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Original Research

Evaluation of patellar tendinopathy using the single leg decline squat test: Is pain location important?

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ABSTRACT

Study design: A cross-sectional study of non-elite volleyball players aged 13–17 years.**Objectives:** To evaluate the presence and location of pain during the single leg decline squat (SLDS) and compare patellar tendon thickness, structure, neovascularisation and symptom severity between SLDS-derived groups.**Methods:** 32 male and 25 female participants attending a 5-day volleyball training camp underwent clinical evaluation by SLDS, describing the location of pain during this test using a pain map. The patellar tendon was examined using ultrasound imaging, performed by an assessor blinded to other assessments. Differences between participants experiencing local patellar tendon pain (PTP), other knee pain (OKP) or no-pain during the SLDS were evaluated.**Results:** Fifteen (26.3%) participants experienced pain during the SLDS. Local PTP was recorded for 12.3% and OKP for 10.5% of right legs. The PTP group was distinguished from the other groups by larger thickness and cross-sectional area of the mid-patellar tendon ($p < 0.001$), more frequent neovascularisation ($p = 0.005$) and greater pain and disability ($p < 0.036$). No differences between OKP and no-pain groups was observed.**Conclusion:** Adolescent non-elite volleyball players reported symptoms indicative of patellar tendinopathy. In this cohort, the SLDS test combined with a pain map was associated with imaging and questionnaire-based outcomes.**Level of evidence:** Diagnosis, Level 2; Cross-sectional study.

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1. Introduction

Patellar tendinopathy has a high prevalence (44.6%) in adult volleyball athletes (Lian et al., 2005), and is the most frequent overuse injury in adolescent, elite athletes (Cassel et al., 2015; Gisslen et al., 2005). In younger athletes, the effects of mechanical loading may compound or interact with those of maturation (Mersmann et al., 2017; Waugh et al., 2012). These factors might expose young athletes to an increased risk of developing overuse injuries such as patellar tendinopathy. In young elite volleyball players, training volume and match exposure are predictors of the development of patellar tendinopathy (Visnes & Bahr, 2013), while other studies highlight the large ground reaction forces (~6 times body weight) during jumping as risk factors (Lian et al., 2005).

Some studies report higher prevalence rates in men, suggesting their greater capacity to generate force may increase their risk of injury (Lian et al., 2005). Most studies have focused on adult, elite athletes, while prevalence of patellar tendinopathy in adolescent athletes, with lower training loads is understudied. Research in non-elite youth may provide insight into potential relationships between maturation, training and susceptibility to injury.

Pain on performing a task that loads the tendon was recently established by Delphi study (Vicenzino et al., 2020) as a core domain for tendinopathy. Although there is no current gold standard test, the single leg decline squat (SLDS) is proposed as a provocation test to discriminate patellar tendon pain (Purdam, 2003). The original test was described using a 25° decline board to increase load on the patellar tendon, avoiding flexion beyond 60°

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(Purdam, 2003). Variations to this test (e.g. knee flexion to only 30°) are also described (Mendonca Lde et al., 2016). As the SLDS loads multiple structures in the knee (Zwerfer et al., 2007), the location of knee pain during the clinical test might provide additional useful information (Hannington et al., 2020). Localised pain over the inferior pole of the patella is described as a key diagnostic sign for patellar tendinopathy (van der Worp et al., 2012). However, a recent study revealed heterogeneity in pain location during the SLDS for elite basketballers with self-reported patellar tendinopathy (Hannington et al., 2020), questioning the ability to make inferences about a pathoanatomical source of pain.

The SLDS offers an alternative to ultrasound imaging and other validated measures of the severity of tendon pain and function, such as the Victorian Institute of Sport Assessment-Patella (VISA-P) (Visentini et al., 1998). It can be easily administered to screen large clinical samples and is free of necessity for advanced training. However, more research is needed to inform its utility as a clinical tool for inferring patellar tendon pain (PTP).

The primary aim of this research was to determine the proportion of non-elite adolescent volleyball athletes experiencing knee symptoms, and reporting no pain, local PTP or other knee pain (OKP) areas during the SLDS test. A secondary aim was to compare ultrasound imaging characteristics (tendon thickness, structure and neovascularisation), as well as the severity of self-reported knee symptoms between no pain, local PTP and OKP groups derived from the SLDS.

2. Materials and methods

2.1. Participants and experimental design

Fifty-seven adolescent volleyball athletes (32 male, 25 female) aged 13–17 years attending a 5-day training program in Brisbane, Australia <<BLIN>> agreed to participate in this cross-sectional study. The study was approved by Griffith <<BLIN>> University ethics committee (Approval # 2017/896). Participants (and legal guardians) signed informed consent prior to testing and the rights of the participants were protected. All assessments were performed over a 3-day period.

Participants completed questionnaires regarding demographics, training history and lead leg for spike jump. The Oslo Sports Trauma Research Center (OSTRC) ten item questionnaire was used to record “knee problems” (pain, ache, stiffness, swelling, instability, locking or other complaints) in one or both knees in the last week (Clarsen et al., 2013). Questions 1–4 were scored from 0 to 25, with the sum of all 4 items ranging from 0 to 100; 0 represents full participation without pain or symptoms (Clarsen et al., 2013). The 4-item severity score was validated in youth basketball players and found to be correlated with clinical evaluation (Owoeye et al., 2018). The VISA-P questionnaire was used to indicate severity of symptoms of patellar tendinopathy, with a score of 100 representing no limitations (Visentini et al., 1998). Left and right limbs were scored separately for questions 1–6 of the VISA-P.

2.2. Clinical evaluation

Standing height and weight were measured, followed by assessment of the SLDS test on each leg in a random order. While balancing on a 25° decline board with arms crossed and trunk vertical, participants flexed their knee to 60° and rated the intensity of pain experienced on a 0–10 numerical scale. If knee pain was present, participants were asked to select the area best matching the location from one of 6 areas illustrated on a pain map (Fig. 1) (van der Worp et al., 2012). Three trials with 30s rest between trials were performed and an average pain rating was used in analysis.

Test responses for each leg were classified as no-pain (pain scores in all trials = 0), patellar tendon pain (PTP, average pain >0 and pain location ‘e’ Fig. 1) or other knee pain (OKP) (average pain >0 and pain locations a-d or f).

2.3. Ultrasonography

A single examiner, blinded to the participant’s clinical evaluation and injury history, performed ultrasound imaging using a GE LOGIQ e with 12–4 MHz transducer (GE Healthcare, Wuxi, China). The participant was positioned in a supine lying position with the test leg (order of legs randomised) supported in 20° knee flexion. The transducer was first oriented longitudinal to the mid-tendon and proximal tendon, then oriented transverse to the mid-tendon, recording 2 or 3 images at each location. Tendon structure was graded as 0, homogenous echogenicity; 1, discrete hypo-echoic areas; 2, well-defined hypo-echoic areas and 3, extended hypo-echoic areas (Sunding et al., 2016). Using Power Doppler, tendon neovascularity was graded as 0, no neovessels; 1, a few solitary blood vessels; 2, multiple vessels; 3, neovascularisation spread to whole depth of tendon. For analysis, grades 1–3 were collapsed to represent abnormal tendon structure or neovascularisation on respective scales. Tendon thickness and cross-sectional area (CSA) were measured from longitudinal and transverse images, at the thickest part of the tendon (Sunding et al., 2016), using OsiriX Lite software and the average values used in analysis. Data extraction was repeated for 15 participants (26.3%) by the same examiner. Reliability was excellent for measurement of thickness (ICC, 95% CI: mid-tendon 0.959, 0.879 to 0.987; proximal tendon 0.974, 0.922 to 0.992) and good for CSA (0.793, 0.407 to 0.928), with coefficients of variation 2.8%, 2.5% and 8.7% and minimal detectable differences of 0.02 cm, 0.03 cm and 0.17 cm² respectively (Weir, 2005).

2.4. Statistics

Frequency and proportion of participants meeting no-pain, PTP and OKP criteria on SLDS were generated for both legs. As right and left legs are coupled and not independent, and because the prevalence of PTP was low for the left leg (n = 2), we performed group comparisons using data from the right leg for all participants. Normality of continuous data was confirmed using Shapiro Wilk tests and is presented as mean ± standard deviation, with the exception of OSTRC and VISA-P questionnaires and training hours, where data did not meet normality assumptions and is presented as median [IQR]. Ultrasonography, demographic and training variables were compared between No-pain, PTP and OKP groups using analysis of variance (ANOVA, post-hoc tests with Bonferroni correction) or Chi-squared (Gisslen et al., 2005) tests. Bias corrected Hedges’ g or odds ratios (and 95% confidence intervals) were calculated for significant pairwise comparisons (Centre for Evaluation and Monitoring, 2020). Kruskal-Wallis ANOVA (post-hoc Wilcoxon test) were used for group comparisons of questionnaires (OSTRC and VISA-P) and training hours. Analyses were tested using Stata 13.0 and a significance level of 0.05 adopted.

3. Results

Fifteen (26.3%) participants reported knee problems in the previous week using the OSTRC questionnaire. Fifteen (26.3%) participants also experienced pain during the SLDS; 22.8% (right leg), 17.5% (left leg) and 14.0% (bilaterally). The location of pain for right and left legs is illustrated in Fig. 2. Pain was described as being over the patellar tendon (PTP) in 7 (12.3%) of right legs and 2 (4%) of left legs.

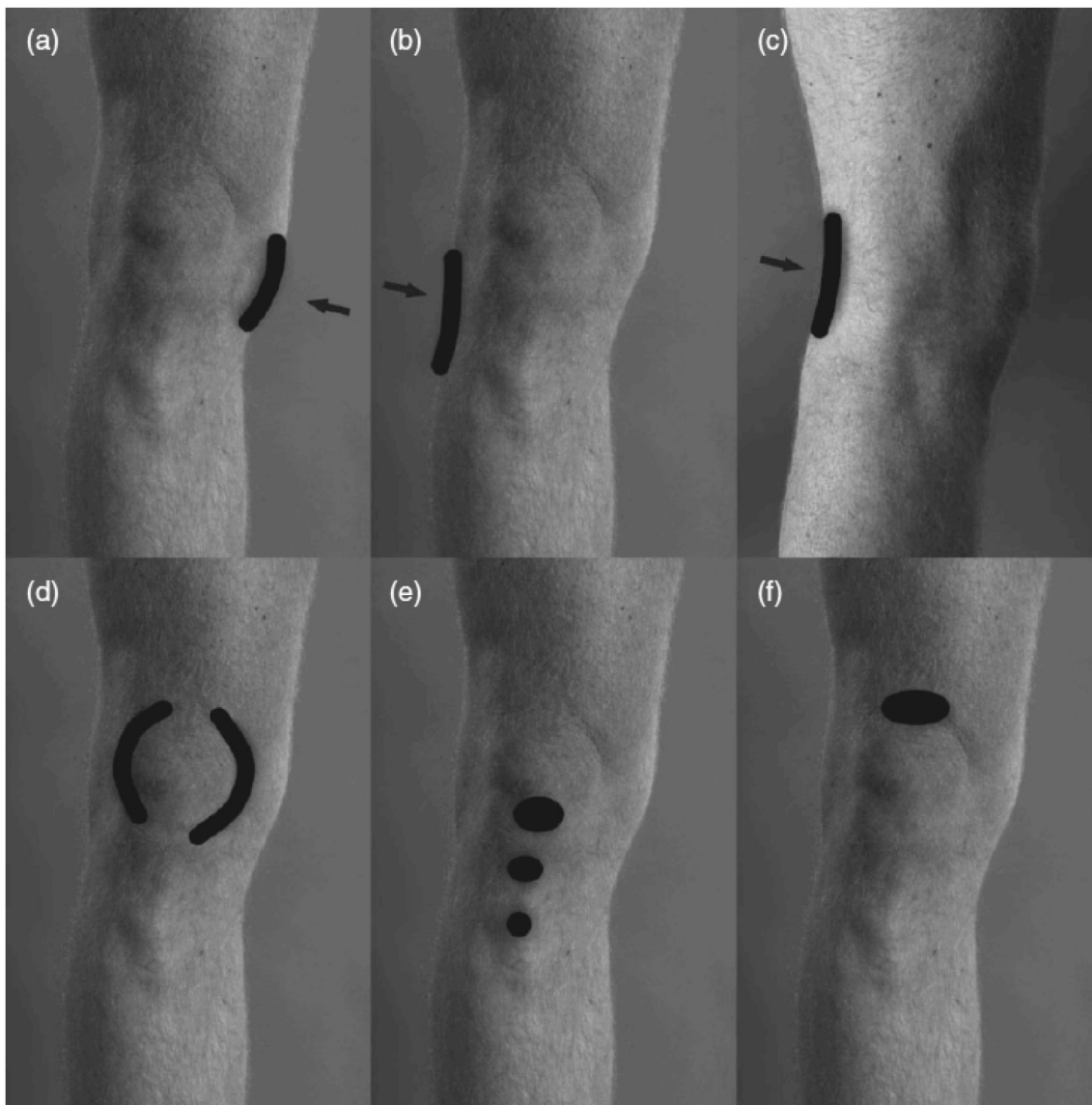


Fig. 1. Pain map used by participants to select the picture that best matches the location of knee pain experienced during or immediately after the squat. Adapted from van der Worp et al. (van der Worp et al., 2012).

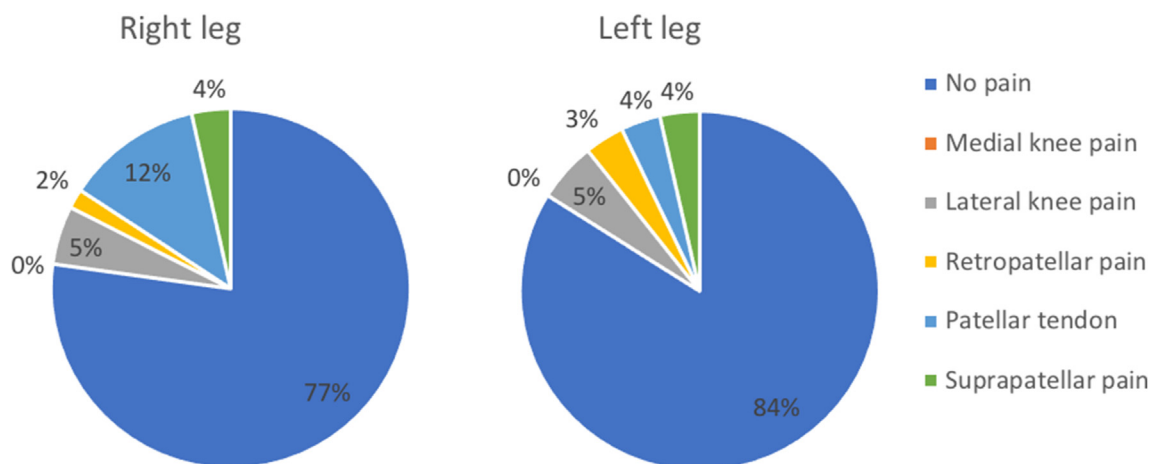


Fig. 2. Frequency of pain in each region of the pain map for the right and left legs. No participants reported posterior knee pain.

Table 1
 Ultrasonography, demographic and training variables and symptom questionnaires for participants with no-pain, localised patellar tendon pain (PTP) or other knee pain (OKP) areas during the single leg decline squat test of the right limb.

	No-pain	OKP	PTP	Sig.
n (%)	44 (77.2%)	6 (10.5%)	7 (12.3%)	
<i>Ultrasonography (Right leg)</i>				
Thickness mid-tendon (cm) [^]	0.33 ± 0.04	0.31 ± 0.02	0.41 ± 0.06	<0.001*
Thickness proximal tendon (cm) [^]	0.47 ± 0.07	0.41 ± 0.04	0.50 ± 0.08	0.053
CSA mid-tendon (cm ²) [^]	0.86 ± 0.14	0.75 ± 0.14	1.07 ± 0.14	<0.001*
Hypo-echogenic areas n (%)~	13 (29.6%)	3 (50%)	5 (71.4%)	0.080
Neovascularisation n (%)~	4 (9.1%)	1 (16.7%)	4 (57.1%)	0.005*
<i>Training variables</i>				
Right lead leg during spike jump	13 (29.6%)	3 (50%)	3 (42.9%)	0.517
Years played volleyball [^]	3.1 ± 1.1	2.0 ± 1.5	2.9 ± 1.3	0.096
Strength training (hrs)#	1 [0,2]	1.5 [1,2]	1 [0,3]	0.923
Athletic training (hrs)#	1.5 [0.5,2.5]	1 [1,1.5]	2 [0,4]	0.738
Ball practice training (hrs)#	2 [1,3.5]	3 [1,7]	4 [0,7]	0.669
<i>Demographic variables</i>				
Male n (%)~	24 (54.6%)	3 (50%)	5 (71.4%)	0.670
Age (years) [^]	15.1 ± 0.9	15.0 ± 0.6	15.3 ± 0.8	0.827
Weight (kg) [^]	64.5 ± 13.9	66.3 ± 15.0	71.7 ± 22.8	0.512
Height (cm) [^]	174.9 ± 9.9	174.1 ± 10.9	176.5 ± 9.8	0.898
<i>Questionnaires</i>				
OSTRC (0–100)#	0 [0,0]	4 [0,16]	28 [0,41]	0.020*
VISA-P (0–100)#	96 [87,100]	77 [76,90]	74 [56,94]	0.013*

CSA: cross-sectional area; OSTRC: Oslo Sports Trauma Research Center; VISA-P: Victorian Institute of Sport Australia-Patellar tendon. Group differences evaluated by analysis of variance[^], Chi-squared 2- or Kruskal-Wallis# with significance indicated by asterisk.

Table 2
 Ultrasonography data for participants with no-pain, localised patellar tendon pain (PTP) or other knee pain (OKP) areas during the single leg decline squat test of the left limb. Results were not analysed due to insufficient cases.

	No-pain	OKP	PTP
n (%)	47 (82.5%)	8 (14.0%)	2 (3.5%)
<i>Ultrasonography</i>			
Thickness mid-tendon (cm)	0.32 ± 0.04	0.32 ± 0.04	0.34 ± 0.005
Thickness proximal tendon (cm)	0.45 ± 0.06	0.46 ± 0.10	0.45 ± 0.03
CSA mid-tendon (cm ²)	0.85 ± 0.16	0.89 ± 0.20	0.82 ± 0.12
Hypo-echogenic areas n (%)	12 (25.5%)	4 (50%)	0 (0%)
Neovascularisation n (%)	6 (12.8%)	2 (25.0%)	0 (0%)

The following group differences are based on SLDS responses for the right leg (Table 1). Ultrasound data for the left leg is provided for reference in Table 2. Comparison of No-pain, PTP and OKP groups demonstrated no significant differences in demographic, anthropometric or training variables. Average years of volleyball play was 3.0 (range 1–6) and the cohort reported a median 2 hrs ball practice in the previous week. A similar proportion of preferred lead legs was observed between groups, while 95% of the cohort was right hand dominant.

Patellar tendon thickness and CSA of the mid-tendon was significantly different between groups ($p < 0.001$) (Fig. 3). Thickness and CSA was significantly ($p < 0.001$) greater for tendons with associated PTP compared to no-pain (Hedges' g : 1.86 (0.96–2.71) and 1.48 (0.63–2.32) respectively) and significantly ($p < 0.002$) greater to OKP groups (2.01 (0.67–3.34) and 2.13 (0.76, 3.49) respectively). No differences ($p > 0.28$) were found between OKP and No-pain groups. Differences in thickness at the proximal patellar tendon did not reach significance ($p = 0.053$). Prevalence of abnormal tendon structure ranged between 29.6% and 71.4%, but was not statistically different between groups ($p = 0.08$). Neovascularisation was significantly different between groups ($p = 0.005$). A higher prevalence of neovascularisation was observed for tendons with associated PTP (57.1%), compared to both no-pain (9.1%, odds ratio (95% CI): 13.3 (2.2, 81.9)) and compared to OKP (16.7%; odds ratio: 6.7 (0.5, 91.3)), which were not different (Table 1). Neovascularisation was observed in male participants only.

VISA-P and OSTRC scores were significantly different between groups ($p = 0.004$ and 0.036 respectively), with more severe symptoms for participants reporting PTP (Table 1).

4. Discussion

Approximately 1 in 4 non-elite volleyball athletes aged 13–17 years reported knee symptoms in the previous week and a similar proportion experienced pain during a SLDS. While these estimates are lower than those of a recent study of elite, adult basketball athletes, where 1 in 2 experienced a positive SLDS test (Hannington et al., 2020), they suggest that knee pain is a common problem in volleyball players of all ages and training background. When the right leg was considered, local pain over the patellar tendon was the most common area when symptoms during the test were reported (12.3%). VISA-P and OSTRC scores and some ultrasonography variables were also different between participants with local PTP compared to participants with other areas of pain.

The results demonstrated significantly greater tendon dimensions at the mid-tendon in the PTP group compared to both OKP and no-pain groups. As no differences in demographic, anthropometric or training variables were found between groups, we are confident that differences in tendon dimensions are not confounded by maturation or recent mechanical loading. Body mass and muscle mass increase substantially from childhood to adulthood, requiring the weight-bearing tendons to tolerate higher loads (Vaughn et al., 2012). This maturational process includes dimensional growth as well as changes in the tendon's intrinsic material properties, such as by increased collagen density. Similar changes are proposed to occur following resistance training in adults, although increases in tendon thickness were not observed in adolescent athletes despite high training volume over a 1.7 year period (Visnes et al., 2015). Contrary to other studies in elite athletes (Cassel et al., 2015; Lian et al., 2005), we did not find a higher prevalence of patellar tendinopathy in males, despite tendon thickness being greater in males than females (mid-tendon 0.35 ± 0.05 and 0.31 ± 0.04 cm respectively, $p < 0.001$).

A systematic review of tendinopathy in children and adolescents found studies using clinical examination typically reported

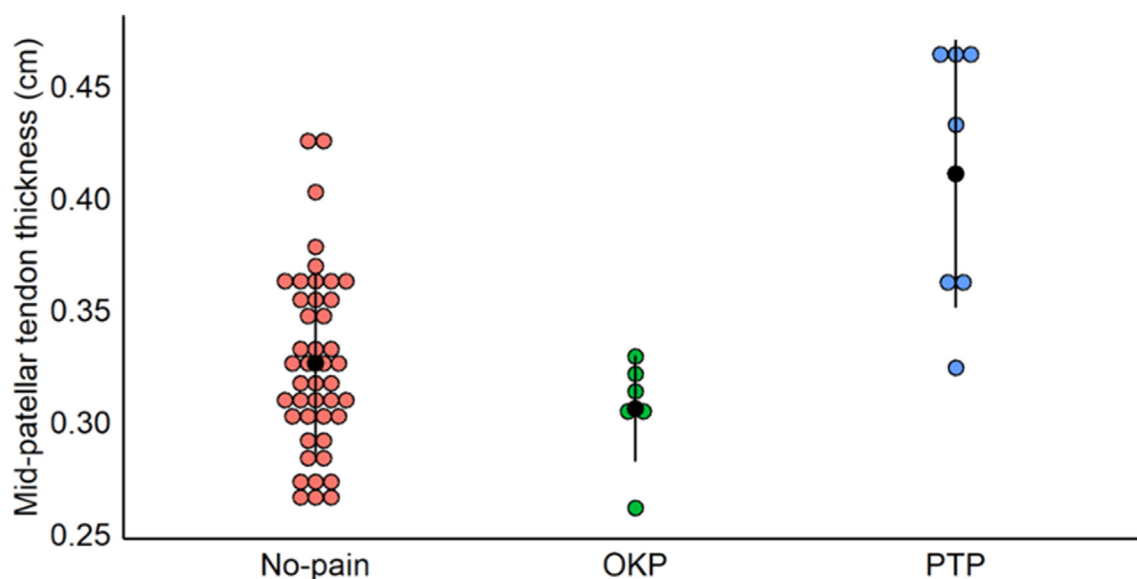


Fig. 3. Mid-tendon thickness of the patellar tendon for participants with no-pain, localised patellar tendon pain (PTP) or other knee pain (OKP) areas during the single leg decline squat test of the right limb. Black circles represent mean and black line represents standard deviation.

lower prevalence rates (2–18%) of tendinopathy than studies using ultrasonography (20–38%) (Simpson et al., 2016). This is consistent with findings that asymptomatic pathology (e.g. presence of hypoechoic areas or neovascularisation) exists in a large proportion of the active adult population (Gisslen et al., 2005). In our cohort, the proportion of participants with structural tendon abnormality was not statistically different between No-pain, PTP and OKP groups. In contrast, neovascularisation was significantly more likely in participants with PTP than both other groups. These findings suggest assessment using Doppler ultrasound examination provides additional information to grey-scale imaging alone when evaluating the adolescent patellar tendon. As previous studies demonstrate intratendinous flow can increase following exercise (Cook et al., 2005), a lack of standardisation of load immediately prior to imaging may have influenced findings.

As patellar tendinopathy begins in childhood and increases in prevalence with age up to 18 years (Simpson et al., 2016), and is difficult to treat, early screening to identify at-risk individuals is needed. The current results support the use of the SLDS test in evaluating patellar tendinopathy when information about the location of symptoms during testing is taken into account. This study is not without limitations. First, the small sample of symptomatic tendons, particularly on the left side will affect robustness of these conclusions. For this reason, evaluation of the diagnostic accuracy of this assessment to identify PTP was not conducted. Second, our physical examination of participants was limited to performance of the SLDS test and not a full clinical examination. Other clinical information is likely to increase the likelihood of a diagnosis of patellar tendinopathy and assist in identification of modifiable factors (Mendonca et al., 2018). A previous study of adult athletes indicated improved diagnostic accuracy for diagnosing patellar tendinopathy when multiple tests were used, compared to that of isolated tests (Mendonca Lde et al., 2016).

5. Conclusion

In a cohort of non-elite volleyball players aged 13–17 years, individuals reporting localised pain over the patellar tendon during the SLDS test were different from participants without pain or other areas of knee pain based on patellar tendon dimensions,

neovascularisation and VISA-P scores. Use of a pain map is proposed to be a critical component of the test. These results support the use of the SLDS test for early screening during the adolescent years, when symptoms of patellar tendinopathy may arise.

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Ethical statement

The study was approved by Griffith University ethics committee (Approval # 2017/896). Participants (and legal guardians) signed informed consent prior to testing and the rights of the participants were protected.

Declaration of competing interest

None.

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