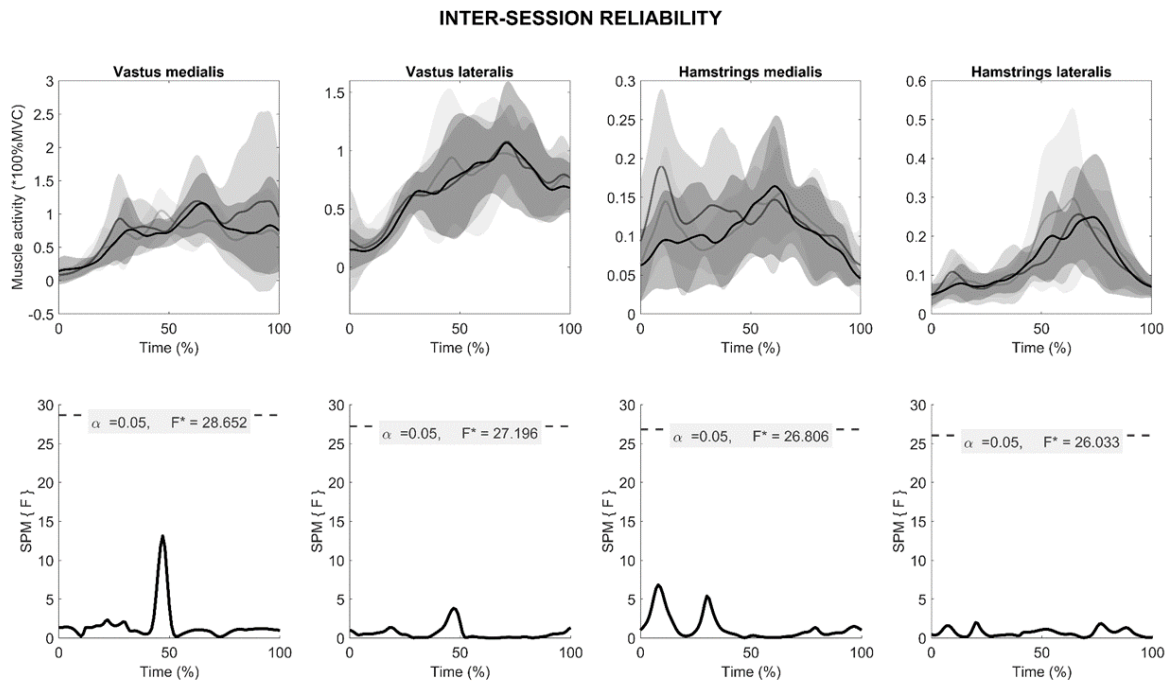
534 **Appendix – Reliability of time-varying EMG data**

535 To investigate the intra- and inter-session reliability of the time-varying EMG data, we did an
536 additional reliability study in 4 uninjured subjects.

537 All subjects (1 female, 3 males, age 28.5 ± 3.5 y height 185.6 ± 9.1 cm, weight 76.9 ± 9.6 kg) were
538 involved in recreational or competitive team sports (soccer, volleyball, handball and basketball)
539 and performed the same protocol as described in the manuscript on 3 different days. They
540 performed the protocol always at the same time during the day and all sessions were performed
541 in the same week (Monday, Wednesday and Friday). All data was processed in the same way
542 as described in the manuscript.

543 As our outcome is 1 dimensional (1D) data (e.g. emg curves) and not traditional 0 dimensional
544 (0D) data (e.g. peak/ mean values), we could not calculate traditional 0D ICC's for investigating
545 reliability. There are no explicit tests of reliability for 1D data yet, thus instead we addressed
546 reliability implicitly using ANOVA. In particular, if inter-session reliability is high, then we
547 would expect intra-subject variability to be greater than inter-session variability, and thus
548 produce a small session effect. Conversely, if inter-session reliability is low, then we would
549 expect a large session effect.

550 To implicitly investigate the inter-session reliability, we used an SPM one-way ANOVA with
 551 1 repeated measure (e.g. session). These analyses showed no significant differences in muscle
 552 activation between the 3 different sessions as the F-curve never exceeded the critical threshold
 553 (see fig. 1). As there was no session-effect, we can conclude that inter-session reliability is high.



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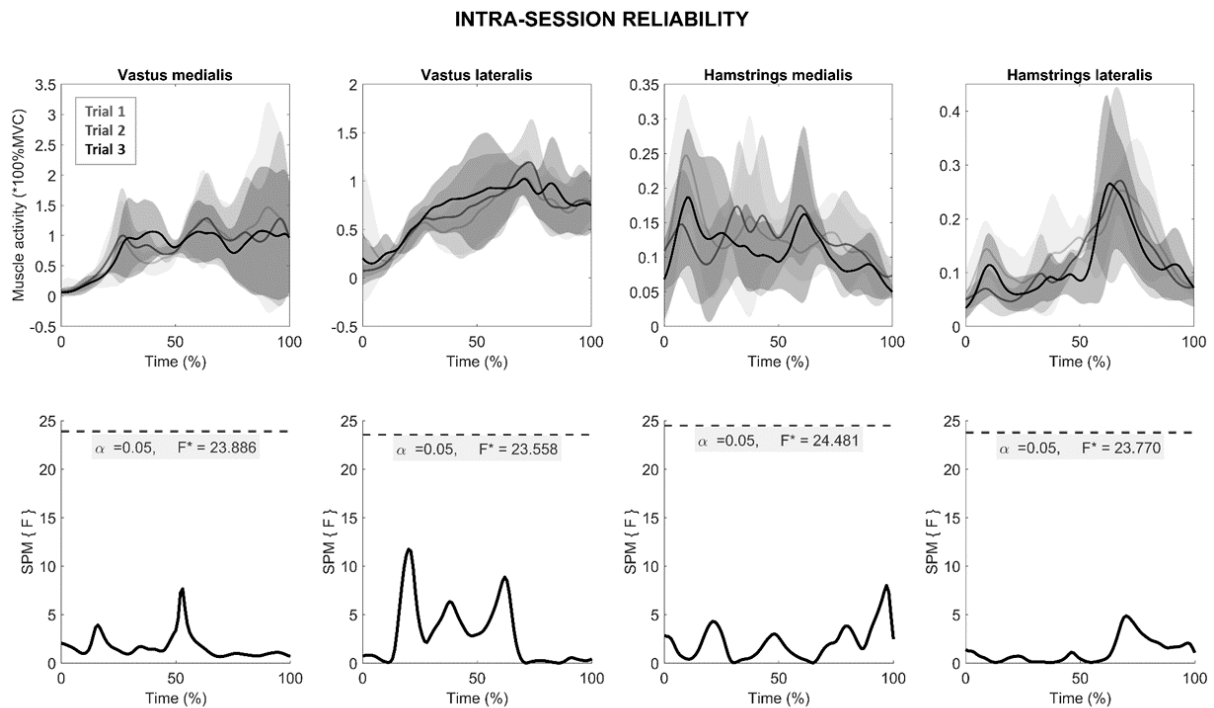
555 **Fig. 1.**

556 *The upper figures illustrate the mean data for session 1 (light grey), session 2 (dark grey) and*
 557 *session 3 (black) for the respective muscles. Standard deviation clouds are represented by the*
 558 *shaded areas. The lower figures illustrate the SPM output. No significant differences were*
 559 *found between sessions as the F-curve never exceeds the critical threshold (horizontal red*
 560 *dashed line).*

561

562 To implicitly investigate intra-session reliability, we performed another SPM one-way ANOVA
 563 with 1 repeated measure (e.g. trial) on the data of session 1. These analyses showed no

564 significant differences in muscle activation between the 3 different trials (see fig. 2). As there
565 was no trial-effect we can conclude that intra-session reliability is high.



566

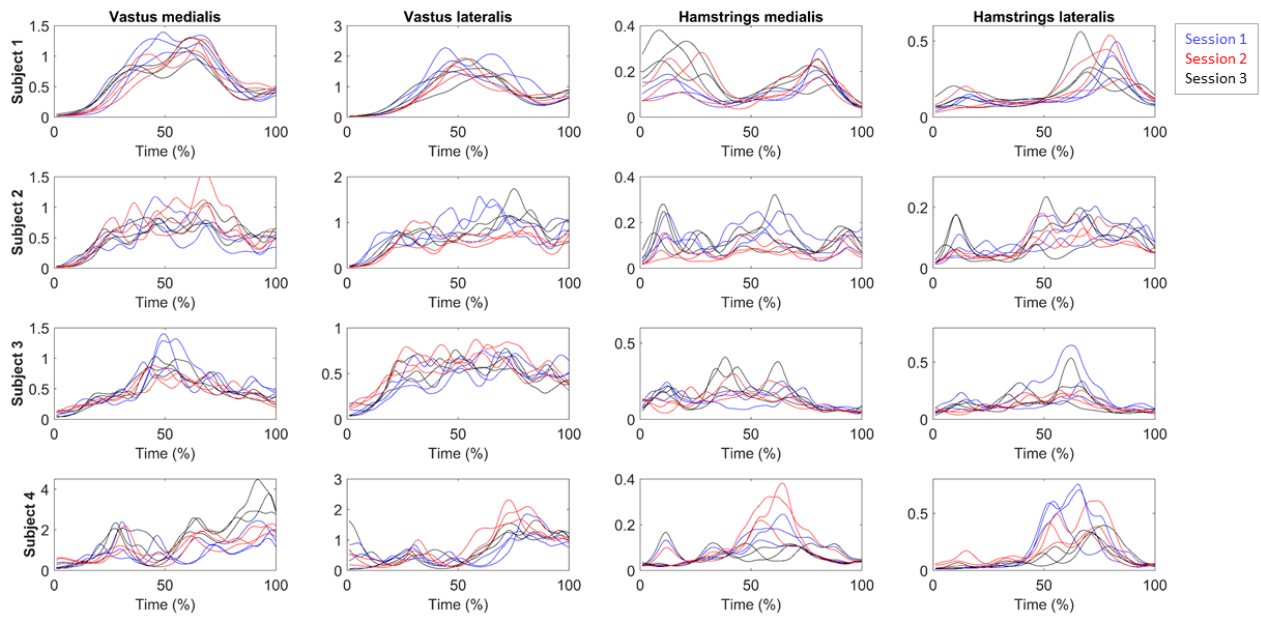
567 **Fig. 2.**

568 *The upper figures illustrate the mean data for trial 1 (light grey), trial 2 (dark grey) and trial 3*
569 *(black) of session 1 for the respective muscles. Standard deviation clouds are represented by*
570 *the shaded areas. The lower figures illustrate the SPM output. No significant differences were*
571 *found between trials as the F-curve never exceeds the critical threshold (horizontal dashed*
572 *line).*

573

574 Below (fig. 3) you can find all the individual emg data of every trial and for every session.

INDIVIDUAL DATA



575

576 **Fig. 3.**

577 *Each row represents the emg data of another subject. All subjects performed 3 trials per session*
578 *(session 1 = blue, session 2= red, session 3 = black).*

579

580 The figures above confirm that the time-varying data of the individual muscles is highly
581 reliable. As ANOVA testing does not exist for vectors, the same approach as described above
582 could not just be repeated on the vector data of the muscle pairs. However, the reliability of
583 muscle co-activation pairs relies on (1) the reliability of measuring each individual muscle
584 within the pairs and (2) the minimization of cross-talk between muscles. Whilst cross-talk
585 between muscles within the pairs would inflate the amount of co-variance, its impact on the
586 muscle pairs is the same as on the individual muscles. In both cases cross-talk has to be avoided
587 as much as possible, which is established by careful electrode placement.

588 We did not investigate inter-rater reliability as we only had 1 rater in this study. We can assume
589 that the rater was very consistent in electrode placement as both the quadriceps and hamstrings

590 muscles were very easy to palpate in our athlete population. Furthermore, the rater is a qualified
591 physiotherapist with extensive palpation experience, and used the SENIAM guidelines to
592 standardize his way of electrode placement.

593