# Title:

A systematic review and meta-analysis of the reliability and validity of sensorimotor measurement instruments in people with chronic low back pain

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#### Abstract

**Background:** Deficits in the sensorimotor system and its peripheral and central processing of the affected body part might be a contributing factor to chronic low back pain (CLBP). Hence, sensorimotor assessment is important. Valid and reliable sensorimotor measurement instruments are needed.

**Objective:** To investigate the reliability and validity of sensorimotor measurement instruments for people with chronic low back pain (CLBP).

**Design:** Systematic review and meta-analysis.

**Methods:** The review was undertaken using the COSMIN guidelines. Databases were searched for studies investigating the clinimetric properties of sensorimotor tests in people with CLBP. The methodological study quality was rated by two independent reviewers using the COSMIN 4-point rating checklist.

**Results:** Ten studies were included covering six sensorimotor measurement instruments with findings for reliability/measurement error, known-groups validity and convergent validity. The methodological quality ranged from poor to good, with only one study rated as good. There was insufficient evidence of enough quality to assess reliability/measurement error or convergent validity. Two-point discrimination, laterality judgement and movement control tests had moderate evidence supporting their ability to distinguish between healthy people and those with CLBP.

**Conclusions:** Two-point discrimination, laterality judgment and movement control tests demonstrate the greatest level of known-groups validity for people with CLBP. However, as the reliability of these measurement tools have yet to be established, this validity data should be interpreted cautiously. Further research is warranted to investigate the clinimetric properties of these sensorimotor techniques.

# Keywords

Chronic low back pain, sensorimotor test, systematic review, meta-analysis

#### Introduction

Chronic low back pain (CLBP) is a major public health problem, with a lifetime prevalence of ~84% (Denteneer et al., 2016, Murray et al., 2013). It is a leading cause of disability worldwide (Murray et al., 2013). Many factors contribute to the development and/or maintenance of CLBP (Denteneer et al., 2016). It has been proposed that deficits in the sensorimotor system (sensorimotor dysfunction) could be a contributing factor (Apkarian et al., 2011, Catley et al., 2014, Moseley and Flor, 2012). As such, there is growing interest in outcome measures and interventions that attempt to measure and improve sensorimotor function in people with CLBP (Ehrenbrusthoff et al., 2016, Elgueta-Cancino et al., 2015, Louw et al., 2015, Louw et al., 2016, Villafane et al., 2015, Vuilleumier et al., 2015).

Sensorimotor function encompasses all sensory and motor elements necessary for an individual to interact with their environment (Shumway-Cook and Woollacott, 2007). This includes the output from the nervous system contributing to motor function and any sensory input contributing to the interpretation of body position and movement (Hodges and Falla, 2015). A range of sensorimotor measurement instruments (SMIs) exist that attempt to measure the construct of sensorimotor dysfunction, defined as a process of altered motor behavior, and/or distorted interpretation or inaccurate input of afferent sensory information (Hodges and Falla, 2015, Pelletier et al., 2015). Some SMIs require expensive specialist equipment and highly skilled technical staff, such as functional magnetic resonance imaging (fMRI). Such techniques are beyond the capacity of routine clinical practice. Thus, there is a need for simple SMIs that are clinically practicable, to facilitate sensorimotor assessment and intervention.

There are a number of clinically practicable SMIs, such as two-point discrimination (TPD), laterality judgement and movement control tests (MCTs) (Catley et al., 2013, Luomajoki, 2012, Moseley, 2006). An essential prerequisite for any clinical test is that it demonstrates sound clinimetric properties (De Vet et al., 2011), particularly, reliability and validity (Atkinson and Nevill, 1998, De Vet et al., 2011). The clinimetric properties of some SMIs have been investigated in healthy people and an array of patient groups (Auld et al., 2011, Stanton et al., 2013, Wand et al., 2014a). The clinimetric properties of some of these SMIs have been explored in people with CLBP but the extent and the quality of the work has not been systematically reviewed. Such a review is needed to guide research and clinical practice in the field. Thus, the aim of this study was to systematically investigate the reliability and validity of simple SMIs in people with CLBP.

#### Methods

The search strategy was developed in accordance with COSMIN recommendations (Terwee et al., 2011) and the PRISMA guidelines (Moher et al., 2010). This systematic review is registered on PROSPERO (Registration number: CRD42015026880).

Structured search strategies were designed using search terms appropriate for each database. Standardised database subject headings such as MeSH terms (in MEDLINE) and Subject Headings (in CINAHL) were used in each database, as appropriate. For the MEDLINE search, the sensitive PubMed search filter proposed by COSMIN for measurement properties was used (Terwee et al., 2009). Search

terms and synonyms were searched separately in four main categories and finally combined into one search string per database. The categories complied with COSMIN guidelines (Terwee et al., 2009) and were defined as:

- 1. Construct: tactile acuity OR sensorimotor dysfunction OR cortical reorganization
- 2. Target population: chronic low back pain
- 3. Measurement instrument: sensorimotor test
- Measurement properties: sensitive COSMIN search filter for measurement properties for in MEDLINE

Electronic searches of databases were conducted by one author (K.E.) until March 30<sup>th</sup> 2015 using MEDLINE via PubMed, CINAHL via EBSCO, Embase via Ovid and Central via Wiley. The search was updated with a time restriction from March 30<sup>th</sup> 2015 to April 30<sup>th</sup> 2016 to identify relevant studies published ad interim. A full description of the search strategies can be found in the supplementary data (Appendix 1: Search strategies for all databases). Identified records were screened by K.E. by title-abstract initially and then by full-text screening. Hand searching of key reference lists was also conducted.

#### **Eligibility Criteria**

Studies were included if: 1) their target population were individuals with CLBP, defined as pain between the 12<sup>th</sup> rib and the buttock creases, persisting for 3 months (Savigny et al., 2009), 2) the SMI investigated claimed to measure a component of sensorimotor dysfunction, 3) the SMI investigated was practicable without sophisticated/expensive instrumentation (e.g. an functional Magnetic Resonance Imaging (fMRI) machine) not easily accessible in a routine clinical setting. An example of an unsophisticated and inexpensive piece of equipment would be a goniometer, 4) the aim was to investigate one or more measurement properties of the SMI under investigation, 5) they were designed to investigate reliability or validity of the SMI, in accordance with the COSMIN taxonomy (Mokkink et al., 2009), 6) the study was published as a full original article in English or German.

Studies were excluded if: 1) they were of an intervention based or single-case design, 2) the SMI investigated required extensive technical skills and/or equipment not found in routine clinical practice (e.g. fMRI, motion analysis systems).

#### **Data Extraction**

According to the COSMIN recommendations for data extraction, the generalisability box of the COSMIN tool was used to extract data on characteristics of the study sample (median/mean age, distribution of sex, important disease characteristics, setting, country, language, sampling strategy, percentage of missing responses). In addition, details of each SMI data collection protocol were summarised and the measurement property results per SMI were extracted separately (De Vet et al. 2011). The extraction process was carried out by the lead author (K.E).

### Methodological Quality Evaluation

The COSMIN four-point scoring checklist (Terwee et al., 2012) was used to assess the methodological quality of included studies. The checklist is a validated tool comprising 10 sections, each assessing a separate measurement property (Mokkink et al., 2010a, Mokkink et al., 2010b).Two reviewers (C.R. and K.E.) with prior experience in using the checklist rated each study. Each item for methodological quality within each section was scored from excellent to poor. The overall score for the measurement property within the study was defined as the lowest rating among all response options within one section, termed as "worst score counts" (Terwee et al., 2012). Where multiple measurement properties were assessed within one study, this study received multiple methodological quality evaluations.

#### **Evaluation of measurement properties**

In the studies included in the review, the results for each SMI measurement property were evaluated against the pre-defined quality for good measurement properties (Terwee et al., 2007), (see table 1 for details). For validity, we investigated the construct validity sub-categories known-groups validity and convergent validity. Known groups validity was defined as an instrument's ability to discriminate between people with and without the target condition or between people having different manifestations of the target condition, respectively (De Vet et al., 2011). Convergent validity was defined as the expected relationship between instruments measuring related constructs (De Vet et al., 2011).

Table 1: Quality criteria for measurement properties

Property	Rating	Quality Criteria
Reliability		
Internal	+	Cronbach's alpha(s) ≥ 0.70
consistency	?	Cronbach's alpha not determined or dimensionally
		unknown
	-	Cronbach's alpha(s) < 0.70
Reliability	+	ICC / weighted Kappa $\ge 0.70$ OR Pearson's r $\ge 0.80$
	?	Neither ICC / weighted Kappa, nor Pearson's r
		determined
	-	ICC / weighted Kappa < 0.70 OR Pearson's r < 0.80
Measurement error	+	MIC > SDC OR MIC outside the LOA
	?	MIC not defined
	-	MIC ≤ SDC OR MIC equals or inside LOA
Validity		
Content validity		All items are considered to be relevant for the
	+	construct to be measured, for the target population,
		and for the purpose of the measurement AND the
		questionnaire is considered to be comprehensive
	?	Not enough information available
		Not all items are considered to be relevant for the
	-	construct to be measured, for the target population,
		and for the purpose of the measurement OR the

		questionnaire is considered not to be
		comprehensive
Construct validity	+	Factors should explain at least 50% of the variance
- Structural validity	?	Explained variance not mentioned
	-	Factors explain < 50% of the variance
- Hypothesis		Correlations with instruments measuring the same
testing	+	construct $\ge$ 0.50 OR at least 75% of the results are
		in accordance with the hypotheses AND correlations
		with related constructs are higher than with
		unrelated constructs
	?	Solely correlations determined with unrelated
		constructs
		Correlations with instruments measuring the same
	-	construct < 0.50 OR < 75% of the results are in
		accordance with the hypotheses AND correlations
		with related constructs are lower than with unrelated
		constructs
Cross-cultural	+	No differences in factor structure OR no important
validity		DIF between language versions
	?	Multiple group factor analysis not applied AND DIF
		not assessed
	-	Differences in factor structure OR important DIF
		between language versions
Criterion validity	+	Convincing arguments that gold standard is "gold"
		AND correlation with gold standard $\ge 0.70$

	?	No convincing arguments that gold standard is
		"gold"
	-	Correlation with gold standard < 0.70
Responsiveness		
		Correlation with changes on instruments measuring
	+	the same construct $\geq$ 0.50 OR at least 75% of the
		results are in accordance with the hypotheses OR
		AUC $\geq$ 0.70 AND correlations with changes in
		related constructs are higher than with unrelated
		constructs
	?	Solely correlations determined with unrelated
		constructs
		Correlation with changes on instruments measuring
	-	the same construct < 0.50 OR < 75% of the results
		are in accordance with the hypotheses OR AUC <
		0.70 AND correlations with changes in related
		constructs are lower than with unrelated constructs
Legend: MIC = minim	nal importa	ant change, SDC = smallest detectable change,
LOA = limits of agree	ment, ICC	C = intraclass correlation coefficient, DIF = differential
item functioning, AUC	C = area u	inder the curve, + = positive rating,? = indeterminate
rating, - = negative ra	iting	
Table taken from CC	)SMIN gui	idelines (Terwee et al., 2011) <sup>1</sup> .

<sup>&</sup>lt;sup>1</sup> Reprinted from the Journal of Clinical Epidemiology 2007;, Terwee CB, Bot SDM, de Boer MR, van der Windt DAWM, Knol DL, Dekker J, Bouter LM, de Vet HCW. Quality criteria were proposed for measurement properties of health status questionnaires, 60:34-42., Copyright (2007), with permission from Elsevier

### Data synthesis: meta-analysis and best evidence synthesis

Where multiple studies with comparable study designs investigated the same SMI and measurement property, a meta-analysis was conducted. For known-groups validity, mean scores and standard deviations from healthy and patient groups were pooled using the statistical package RevMan (Version 5) by means of forest plots (fixed effects model) to establish a pooled difference between groups. Heterogeneity was quantified using the I<sup>2</sup> (Higgins et al., 2003). Following the COSMIN recommendations, studies with a poor methodological score were excluded from quantitative pooling (Mokkink et al., 2009). Where quantitative pooling was not appropriate, a 'best evidence synthesis" approach was used, (see Table 2) (Guyatt et al., 2011, Schünemann et al., 2011).

Table 2: Level of Evidence for the quality of the measurement property

Level	Rating*	Criteria
strong	+++ or	Consistent findings in multiple
		studies of good
		methodological quality OR in one
		study of excellent
		methodological quality
moderate	++ or	Consistent findings in multiple
		studies of fair
		methodological quality OR in one
		study of good
		methodological quality
limited	+ or -	One study of fair methodological
		quality
conflicting	+/-	Conflicting findings
unknown	?	Only studies of poor
		methodological quality
Legend: * + = positive rating	g,? = indetermina	ate rating, - = negative rating
Table taken from COSMIN	guidelines (Terw	ee et al., 2011) <sup>2</sup>

<sup>&</sup>lt;sup>2</sup> Reprinted from the Journal of Clinical Epidemiology 2007; Terwee CB, Bot SDM, de Boer MR, van der Windt DAWM, Knol DL, Dekker J, Bouter LM, de Vet HCW. Quality criteria were proposed for measurement properties of health status questionnaires, 60:34-42., Copyright (2007), with permission from Elsevier

### Results

#### **Study Selection**

Initially, 4,285 studies were identified, of which 407 were excluded as duplicates and another 3,839 were excluded following title and abstract screening. Fifty studies were included for full-text assessment from which nine studies evaluating six SMIs were included. In the updated search, 686 studies were initially identified, however, only one additional relevant study was included in the final study list. Thus, in total, 10 studies (Table 3) evaluating six SMIs were included (Figure 1), within which three measurement properties were investigated: reliability/measurement error, known-groups validity, and convergent validity. Details for the data collection protocols for each study are summarised in supplementary data (Appendix 2: Individual study data collection protocols).

The findings for each measurement property per SMI from the individual studies are quantified in supplementary data (Appendices 4-8).



Figure 1: Flow chart of literature search and study selection process.

		Patient Characteristics		
Author (Year)	a) Instrument b) Measurement	a) Mean Age (SD) (years)	a) Pain Severity Mean (SD)	a) Setting b) Country
	Property	b) Distribution of Sex	b) Disability Mean (SD)	c) Language
	c) n			d) Sampling
				e) %of missing responses
Linder et al (2015)	a) Laterality Judgement	a) CLBP: 44.9 (11.0)	a) VAS Scores:	a)PT clinics
	b) Known-Groups	HC: 43.3 (9.6)	55.3 (17.8)	b)Sweden
	Validity	b) CLBP: $\bigcirc = 20$	b) ODI Scores:	c)Swedish
	c) CLBP: n=30	් = 10 ර = 10	25.1 (13.1)	d)CLBP: consecutive
	HC: n=30	HC: $\begin{array}{c} \varphi \\ \varphi \end{array} = 20 \\ 10 \end{array}$		HC: convenience
		d' = 10		e) n=1 in CLBP group
Nishigami et al	a) TPD	a) CLBP normal BI:	a) VAS Scores normal BI:	a) Orthopedic clinic
(2015)	BID	65.1 (11.2)	48.3 (21.8)	b) Japan
	b) Known-Groups	CLBP expanded BI:	VAS Scores expanded	c) Japanese
	Validity	56.7 (16.7)	BI:	d) Not stated
	c) CLBP: n=42	CLBP shrink BI:	42.5 (24.5)	e) Not stated
	HC: n=17	62.0 (12.4)	VAS Scores shrink BI:	
		HC:	42.0 (23.5)	
		63.4 ± 12.2	b) RMDQ Scores normal	
		c) CLBP: 우 = 26	BI:	
		∂ <sup>^</sup> = 16	7.0 (2.4)	
		HC: $\bigcirc = 8$	RMDQ Scores	
		⊖ <sup>′</sup> = 9	expanded BI:	
			6.2 (3.4)	
			RMDQ shrink BI:	
			6.8 (4.4)	

		Patient Characteristics		
Author (Year)	a) Instrument b) Measurement Property c) n	a) Mean Age (SD) (years) b) Distribution of Sex	a) Pain Severity Mean (SD) b) Disability Mean (SD)	a) Setting b) Country c) Language d) Sampling e) %of missing responses
Wand et al (2014)	<ul> <li>a) FreBAQ</li> <li>b) Known-Groups Validity</li> <li>c) CLBP: n=51 HC: n=51</li> </ul>	<ul> <li>a) CLBP: 41.7 (14.0) HC: 38.7 (13.4)</li> <li>b) CLBP: ♀ = 21 ♂ = 30 HC: ♀ = 20 ♂ = 31</li> </ul>	<ul> <li>a) NRS Scores (0-100): 48.2 (17.8)</li> <li>b) RMDQ Scores: 10.1 (5.9)</li> </ul>	<ul> <li>a) Community PT practice; Department of pain management , The Sir Charles Gairdner Hospital, Perth, Western Australia</li> <li>b) English</li> <li>c) CLBP: convenience HC: convenience</li> <li>d) Not stated</li> </ul>
Bowering et al	a) Laterality Judgement	a) Complete sample:	a) Not stated	a) Online study
(2014)	b) Known-Groups	37 (13)	b) Not stated	b) Australia
	Validity	b) Complete sample:		c) English
	n=117 History of back pain: n= 462 HC: n= 429	♀ = 684; ♂ = 324		e) 181 datasets excluded
Stanton et al	a) TPD	a) CLBP: 45 (14)	a) Not stated	a) Royal North Shore
(2013)	b) Known Groups-		b) Physical component of	Hospital, Sydney
	Validity Convergent Validity	D) CLBP: $\Upsilon = 14$	SF-36 (U-30): 10 7 (7 4)	D) AUSTRAIIA
	c) CL RP: n=17	⊖ − 3 HC: ○ = 11	19.7 (7.4)	d) CI BP data from Bray
	HC: n=18	∂ = 7		and Moseley (2011)

		Patient Characteristics		
Author (Year)	a) Instrument b) Measurement Property c) n	a) Mean Age (SD) (years) b) Distribution of Sex	a) Pain Severity Mean (SD) b) Disability Mean (SD)	a) Setting b) Country c) Language d) Sampling
	-			e) %of missing responses
				HC: convenience e) Not stated
Bray and Moseley (2011)	<ul> <li>a) Laterality Judgement</li> <li>b) Known-Groups Validity</li> <li>c) CLBP: n=21 HC: n=14</li> </ul>	<ul> <li>a) CLBP: 44 (13) HC: 43 (7)</li> <li>b) CLBP: ♀ = 15; ♂ = 6 HC: ♀ = 9; ♂ = 5</li> </ul>	a) VAS Scores (0-100): 37 (21) b) Not stated	<ul> <li>a) Private PT practice</li> <li>b) United Kingdom (ethical approval)</li> <li>c) English</li> <li>d) Convenience sample</li> <li>e) Not stated</li> </ul>
Luomajoki and Moseley (2011)	<ul> <li>a) TPD Movement Control Tests</li> <li>b) Known-Groups Validity</li> <li>c) CLBP: n=45 HC: n=45</li> </ul>	a) CLBP: 43 (15) HC: 41 (10) b) CLBP: ♀ = 25; ♂ = 20 HC: ♀ = 25; ♂ = 20	a) Not stated b) RMDQ Scores: 9 (5)	<ul> <li>a) Private PT practice</li> <li>b) Switzerland</li> <li>c) German</li> <li>d) Convenience</li> <li>e) Not stated</li> </ul>
Wand et al (2010)	<ul> <li>a) TPD Graphesthesia</li> <li>b) Known-Groups Validity</li> <li>c) CLBP: n=19 HC: n=19</li> </ul>	a) CLBP: 41 (12.5) HC: 34 (12.1) b) CLBP: ♀ = 11 ♂ = 8 HC: ♀ = 14 ♂ = 5	<ul> <li>a) NRS Scores (0-10) usual pain: 3.9 (2.1)</li> <li>b) Physical component of SF-36 (0-30): 21.8 (5.0)</li> </ul>	<ul> <li>a) District General hospital, Perth, Western Australia</li> <li>b) Australia</li> <li>c) English</li> <li>d) CLBP: convenience HC: convenience</li> <li>e) 2 subjects with ambiguous pain scores and missing scores respectively, treated as missing values</li> </ul>

		Patient Characteristics		
Author (Year)	a) Instrument	a) Mean Age (SD)	a) Pain Severity Mean	a) Setting
	b) Measurement	(years)	(SD)	b) Country
	Property	b) Distribution of Sex	b) Disability Mean (SD)	c) Language
	c) n			d) Sampling
				e) %of missing responses
Moseley (2008)	a) TPD	a) CLBP: 43.83 (11.12)	a) VAS Scores:	a) Not stated
	BID	HC: not stated	47.2 (12.54)	b) UK
	b) Known-Groups	b) CLBP: ♀ = 3	b) Not stated	c) English
	Validity	ె = 3		d) Consecutive
	c) CLBP: n= 6	HC: ♀ = 5		e) Not stated
	HC: n=10	্ৰ = 5		
Luomajoki (2008)(	a) Movement Control	a) LBP: 41 (15)	a) Not stated	a) Outpatient PT clinics,
	Tests	HC: 37 (12)	b) RMDQ Scores:	Canton Aargau
	b) Known-Groups	b) LBP: ♀ = 72	8(5)	b) Switzerland
	Validity	ే = 36		c) German
	c) LBP: n=108	HC: ♀ = 58		d) CLBP: consecutive
	HC: n= 102	্ৰ = 44		HC: convenience
				e) Not stated
Separate Patient Ch	paracteristics of Studies inve	estigating Reliability (Subsa	mples of CLBP patients)	
Wand et al (2014)	a) FreBAQ	a) 42 (14)	a) Back Pain Intensity (0-	a) Community PT practice
	b) Reliability	b) ♀ = 12; ♂ = 14	100):	b) Australia
	c) CLBP: n= 26		47.7 (14.4)	c) English
			b) RMDQ Scores:	d) Not stated
			10.6 (6.0)	<ul> <li>e) N=1 did not return</li> </ul>
				second questionnaire;
				handled as missing item
Bray and Moseley	a) Laterality Judgement	a) 46 (16)	a) VAS Scores:	a) Private PT practice
(2011)	b) Reliability	b)♀ = 1; ♂ = 4	46 (23)	b) UK
	c) CLBP: n= 5		b) ODI Scores:	c) English
			25.1 (13.1)	d) Convenience
				e) Not stated

		Patient Characteristics		
Author (Year)	a) Instrument b) Measurement Property c) n	a) Mean Age (SD) (years) b) Distribution of Sex	a) Pain Severity Mean (SD) b) Disability Mean (SD)	a) Setting b) Country c) Language d) Sampling e) %of missing responses
Linder et al (2015)	a) Laterality Judgement b) Measurement Error Reliability c) CLBP: n= 30	a) 44.9 (11) b) ♀ = 20; ♂ = 10	<ul> <li>a) VAS Scores: 55.3 (17.8)</li> <li>b) ODI Scores: 25.1 (13.1)</li> </ul>	<ul> <li>a) PT clinics</li> <li>b) Sweden</li> <li>c) Swedish</li> <li>d) Convenience</li> <li>e) Not stated</li> </ul>
Legend: CLBP= Ch Image Drawings; Fr 36-Item Short Form Pain Intensity (0-100	ronic Low Back Pain; HC= I eBAQ= Fremantle Back Aw Health Survey (SF-36); RM 0); Data are presented as M	Healthy Controls; ♂= male; /areness Questionnaire; OD IDQ = Roland Morris Disabi /lean (SD) unless otherwise	♀=female; TPD = Two-Point I = Oswestry Disability Score lity Questionnaire; NRS = Nu stated.	Discrimination; BID= Body ; PT= Physiotherapy; SF-36 = merical Rating scale; Back

# Methodological quality evaluation of the studies

Across all 10 included studies, four methodological quality evaluations concerning reliability and/or measurement error were undertaken and received a poor methodological quality rating. Sixteen methodological quality evaluations of known-groups or convergent validity were conducted with one rated as good, eleven as fair and four as poor (see Table 4 and 5<sup>3</sup>).

<sup>&</sup>lt;sup>3</sup> Reprinted from the Journal of Clinical Epidemiology, 63, Mokkink LB, Terwee CB, Patrick DL, Alonso J, Stratford PW, Knol DL, Bouter LM, de Vet HCW., International consensus on taxonomy, terminology, and definitions of measurement properties: results of the COSMIN study, 737-745., Copyright (2010), with permission from Elsevier

	Desig	n Requi	rements F	Reliability	,	Statistical Methods									
Instrument				_	ts				to	. d	i			5d?	
and		f missing	on of how	ncluded in	asuremeni	suo		n the	construct	ons similar ts? e.g. ty	tant flaws	elation ated?	s: Was	a calculate	eme
Measure-		itage o	scriptic	e size ir	vo mea	listrati	ated?	table i	on the terval	onditic	impor	scores ss corr	s/ I score	es: I kappa	es: ing sch
ment	score	e percen	ere a des	e sample	it least tv	he admir	e time st	atients s	e time in	he test c h measu	): here any	L: tinuous : intracla: ient (ICC)	2: notomou al/ordina	3: inal score weighted	t: inal score e weight
Property	overall	ltem 1: Was th	ltem 2: Was th	ltem 3: Was th	ltem 4: Were a	ltem 5: Were t	ltem 6: Was th	ltem 7: Were p	interim Item 8: Was th	ltem 9: Were t for bot	Were t	ltem 1. for con Was ar coeffici	for dich nomina	ltem 15 for ord Was a y	ltem 1 <sup>4</sup> for ord Was th
Laterality															
Judgement															
Reliability															
Linder et al	poor	good	fair	poor	excell.	excell.	excell	good	excell.	fair	fair	excell.	n.a.	n.a.	n.a.
(2015)															
Bray and	poor	good	good	poor	excell.	excell.	excell	good	excell.	good	fair	good	n.a.	n.a.	n.a.
Moseley															
(2011)															
FreBaQ							I	1							
Reliability															
Wand et al	poor	good	fair	poor	excell.	good	excell.	poor	excell.	fair	fair	excell.	n.a.	n.a.	n.a.
(2014)															

# Table 4: Methodological quality evaluation: Reliability and Measurement Error Studies

	Design Requirements Measurement Error         S												Sta	tisti	ical	Met	thods	;	 											
Instrument				(	c		nts										r	ype			rs in									
and		fmissing			i papulad		asuremei		suo		stated?		n the				ons simila	ts? e.g. t			tant flaw		or of	Smallest						
Measure-		entage o		iescriptic	ole size ir		two mea		ninistrati		interval		s stable i		interval		t conditio	suremen	400		ny impor	Ë	dard Errc	t (SEM), :						
ment	score	e nerci	_	ere a o	e samp		ıt least		he adn		e time		atient		e time		he test	h mea:	inictro.	ö	here a	l:.for C	e Stan	remen	able					
Property	overall	ltem 1: Was th	Item 2:	wastn	ltem 3: Was th	Item 4:	Were a	Item 5:	Were t	Item 6:	Was th	Item 7:	Were p	Item 8:	Was th	Item 9:	Were t	for bot	ملالم عم	ltem 10	Were t	ltem 11	Was th	Measui	Detecta					
Laterality																														
Judgement																														
Measure-																														
ment Error																														
Linder et al	poor	good	fair		poor	exce	ell.	exc	ell.	exce	ell.	good	b	exc	ell.	fai	r			fair		goo	d							
(2015)																														
FreBaQ= Fre	emantle	Back A	warene	ess C	Questior	naire	; CTT	T= Cla	assica	al Tes	t Tł	heory	/; exe	cell.	= exc	elle	nt; n	i.a.= r	not	арр	lica	ble								

	Desig	n Require	monts								Statistical
	Desig	in require	inento								Methods
Instrument and Measurement Property	overall score	Item 1: Percentage of missing items given?	Item 2: Description of how missing items were	Item 3: Adequate sample size included in the	Item 4: Formulation of hypotheses regarding	Item 5: Was the expected direction of correlations	Item 6: Was the expected absolute or relative	Item 7: For CV: was an adequate description of the	Item 8: For CV: Were the measurement properties of the	Item 9: Were there any important flaws in the	Item 10: Were design and statistical methods adequate for the hypotheses to be tested?
		1	1		1			1		1	Ι
Nishigami et al (2015)											
TPD measured as side-to-side differences	fair	good	fair	good	fair	good	good	n.a.	n.a.	fair	excell.
Stanton et al (2013)	fair	good	fair	fair	good	good	good	n.a	n.a.	fair	excell.
Luomajoki and Moseley (2011)	fair	good	fair	good	excell.	excell.	good	n.a.	n.a.	excell.	good
		guuu	iali	iali	excen.	CAUCH.	guuu	11.a.	11.a.	iaii	

# Table 5: Methodological quality evaluation: Known-Groups and Convergent Validity Studies

	Design Requirements													
Instrument and Measurement Property	overall score	ltem 1: Percentage of missing items given?	Item 2: Description of how missing items were	Item 3: Adequate sample size included in the	Item 4: Formulation of hypotheses regarding	Item 5: Was the expected direction of correlations or mean differences included in the	Was the expected absolute or relative	frem 7: For CV: was an adequate description of the	Item 8: For CV: Were the measurement properties of the	Item 9: Were there any important flaws in the	Item 10: Were design and statistical methods adequate for the hypotheses to be tested?			
Moseley (2008)	poor	good	good	poor	fair	good	good	n.a.	n.a.	fair	poor			
TPD CV														
Stanton et al (2013)	poor	good	fair	poor	excell.	excell.	good	excell.	fair	poor	excell.			
MCT KGV														
Luomajoki Moseley (2011)	fair	good	fair	good	excell.	excell. excell.		good n.a.		fair	good			
Luomajoki et al (2008)	fair	good	fair	excell.	good	ood good		good n.a.		fair	good			
Graphestesia KGV						1					I			
Wand et al (2010)	fair	good	fair	fair	excell.	excell.	good	n.a.	n.a.	fair	excell.			
Laterality Judgement KGV		1			1	1	1	1	1		1			
Linder et al (2015)	fair	good	fair	good	fair	good	good	n.a.	n.a.	excell. #	excell.			
Bowering et al (2014)	good	excell.	excell.	excell.	good	excell.	good	n.a.	n.a.	excell.	excell.			

	Design Requirements												
	Desig	n nequirei	liento								Methods		
Instrument and Measurement Property	overall score	Item 1: Percentage of missing items given?	Item 2: Description of how missing items were	Item 3: Adequate sample size included in the	Formulation of hypotheses regarding	Item 5: Was the expected direction of correlations	Item 6: Was the expected absolute or relative	ltem 7: For CV: was an adequate description of the	Item 8: For CV: Were the measurement properties of the	Item 9: Were there any important flaws in the	Item 10: Were design and statistical methods adequate for the hypotheses to be tested?		
Bray and Moseley ( 2011)	fair	good	fair	fair	excell.	excell.	good	n.a.	n.a.	fair	excell.		
Laterality Judgement CV		<u> </u>			1								
Stanton et al (2013)	poor	good	fair	poor	excell.	excell.	good	excell.	fair	poor	excell.		
BIDs KGV		1	I		1			I					
Nishigami et al (2015)	fair	good	fair	good	fair	Good	good	n.a.	n.a.	fair	excell.		
Moseley (2008)	poor	good	good	poor	fair	air good		n.a.	n.a.	fair	poor		
FreBaQ KGV		1	1	1		1	1						
Wand et al (2014)	fair	good	fair	excell.	fair	good	good	n.a.	n.a.	excell.	excell.		
TPD = Two-Point Discrin	TPD = Two-Point Discrimination; KGV= Known-Groups Validity; MCT= Movement Control Test; CV=Convergent Validity; BID= Body Image Drawings;												
FreBaQ= Fremantle Back Awareness Questionnaire; excell.= excellent; n.a.= not applicable													

## **Reviewer Agreement**

The inter-rater agreement for methodological quality between raters was good (Altman, 1991) (absolute agreement = 73%, Cohen's Kappa  $\kappa$  = 0.62 (95% CI 0.54, 0.70). Initial disagreement was resolved by consensus (see supplementary data Appendix 3: Reviewer Agreement).

## **Measurement properties**

## **Reliability and Measurement Error**

Studies were identified that investigated the reliability of laterality judgement and the FreBaQ.

# Laterality Judgement: reliability

Two studies (Bray and Moseley, 2011, Linder et al., 2015) (CLBP = 10 and CLBP=25/22), both of poor methodological quality, investigated the reliability of laterality judgement. Intraclass correlation coefficients (ICCs) for response time and accuracy were provided (see supplementary data: Appendix 5).

ICC values ranged from 0.51 to 0.91 for reaction time and from 0.69 to 0.92 for accuracy. Thus, the level of reliability could be considered acceptable for accuracy, but not for reaction time against the predefined acceptable level (ICC  $\geq$ 0.70).

As this body of evidence consisted of only poor quality studies, the evidence for the reliability of laterality judgement was classified as unknown.

Linder et al. (2015) investigated measurement error reporting the coefficients of variation (CV) for reaction time and accuracy of repeated measurements between time point one and two (CLBP= 25) and between time point two and three (CLBP= 22). For reaction time, the CV reduced from 19.6 % to 6.2 % whilst for accuracy it remained stable at 6.46 % to 6.77%. Data on minimally important change (MIC) were not provided. As this body of evidence consists of only one poor methodological quality study, the evidence for the measurement error of laterality judgement was classified as unknown.

#### Fremantle Back Awareness Questionnaire (FreBaQ): reliability

One study (Wand et al., 2014b), of poor methodological quality (n=26), investigated the test-retest reliability of the FreBaQ over a period of one week in people with CLBP (see supplementary data: Appendix 8). The test-retest performance was ICC<sub>2.1</sub> (95% CI) =0.652(0.307-0.848) (Agreement) and ICC<sub>2.1</sub> (95% CI)=0.667(0.317-0.857) (Consistency). This was below the COSMIN threshold of  $\geq$ 0.70.

However, as this body of evidence consists of only one poor quality study, the evidence for the test-retest reliability of the FreBaQ was classified as unknown.

#### **Known-Groups Validity**

Studies investigating the known-groups validity of TPD, graphaesthesia, laterality judgement, MCTs and the FreBaQ were found.

#### Two Point Discrimination (TPD): Known-Groups Validity

Five studies (CLBP=129, HC=91) investigated the known-groups validity of TPD were found. Four were of fair quality (Loumajoki and Moseley, 2011, Nishigami et al., 2015, Stanton et al., 2013, Wand et al., 2010) and one of poor methodological guality (Moseley, 2008) (see supplementary data: Appendix 4). Four of these studies were broadly consistent regarding their measurement protocol, in particular regarding the region of the lower back assessed and the horizontal TPD measurement approaches (Luomajoki and Moseley, 2011, Moseley, 2008, Stanton et al., 2013, Wand et al., 2010). The other study used side-to-side difference of TPD threshold as the outcome measure (Nishigami et al., 2015) and categorised patients according to their perceived body image. Of the four comparable studies, three identified a statistically wider TPD threshold for people with CLBP whilst the one study of poor methodological quality noted no difference between patients and healthy controls. When the three fair quality studies were statistically pooled people with CLBP demonstrated a statistically wider TPD threshold of 16 mm than healthy people (figure 2). No evidence of heterogeneity was found when the data was statistically pooled (Higgins and Green, 2011). Thus, there was moderate evidence from three studies of fair methodological quality that TPD possesses known-groups validity.

# Figure 2: Forest Plot comparing TPD regarding its known-groups validity to distinguish between people with CLBP and healthy controls

		clbp		healt	hy con	trol		Mean Difference		Mean Difference	Э	
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% CI		IV, Fixed, 95% 0		
Luomajoki Moseley	61	13	45	44	10	45	55.5%	17.00 [12.21, 21.79]		-		
Stanton	59.8	11.7	17	45.3	5.1	18	34.9%	14.50 [8.46, 20.54]				
Wand	62	21.6	19	44.2	13.7	19	9.6%	17.80 [6.30, 29.30]				
Total (95% CI)			81			82	100.0%	16.20 [12.64, 19.77]		•		
Heterogeneity: Chi <sup>2</sup> = 0.49, df = 2 (P = 0.78); l <sup>2</sup> = 0% Test for overall effect: Z = 8.90 (P < 0.00001)									-100 -50 Favo	0 urs [clbp] Favour	50 s (healthy co	100 ontrol]

Legend: IV = inverse variance; CI= confidence interval. Note: only data from horizontal TPD measurements were included in this analysis, leaving out values from vertical measurements from Luomajoki and Moseley (2011).

Nishigami et al. (2015) also reported a statistically significant difference between groups regarding the side-to-side differences of TPD thresholds (CLBP=42, HC=17). Thus, there is limited evidence from one study of fair methodological quality that TPD side-to-side difference possesses known-groups validity.

#### Laterality Judgment: Known Groups Validity

Three studies (Bowering et al., 2014, Bray and Moseley, 2011, Linder et al., 2015) (168=CLBP, 473=HC) assessed the known-groups validity of laterality judgement, two of fair (Bray and Moseley, 2011, Linder et al., 2015) and one (Bowering et al., 2014) of good methodological quality (see supplementary data: Appendix 5). Two studies (Bowering et al., 2014, Bray and Moseley, 2011) found a significant difference regarding laterality judgement accuracy, but only one concerning reaction time (Bowering et al., 2014), between people with CLBP and healthy controls, whilst one study found neither a statistical difference for reaction time nor accuracy (Linder et al., 2015). When the three studies were quantitatively pooled, people with CLBP were, on average, 9% less accurate and 0.1 seconds slower than healthy controls, both statistically significant. (figure 3a and 3b). The level of heterogeneity was substantial with I<sup>2</sup> values of 65% and 90% for reaction time and accuracy respectively, therefore, the findings should be interpreted cautiously (Higgins and Green, 2011). Hence, there was moderate evidence from one study of good and two

of fair methodological quality that demonstrated known-groups validity in laterality judgement accuracy and reaction time.

## Figure 3a: Forest Plot Laterality Judgement Reaction Time



## Figure 3b: Forest Plot Laterality Judgement Accuracy

		clbp		healt	thy cont	rol		Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% CI	IV, Fixed, 95% CI
Bowering	76.5	3.4	117	85.9	0.08	429	98.5%	-9.40 [-10.02, -8.78]	
Bray Moseley	61.3	18.17	21	87	19.92	14	0.2%	-25.70 [-38.71, -12.69]	
Linder	90.1	9.05	30	88.82	12.27	30	1.3%	1.28 [-4.18, 6.74]	+
Total (95% CI)			168	043-17		473	100.0%	-9.30 [-9.91, -8.69]	
Heterogeneity: Chi² = 20.65, df = 2 (P < 0.0001); l² = 90% Test for overall effect: Z = 29.81 (P < 0.00001)									-100 -50 0 50 100 Favours [clbp] Favours [healthy control]

Legend: IV = inverse variance; CI= confidence interval. For Linder et al. (2015) combined data for left/right laterality judgements were used for quantitative pooling; for Bowering et al (2014) only the data from "current back pain" patients were used. The data from Bowering et al (2014) was calculated from a graph using the software DigitizeIt<sup>®</sup>.

# Movement Control Tests (MCTs): Known-Groups Validity

Two studies of fair methodological quality (Luomajoki et al., 2008, Luomajoki and Moseley, 2011) (CLBP=91, HC=147) investigated the known-groups validity of MCTs

(see supplementary data: Appendix 6). Both studies independently, and when quantitatively pooled (figure 4), found a statistically poorer performance in the MCT performance by people with CLBP compared to healthy controls. The degree of heterogeneity was low and, likely, of no importance (Higgins and Green, 2011). Thus, there is moderate evidence from two studies of fair methodological quality that MCTs demonstrate known-groups validity.

# Figure 4: Forest Plot comparing movement control tests regarding their knowngroups validity to distinguish between people with CLBP and healthy controls

		clbp		healt	hy cont	trol		Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% CI	IV, Fixed, 95% CI
Luomajoki et al	2.37	1.34	46	0.75	1.03	102	56.6%	1.62 [1.18, 2.06]	
Luomajoki Moseley	3	1.1	45	1	1.3	45	43.4%	2.00 [1.50, 2.50]	
Total (95% CI)			91			147	100.0%	1.78 [1.46, 2.11]	◆
Heterogeneity: Chi² = Test for overall effect:	1.27, df Z = 10.6	= 1 (P 67 (P <	= 0.26) 0.0000	-4 -2 0 2 4 Favours [clbp] Favours [healthy control]					

Legend: IV = inverse variance; CI= confidence interval

## Graphesthesia: Known-Groups Validity

One study (Wand et al., 2010) (CLBP=19, HC=19) of fair methodological quality evaluated the known-groups validity of graphesthesia (see supplementary data: Appendix 7). Performance, as adjudged by letter recognition error rates, was poorer in patients (mean difference; 6.1, 95% CI: 1.3 to 11.0). Thus, there was limited evidence from one study of fair methodological quality that graphesthesia demonstrates known-groups validity.

#### FreBaQ: Known-Groups Validity

One study, which investigated the FreBaQ reliability also (Wand et al., 2014b) looked at the FreBaQ known-groups validity (see supplementary data: Appendix 8). Due to the larger sample size gathered for this specific question (n=52 CLBP, n=52 HC) the study's methodological quality was rated fair (see supplementary data: Appendix 8). Data were presented as medians and inter-quartile ranges. The FreBaQ score (median [range]) for the patient and control group was 11 [0-26] and 0 [0–6], respectively, indicating poorer performance by patients (p<0.05). Thus, there is limited evidence from one study of fair methodological quality that the FreBaQ demonstrates known-groups validity.

#### **Convergent Validity**

Studies that investigated the convergent validity of TPD, laterality judgement and BID were found.

#### Two Point Discrimination and Laterality judgment: Convergent Validity

One study (n=17) of poor methodological quality investigated the convergent validity of TPD against laterality judgement in people with CLBP (Stanton et al., 2013) (see supplementary data: Appendices 4 and 5). An increase in the TPD threshold by 1 mm was associated with a decrease in accuracy of 0.6% ( $\beta$ = - 0.06, 95% CI: 0.80 to 0.43). However, as this body of evidence consists of only one poor quality study, the quality of evidence for convergent validity of TPD and laterality judgment was classified as unknown.

#### Two-Point Discrimination and Body Image Drawings: Convergent Validity

Two studies, one of fair (Nishigami et al., 2015) (CLBP= 42) and one of poor methodological quality (Moseley, 2008) (CLBP=6) reported aspects of convergent validity for BIDs compared to TPD in a qualitative manner. Nishigami et al. (2015) displayed TPD values for participants with CLBP who drew either a normal, expanded or shrunken body image. These data suggested that TPD was increased in patients who reported an expanded BID. Similarly, Moseley (Moseley, 2008) reported an increase of the TPD threshold corresponding to the zone of the absence or disruption of the BIDs. However, as neither study quantified the relationship between TPD and BIDs, no evidence statement can be drawn regarding the convergent validity of BIDs.

#### Discussion

The aim of this study was to systematically investigate the reliability and validity of SMIs in people with CLBP. Ten studies investigating the following six SMIs were included; TPD, laterality judgment, MCTs, BIDs, FreBaQ, and graphesthesia. The methodological quality ranged from poor to good with only one study rated good (Bowering et al., 2014). The SMIs with the strongest support in the literature were TPD, laterality judgment and MCTs. There was moderate evidence to support the known-groups validity of these three SMIs. However, in general, there was a lack of high-quality studies investigating the clinimetric properties of SMIs for people with
CLBP. Hence, data collected using these techniques should be interpreted cautiously.

Only three studies assessed reliability comprising two SMIs, the FreBaQ and laterality judgement. All three studies were graded as methodologically poor, primarily due to small sample sizes. Thus, the level of evidence for the reliability and/or measurement error of FreBaQ and laterality judgement was unknown. For all other outcome measures, there were no studies to inform an evidence statement regarding reliability or measurement error. Regarding convergent validity, there were four poor quality studies, investigating the convergent validity of either TPD, laterality judgement or BIDs. Thus, the level of convergent validity for these SMIs was considered unknown, and for the other outcome measures, there were no studies to inform an evidence statement. The state of the evidence only allowed for statements to be made with respect to the known-groups validity of the SMIs.

TPD, laterality judgement and MCTs demonstrated the strongest evidence of knowngroups validity. Regarding TPD, our meta-analysis demonstrated a mean difference between healthy controls and people with CLBP of 16mm. This is broadly in keeping with a previous meta-analysis by Catley et al. (2014) which compared TPD performance between healthy controls and people with CLBP reporting a mean difference of 11.7mm (95% CI:5.5 mm to 17.8 mm). The differences between our results and those of Catley et al. (21) are explained by the exclusion of the vertical TPD measurements from Luomajoki and Moseley (Loumajoki and Moseley, 2011) from the present meta-analysis and by the exclusion of the results from Moseley et al. (2008) as this study was rated of poor methodological quality in our review. Metaanalysis for laterality judgement and MCTs demonstrated evidence for known-groups

validity of both outcome measures. However, there are no previous meta-analyses with which to compare our findings.

There was less evidence regarding graphesthesia, the FreBAQ and BIDs. One study of fair methodological quality implied a degree of known-groups validity for graphesthesia which was in line with results from studies investigating this technique in other clinical conditions, such as Parkinson disease (Jobst et al., 1997) and corticobasal degeneration (Drago et al., 2010). The results from one study of fair methodological quality demonstrated a degree of known-groups validity for the FreBaQ. A further study (Wand et al., 2016), published after the search cut-off date for the present review, investigated the psychometric properties of the FreBAQ by means of a Rasch analysis in 255 people with CLBP demonstrating adequate internal consistency with a person reliability index of 0.74 and a Cronbach's Alpha Value of 0.80. As adequate internal consistency is an essential prerequisite for questionnaires which intend to measure a single underlying construct by multiple items (Terwee et al., 2007), the results provide a basis for further psychometric evaluation. There was very limited evidence upon which to make any recommendations regarding the use of BIDs in people with CLBP.

## **Review limitations**

The search was restricted to full peer-reviewed published articles to enhance quality control, thus relevant conference papers/grey literature may have been excluded. Only one author undertook the screening and selection process increasing the risk of inadvertently excluding relevant studies. Additionally, only one reviewer extracted the

data from the included studies, which increases the risk of errors in the extraction process.

There were variations in the data collection protocols reported in the reviewed studies. In some cases, the variations were quite marked. For example, Nishigami et al. (2015) measured TPD performance as side-to-side differences, which was very different to the TPD protocols used in the other included studies. However, the protocol differences between the other studies were more subtle (see supplementary data: Appendix 2). These variations in data collection protocols may have reduced the comparability of the studies. In addition, the studies included in this review tended to have small sample sizes, which can lead to over-inflated effect sizes. This may have influenced the results of our meta-analyses.

The level of heterogeneity was substantial when the studies for laterality judgment were pooled for both reaction time and accuracy, thus these meta-analyses should be interpreted cautiously.

A key issue affecting the strengths of recommendations which can be drawn from this systematic review was the quality of included studies. Only one included study was rated as being of good methodological quality. In addition, the degree of reliability could not be established for any of the measures investigated in this review. Reliability is an important prerequisite of validity. Thus, the validity data presented should be interpreted cautiously. There is a need for higher quality studies investigating the clinimetric properties of SMIs for people with CLBP, with particular attention to recruiting adequate sample sizes.

## Conclusion

There was a lack of high quality studies investigating the clinimetric properties of SMIs in people with CLBP. The methodological quality of the studies were predominately rated as poor or fair, with a small sample size frequently the reason for a low rating. The strongest body of evidence currently exists for TPD, laterality judgment and MCT with respect to known-groups validity. However, as the reliability of these measurement tools have yet to be established, this validity data should be interpreted cautiously. There is an urgent need to undertake high quality studies investigating the clinimetric performance of SMIs for people with CLBP in order to guide clinical and research practice. Given the state of the evidence, data collected using these SMIs should be treated cautiously.

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#### Supplementary Data (e-Supplements):

#### Appendix 1: Search Strategies for all databases

Search Strategy Medline with search filter for measurement properties

#1 "Sensory acuity"[tw] OR "Sensory perception"[tw] OR "sensory threshold"[tw] OR "Tactile acuity"[tw] OR "Tactile threshold"[tw] OR "tactile perception"[tw] OR "tactile discrimination"[tw] OR "tactual discrimination"[tw] OR "pressure sensitivity"[tw] OR "pressure sensibility"[tw] OR "proprioceptive"[tw] OR "acuity"[tw] OR "touch sensitivity"[tw] OR "tactile sensation\*"[tw] OR "tactile sensitivity"[tw] OR "tactile sensibility"[tw] OR "depth-sense threshold"[tw] OR "perception threshold"[tw] OR "Discrimination sensation"[tw] OR "discriminative sensations"[tw] OR "Touch perception"[tw] OR "sensorimotor performance"[tw] OR "sensorimotor competence"[tw] OR "distorted body image"[tw] OR "Body schema"[tw] OR "physical self-awareness"[tw] OR "primary somatosensory cortex"[tw] OR "primary sensory cortex"[tw] OR "sensory-motor incongruence"[tw] OR "S1"[tw] OR "S1 representation"[tw] OR "Cortical reorganisation"[tw] OR "Cortical reorganization"[tw] OR "Consciousness"[tw] OR Neuroimaging[tw] OR "Neuronal plasticity"[tw] OR "cortical body map"[tw] OR "Touch Perception"[MeSH Terms] OR "Touch/physiology"[MeSH Terms] OR "Recognition (Psychology)"[MeSH Terms] OR "Pain Perception"[MeSH Terms] OR "Discrimination Learning"[MeSH Terms] OR "Discrimination (Psychology)/physiology\*"[MeSH Terms] OR "Perception/physiology"[MeSH Terms] OR "proprioception/physiology"[MeSH Terms] OR "Pain Threshold" [MeSH Terms] OR "Pattern Recognition, Physiological" [MeSH Terms] OR "Brain mapping" [MeSH Terms]

#2 back pain[tw] OR backache[tw] OR lumbago[tw] OR sciatic[tw] OR sciatica[tw]
OR "low back disorder "[tw] OR "low back pain"[tw] OR "Chronic low back pain"[tw]
OR "lower back pain"[tw] OR "non-specific low back pain"[tw] OR "NSCLBP"[tw] OR
"back injury"[tw] OR "lumbar spine dysfunction"[tw] OR "Back Pain"[MeSH Terms]
OR Sciatica[MeSH Terms] OR "Low Back Pain"[MeSH Terms] OR "Low Back
Pain/physiopathology\*"[MeSH Terms]

#3 Sensorymotor\*[tw] OR Sensorimotor\*[tw] OR "Sensory-motor\*"[tw] OR Sensomotor[tw] OR Sensomotoric[tw] OR "sensori-motor"[tw] OR "sensoryperceptual-motor\*"[tw] OR "sensory discrimination"[tw] OR "tactile stimulation"[tw] OR "Tactile assessments" [tw] OR "tactile perceptual tasks" [tw] OR "tactile tests" [tw] OR "sensory tests"[tw] OR "sensory testing"[tw] OR "somatosensory task"[tw] OR "somatosensory testing"[tw] OR "Test"[tw] OR "Testing"[tw] OR task[tw] OR "Somatosensory Cortex"[MeSH Terms] OR "Motor Cortex"[MeSH Terms] OR "Physical Stimulation" [MeSH Terms] OR "Two point discrimination" [tw] OR "Twopoint discrimination"[tw] OR "Two-point-discrimination"[tw] OR "two-point thresholds"[tw] OR "TPD threshold"[tw] OR "2-point discrimination"[tw] OR "2-PD"[tw] OR "TPD"[tw] OR "T.P.D"[tw] OR "discrimination threshold" OR Graphaestesia[tw] OR graphesthesia[tw] OR graphestesia[tw] OR "lumbopelvic motor control"[tw] OR "Movement Control"[tw] OR "Lumbopelvic control"[tw] OR "Movement test"[tw] OR "Motor Activity/physiology\*"[MeSH Terms] OR "Lumbosacral Region"[MeSH Terms] OR "Motor Skills\*"[MeSH Terms] OR "Movement/physiology"[MeSH Terms] OR "Movement Disorders" [MeSH Terms] OR "Body image drawing" [tw] OR "motor imagery"[tw] OR "motor imagery task"[tw] OR "Body schema"[tw] OR "bodyperception"[tw] OR "Body image assessment"[tw] OR "Body image perception"[tw] OR "body image"[MeSH Terms] OR "motor imagery"[tw] OR "left/right judgment"[tw]

OR "left/right judgement"[tw] OR "Recognition (Psychology)"[MeSH Terms] OR "Functional Laterality"[MeSH Terms]

#4 instrumentation[sh] OR methods[sh] OR Validation Studies[pt] OR Comparative Study[pt] OR "psychometrics"[MeSH] OR psychometr\*[tiab] OR clinimetr\*[tw] OR clinometr\*[tw] OR "outcome assessment (health care)"[MeSH] OR outcome assessment[tiab] OR outcome measure\*[tw] OR "observer variation"[MeSH] OR observer variation[tiab] OR "Health Status Indicators"[MeSH] OR "reproducibility of results"[MeSH] OR repoducib\*[tiab] OR "discriminant analysis"[MeSH] OR reliab\*[tiab] OR unreliab\*[tiab] OR valid\*[tiab] OR coefficient[tiab] OR homogeneity[tiab] OR homogeneous[tiab] OR "internal consistency"[tiab] OR (cronbach\*[tiab] AND (alpha[tiab] OR alphas[tiab])) OR (item[tiab] AND (correlation\*[tiab] OR selection\*[tiab] OR reduction\*[tiab])) OR agreement[tiab] OR precision[tiab] OR imprecision[tiab] OR "precise values"[tiab] OR test-retest[tiab] OR (test[tiab] AND retest[tiab]) OR (reliab\*[tiab] AND (test[tiab] OR retest[tiab])) OR stability[tiab] OR interrater[tiab] OR inter-rater[tiab] OR intrarater[tiab] OR intrarater[tiab] OR intertester[tiab] OR inter-tester[tiab] OR intratester[tiab] OR intratester[tiab] OR interobserver[tiab] OR inter-observer[tiab] OR intraobserver[tiab] OR intra-observer[tiab] OR intertechnician[tiab] OR inter-technician[tiab] OR intratechnician[tiab] OR intra-technician[tiab] OR interexaminer[tiab] OR interexaminer[tiab] OR intraexaminer[tiab] OR intra-examiner[tiab] OR interassay[tiab] OR inter-assay[tiab] OR intraassay[tiab] OR intra-assay[tiab] OR interindividual[tiab] OR inter-individual[tiab] OR intraindividual[tiab] OR intra-individual[tiab] OR interparticipant[tiab] OR inter-participant[tiab] OR intraparticipant[tiab] OR intraparticipant[tiab] OR kappa[tiab] OR kappa's[tiab] OR kappas[tiab] OR repeatab\*[tiab] OR ((replicab\*[tiab] OR repeated[tiab]) AND (measure[tiab] OR measures[tiab] OR

findings[tiab] OR result[tiab] OR results[tiab] OR test[tiab] OR tests[tiab])) OR generaliza\*[tiab] OR generalisa\*[tiab] OR concordance[tiab] OR (intraclass[tiab] AND correlation\*[tiab]) OR discriminative[tiab] OR "known group"[tiab] OR factor analysis[tiab] OR factor analyses[tiab] OR dimension\*[tiab] OR subscale\*[tiab] OR (multitrait[tiab] AND scaling[tiab] AND (analysis[tiab] OR analyses[tiab])) OR item discriminant[tiab] OR interscale correlation\*[tiab] OR error[tiab] OR errors[tiab] OR "individual variability"[tiab] OR (variability[tiab] AND (analysis[tiab] OR values[tiab])) OR (uncertainty[tiab] AND (measurement[tiab] OR measuring[tiab])) OR "standard error of measurement"[tiab] OR sensitiv\*[tiab] OR responsive\*[tiab] OR ((minimal[tiab] OR minimally[tiab] OR clinical[tiab] OR clinically[tiab]) AND (important[tiab] OR significant[tiab] OR detectable[tiab]) AND (change[tiab] OR difference[tiab])) OR (small\*[tiab] AND (real[tiab] OR detectable[tiab]) AND (change[tiab] OR difference[tiab])) OR meaningful change[tiab] OR "ceiling effect"[tiab] OR "floor effect"[tiab] OR "Item response model"[tiab] OR IRT[tiab] OR Rasch[tiab] OR "Differential item functioning"[tiab] OR DIF[tiab] OR "computer adaptive testing"[tiab] OR "item bank"[tiab] OR "cross-cultural equivalence"[tiab]

#### #5 #1 AND #2 AND #3 AND #4

#6 #5 NOT ("addresses"[pt] OR "biography"[pt] OR "case reports"[pt] OR "comment"[pt] OR "directory"[pt] OR "editorial"[pt] OR "festschrift"[pt] OR "interview"[pt] OR "lectures"[pt] OR "legal cases"[pt] OR "legislation"[pt] OR "letter"[pt] OR "news"[pt] OR "newspaper article"[pt] OR "patient education handout"[pt] OR "popular works"[pt] OR "congresses"[pt] OR "consensus development conference"[pt] OR "consensus development conference, nih"[pt] OR "practice guideline"[pt]) NOT ("animals"[MeSH Terms] NOT "humans"[MeSH Terms]) Search Strategy Embase

**#1** (Sensory acuity or Sensory perception or sensory threshold or Tactile acuity or Tactile threshold or tactile perception or tactile discrimination or tactual discrimination or pressure sensitivity or pressure sensibility or proprioceptive or acuity or touch sensitivity or tactile sensation\$ or tactile sensitivity or tactile sensibility or depth-sense threshold or perception threshold or Discrimination sensation or discriminative sensations or Touch perception or sensorimotor performance or sensorimotor competence or distorted body image or Body schema or physical self-awareness or primary somatosensory cortex or primary sensory cortex or sensory-motor incongruence or S1 or S1 representation or Cortical reorganization or Cortical reorganisation or Neuroimaging or Neuronal plasticity or cortical body map).mp. or exp touch/ or exp recognition/ or exp nociception/ or exp proprioception/ or exp perceptive threshold/

**#2** (back pain or backache or lumbago or sciatic or sciatica or low back disorder or low back pain or chronic low back pain or lower back pain or non-specific low back pain or NSCLBP or back injury or lumbar spine dysfunction).mp.

**#3** (Sensorymotor\* or Sensorimotor\* or sensory-motor\* or sensori-motor\* or Sensomotor or Sensomotoric or sensory-perceptual-motor\* or sensory discrimination or tactile stimulation or Tactile assessments or tactile perceptual tasks or tactile tests or sensory tests or sensory testing or somatosensory task or somatosensory testing or test or testing or task or Two point discrimination or two-point thresholds or TPD threshold or 2-point discrimination or 2-PD or TPD or "T.P.D" or discrimination threshold or graphaestesia or graphesthesia or graphestesia or lumbopelvic motor control or movement control or Lumbopelvic control or Movement test or

Lumbosacral Region or Body image drawing or motor imagery or motor imagery task or Body schema or body-perception or Body image assessment or Body image perception or "left/right judgment" or "left/right judgement").mp. or exp motor dysfunction/ or exp "movement (physiology)"/ or exp recognition/ or exp body image/ or exp somatosensory cortex/ or exp motor cortex/ or exp motor activity/ or exp motor performance/

## #4 #1 AND #2 AND #3

**#5 4 NOT** (editorial or letter or conference abstract or conference paper or conference proceeding or conference review).pt. not (exp animal/ not exp human/)

#### Search Strategy CINAHL

**S1** TX Sensory acuity OR TX Sensory perception OR TX sensory threshold OR TX Tactile acuity OR TX Tactile threshold OR TX tactile perception OR TX tactile discrimination OR TX tactual discrimination OR TX pressure sensitivity OR TX pressure sensibility OR TX proprioceptive OR TX acuity OR TX touch sensitivity OR TX tactile sensation\* OR TX tactile sensitivity OR TX tactile sensibility OR TX depthsense threshold OR TX perception threshold OR TX Discrimination sensation OR TX discriminative sensations OR TX Touch perception OR TX sensorimotor performance OR TX sensorimotor competence OR TX distorted body image OR TX Body schema OR TX physical self-awareness OR TX primary somatosensory cortex OR TX primary sensory cortex OR TX sensory-motor incongruence OR TX "S1" OR TX "S1 representation" OR TX Cortical reorganization OR TX Cortical reorganization OR TX Consciousness OR TX Neuroimaging OR TX Neuronal plasticity OR TX cortical body map OR MH "Touch/PH" OR MH "Recognition Psychology" OR MH "Pain+" OR MH "Perception+" OR MH "Perception/PH" OR MH "Proprioception+/PH" OR MH "Pain Threshold" OR MH "Brain Mapping"

**S2** TX back pain OR TX backache OR TX lumbago OR TX sciatic OR TX sciatica OR TX low back disorder OR TX low back pain OR TX Chronic low back pain OR TX lower back pain OR TX non-specific low back pain OR TX NSCLBP OR TX back injury OR TX lumbar spine dysfunction OR MH "Back Pain+" OR MH "Sciatica" OR MH "Low Back Pain" OR MH "Low Back Pain/PP"

**S3** TX Sensorymotor\* OR TX Sensorimotor\* OR TX Sensory-motor\* OR TX Sensomotor OR TX Sensomotoric OR TX sensori-motor OR TX sensory-perceptualmotor\* OR TX sensory discrimination OR TX tactile stimulation OR TX Tactile assessments OR TX tactile perceptual tasks OR TX tactile tests OR TX sensory tests

OR TX sensory testing OR TX somatosensory task OR TX somatosensory testing OR TX Test OR TX Testing OR TX task OR MH "Physical Stimulation+" OR TX Two point discrimination OR TX Two-point discrimination OR TX Two-point-discrimination OR TX two-point thresholds OR TX TPD threshold OR TX 2-point discrimination OR TX 2-PD OR TX TPD OR TX T.P.D OR TX discrimination threshold OR TX Graphaestesia OR TX graphesthesia OR TX graphestesia OR TX lumbopelvic motor control OR TX Movement Control OR TX Lumbopelvic control OR TX Movement test OR MH "Motor Activity+/PH" OR MH "Motor Skills+" OR MH "Movement+/PH" OR MH "Movement Disorders+" OR TX Body image drawing OR TX motor imagery OR TX motor imagery task OR TX Body schema OR TX body-perception OR TX Body image assessment OR TX Body image perception OR MH "Recognition Psychology" OR MH "Body Image+" OR TX motor imagery OR TX left/right judgment OR TX left/right judgement

## S4 S1 AND S2 AND S3

**S5 S4 NOT** (PT biography OR PT case study OR PT commentary OR PT directories OR PT editorial OR PT interview OR PT legal case OR PT letter OR PT consumer/patient teaching materials OR PT practice guidelines) NOT ((MH "Animals+") NOT (MH "Human"))

#### Search Strategy CENTRAL

**#1** "Sensory acuity" OR "Sensory perception" OR "sensory threshold" OR "Tactile acuity" OR "Tactile threshold" OR "tactile perception" OR "tactile discrimination" OR "tactual discrimination" OR "pressure sensitivity" OR "pressure sensibility" OR proprioceptive OR acuity OR "touch sensitivity" OR "tactile sensation\*" OR "tactile sensibility" OR "tactile sensibility" OR "tactile sensibility" OR "tactile sensibility" OR "tactile sensitivity" OR "tactile sensitivity" OR "tactile sensitivity" OR "tactile sensibility" OR "depth-sense threshold" OR "perception threshold" OR "Discrimination sensation" OR "discriminative sensations" OR "Touch perception" OR "sensorimotor performance" OR "sensorimotor competence" OR "distorted body image" OR "Body schema" OR "physical self-awareness" OR "primary somatosensory cortex" OR "primary sensory cortex" OR "sensory-motor incongruence" OR "S1" OR "S1 representation" OR "Cortical reorganisation" OR "Cortical reorganization" OR Consciousness OR Neuroimaging OR "Neuronal plasticity" OR "cortical body map"

**#2** MeSH descriptor: [Touch Perception] explode all trees

#3 MeSH descriptor: [Touch] explode all trees and with qualifier(s): [Physiology -PH]

**#4** MeSH descriptor: [Recognition (Psychology)] explode all trees

- **#5** MeSH descriptor: [Pain Perception] explode all trees
- **#6** MeSH descriptor: [Discrimination Learning] explode all trees
- **#7** MeSH descriptor: [Discrimination (Psychology)] explode all trees and with qualifier(s): [Physiology PH]

#8 MeSH descriptor: [Perception] explode all trees and with qualifier(s):[Physiology - PH]

**#9** MeSH descriptor: [Proprioception] explode all trees and with qualifier(s): [Physiology - PH]

**#10** MeSH descriptor: [Pain Threshold] explode all trees

**#11** MeSH descriptor: [Pattern Recognition, Physiological] explode all trees

**#12** MeSH descriptor: [Brain Mapping] explode all trees

# #13 #1 OR #2 OR #3 OR #4 OR #5 OR #6 OR #7 OR #8 OR #9 OR #10 OR #11 OR #12

**#14** "back pain" OR backache OR lumbago OR sciatic OR sciatica OR "low back disorder" OR "low back pain" OR "Chronic low back pain" OR "lower back pain" OR "non-specific low back pain" OR "NSCLBP" OR "back injury" OR "lumbar spine dysfunction"

**#15** MeSH descriptor: [Back Pain] explode all trees

**#16** MeSH descriptor: [Sciatica] explode all trees

**#17** MeSH descriptor: [Low Back Pain] explode all trees

**#18** MeSH descriptor: [Low Back Pain] explode all trees and with qualifier(s): [Physiopathology - PP]

## #19 #14 OR #15 OR #16 OR #17 OR #18

**#20** Sensorymotor\* OR Sensorimotor\* OR "Sensory-motor\*" OR Sensomotor OR Sensomotoric OR "sensori-motor" OR "sensory-perceptual-motor\*" OR "sensory discrimination" OR "tactile stimulation" OR "Tactile assessments" OR "tactile perceptual tasks" OR "tactile tests" OR "sensory tests" OR "sensory testing" OR "somatosensory task" OR "somatosensory testing" OR "Test" OR "Testing" OR task OR "Two point discrimination" OR "Two-point discrimination" OR "Two-pointdiscrimination" OR "two-point thresholds" OR "TPD threshold" OR "2-point discrimination" OR "2-PD" OR "TPD" OR "T.P.D" OR "discrimination threshold" OR Graphaestesia OR graphesthesia OR graphestesia OR "lumbopelvic motor control" OR "Movement Control" OR "Lumbopelvic control" OR "Movement test" OR "Body image drawing" OR "motor imagery" OR "motor imagery task" OR "Body schema" OR "body-perception" OR "Body image assessment" OR "Body image perception" OR "motor imagery" OR "left/right judgment" OR "left/right judgement"

**#21** MeSH descriptor: [Somatosensory Cortex] explode all trees

#22 MeSH descriptor: [Motor Cortex] explode all trees

#23 MeSH descriptor: [Physical Stimulation] explode all trees

**#24** MeSH descriptor: [Motor Activity] explode all trees and with qualifier(s): [Physiology - PH]

**#25** MeSH descriptor: [Lumbosacral region] explode all trees

#26 MeSH descriptor: [Motor Skills] explode all trees

**#27** MeSH descriptor: [Movement Disorders] explode all trees

#28 MeSH descriptor: [Body Image] explode all trees

#29 MeSH descriptor: [Functional Laterality] explode all trees

**#30** MeSH descriptor: [Movement] explode all trees and with qualifier(s): [Physiology - PH]

**#31** MeSH descriptor: [Recognition (Psychology)] explode all trees

#32 #20 OR #21 OR #22 OR #23 OR #24 OR #25 OR #26 OR #27 OR #28 OR #29 OR #30 OR #31

#33 #13 AND #19 AND #32

Author (Year)	a. b	Instrument Assessor (Profession)	Patient position	Instructions/ Feedback	Administration
	C. d	Equipment Rehearsal			
Linder et al (2015)	a. b. c.	Laterality Judgement PT Program: Recognise Online <sup>™</sup> – Difficulty setting "Vanilla" – Images displayed against plain background, randomly rotated at 0°, 90° or 180° to either the left or right distributed equally regarding laterality and rotation. 2 instructional images before each part of testing	seated comfortably with elbows at 90°, palms facing downwards, either left or right hand used on keys A and D or left/right arrows	Instructions: Session 1: Oral instruction by PT Session 2 and 3: Written instructions Feedback: NA	60 trunk images; Participants should determine, whether the depicted trunk was moved, laterally, flexed or rotated to the left or right. After 5 s new image if no selection was made.
Bowering et al (2014)	a b c	<ul> <li>Laterality Judgement</li> <li>NA</li> <li>Program: Recognise</li> <li>Online<sup>™</sup> – Images of the back and control images, which contained the back with another body part;</li> <li>"Neutral Images" - displayed against plain background, randomly rotated at +90°, -90° or 180° to either the left or right, distributed equally</li> </ul>	Seated on a comfortable chair in front of the computer; hand on keys A and D	Instructions: Written instructions Feedback: Not provided during task	40 images; 2 identical testing blocks with 2 minute break between; after 8 s a new image was presented if no selection was made.

Appendix 2: Individual study data collection protocols (adapted from (Pin, 2014)

Author	a.	Instrument	Patient position	Instructions/	Administration
(Year)	b.	Assessor (Profession)		Feedback	
	C.	Equipment			
	d.	Rehearsal			
Brav and	d a.	regarding laterality and rotation. 2 instructional images displaying a left or right hand; Participants had to press the "a" key for left and "d" for right Laterality Judgement	Participants	NA	56 photographs randomly
Moseley (2011)	d. c.	NA Program: Recognise <sup>™</sup> ; 28 photographs of a male model in various positions; trunk rotated right between 5° and 90°; photographs were digitally mirrored to construct identical pictures of the same model in various degrees of left rotation ;56 pictures integrated into Recognise <sup>™</sup> Each trial preceded by practice trial of 80 pictures	positioned themselves so that they were comfortable; Index and middle finger of the dominant hand placed on key "a" for left and "d" for right.		displayed; 40 images per trial;2 blocks with 3minute break between; Participants had to sit quietly
Stanton et			Late	rality Judgement	
ai (2013)		TDD		om Bray and Moseley (2011)	Dilataral according to Opling a
ivisnigami	a.		NA	Subjects were instructed	Bilateral assessment; Caliper
	р.	NA		to say one when they	position: perpendicular with the
(2015)				perceived one point and	spine, transverse process of the

Author	a.	Instrument	Patient position	Instructions/	Administration
(Year)	b	Assessor (Profession)	-	Feedback	
	C.	Equipment			
	d	Rehearsal			
	d.	Plastic ruler with 1 mm precision NA		"two" when they perceived two.	most severe pain level and same opposite level centered between the two tips of the caliper Pressure: until first blanching of the skin; Testing Order: Ascending: Starting at 0 mm, 5 mm steps until subjects identified "2 points"; Descending: Starting from 10 cm; 5 mm steps until subjects identified "1 point"; Repetitions: 2 ascending, 2 descending tests; Values of 1 ascending and descending run per side were averaged; Determination of TPD threshold: Side-to side- difference: TPD value higher pain side – TPD value lower pain side
Stanton et al (2013)	a b c d	<ul> <li>TPD</li> <li>NA</li> <li>Plastic caliper ruler</li> <li>Sensory testing with monofilaments to assess potential hypoesthesia</li> </ul>	NA	NA	Bilateral assessment; Caliper position: horizontally on both sides of the back; between the first lumbar vertebra and iliac crest; Pressure: Supra-sensory threshold; non-noxious; Testing Order: Ascending: Starting from 10 mm; 5 mm increments Descending: Starting from 100 mm, 5 mm increments; Repetitions: 1 ascending, 1 descending test;

Author (Year)	a. Instrument b. Assessor (Profession) c. Equipment	Patient position	Instructions/ Feedback	Administration
	d. Rehearsal			
				Determination of TPD threshold: Mean of ascending/descending test per side; Average of right and left mean actual TPD threshold
Luomajoki et al (2011)	a. TPD b. Physiotherapist c. Plastic caliper ruler d. NA	NA	NA	Bilateral Assessment; Caliper position: Horizontally and vertically between the first lumbar vertebra and iliac crest; Pressure: NA; Testing Order: Ascending: Starting from 10 mm; 5 mm increments Descending: Starting from 100 mm, 5 mm increments; Repetitions: 1 ascending, 1 descending test; Catch trials to prevent guessing (expanding the calipers instead of contracting or vice versa); Determination of TPD threshold: Average of ascending/descending test
Wand et al (2010)	<ul> <li>a. TPD</li> <li>b. NA</li> <li>c. Lafayette two-point aesthesiometer, (Lafayette Instruments, Lafayette, IN, USA), 1 mm precision</li> <li>d. NA</li> </ul>	Positioned comfortably in prone lying on an examination table with back exposed Pillow under stomach to flatten the lumbar spine	Subjects were instructed to say 'one' when they felt one point and 'two' when they felt two points.	Bilateral Assessment; Caliper position: Parallel to the spine; L3 transverse process in the centre of the caliper; Pressure: until first blanching of the skin; Testing Order: Ascending: Starting at 0 mm, 2 mm steps until subjects identified "2 points"; Descending: Starting from a start point "well

Author (Year)	a. Instrument b. Assessor (Profession) c. Equipment d. Rebearsal	Patient position	Instructions/ Feedback	Administration
		and to standardized lumbar position.		above the initial ascending threshold value"; 2 mm steps until patients identified "1 point"; Repetitions: NA; Determination of TPD threshold: Testing continued around initial values using ascending and descending sequences until a consistent response was obtained; Catch trials used to prevent guessing
Moseley (2008)	a. TPD b. NA c. Mechanical caliper with 1mm precision d. NA	prone	Subjects were instructed to say 'one', when one point was felt, 'two', when two points were felt.	Bilateral Assessment; caliper position: 16 levels from the fourth thoracic vertebra to the bottom of the gluteal folds; Medial point was 1, 2, 3 cm from midline; Pressure: until first blanching of the skin; Testing Order: Ascending: Starting at 0 mm, gradually increasing until patients identified "2 points"; Descending: NA; Level was randomised and counterbalanced; side was alternated until 6 measures (3 each side) were obtained; Repetitions: 3 measures per level and side; Determination of TPD threshold: Average of 1 ascending and descending series

Author (Year)	a. Instrument b. Assessor (Profession) c. Equipment	Patient position	Instructions/ Feedback	Administration
	d. Rehearsal			
				Average of 3 measures per level and side were used for analysis
Luomajoki et al (2008)	<ul> <li>a. MCT</li> <li>b. PTs, on average 7 years working experience; not blinded to subject's group</li> <li>c. NA</li> <li>d. Subjects did not know the tests before</li> </ul>	Depending on test starting position; Subjects wore underwear to allow inspection of the entire spine, hips and lower extremities	Verbal standardized instructions specified for each test; If the subject did not understand how to perform the test, the examiner explained and demonstrated it again;	<ul> <li>Testing order: Waiter's bow, Pelvic Tilt, One leg stance, Single knee extension, Quadruped position:</li> <li>Rocking backward</li> <li>Rocking forward</li> <li>Prone lying active knee flexion; Repetitions: 3 trials permitted; Rating Protocol: Clear movement dysfunction was rated as "not correct" and scored "1"; correct movements were scored"0"; If the movement control improved by instruction and correction, it was considered not a relevant movement dysfunction.</li> </ul>
Luomajoki and Moseley (2011)	a. TPD b. Trained PT c. NA d. NA	Depending on test starting position;	Subjects were given a picture in which a model demonstrated the target alignment of the pelvis and lumbar spine as a reference	Testing Order: Battery of six tests, referencing Luomajoki et al (2008); Repetitions: NA; Reference to Luomajoki et al (2008); Rating Protocol: Scores ranging from 6-0; 6 demonstrated the poorest movement control performance

Author	a.	Instrument	Patient position	Instructions/	Administration
(Year)	b.	Assessor (Profession)		Feedback	
	C.	Equipment			
	d.	Rehearsal			
Wand et al	а	Graphesthesia	positioned	Subjects were asked to	Testing order:
(2010)	b	NA	comfortably in	identify the letters drawn	Letters were drawn on three sites
	C.	Blunt end of monofilament	prone lying on an	on the back	centred on the tips of the L1, L3
	d	Subjects were first shown a	examination table		and L5 transverse processes; did
		wall chart o upper case	With back		not extend across the midline; the
		alphabet, letters and were	Dillow under		there was no everlep in the area of
		the way the letters would be	stomach to flatten		skin in which the letters were
		drawn They were then	the lumbar spine		drawn between the 3 sites: 20
		shown a diagram of the	and to		random letters at each level of the
		lumbar spine depicting the	standardised		3 sites: 3 sites were tested in
		orientation and location of	lumbar position.		random order: Error rate out of 60
		the letters			was calculated for each side of the
					back
Moseley et	a.	BID	Subjects stood in	'Concentrate on your back.	On request after instruction
al (2008)	b.	NA	front of a waist	Add to this drawing by	
	C.	a line drawing showing the	high bench	following the outline of	
		posterior surface of the back		your own back as you	
		with only the top and bottom		track it in your mind.	
	ام	of the picture drawn		Concentrate on where you	
	α.	NA		draw in the wortebre that	
				vou can fool. Do this	
				without touching your	
				back Your drawing should	
				relate to your own sense	
				of vour back. Don't draw	
				any part you can't sense.	

Author (Year)	a. Instrument b. Assessor (Profession)	Patient position	Instructions/ Feedback	Administration
	c. Equipment d. Rehearsal			
			Do not draw what you think your back looks like – draw what it feels like."	
Nishigami et al (2015)	<ul> <li>a. BID</li> <li>b. NA</li> <li>c. a line drawing showing the posterior surface of the back with only the top and bottom of the picture drawn (acc. to Moseley et al (2008))</li> <li>d. NA</li> </ul>	Subjects were asked to sit in a chair	"Concentrate on your back. Add to this drawing by following the outline of your own back as you track it in your mind. Concentrate on where you feel your back to be. Also draw in the vertebra that you can feel. Do this without touching your back. Do not draw any part you cannot sense. Do not draw what you think your back looks like- draw what it feels like."	On request after instruction
Wand et al (2014)	a. FreBAQ b. NA c. NA d. NA	NA	Written instructions	First test: On site; Second test: Take home copy; filled out and posted one week later
Legend: NA body image	= information not available; PT = drawings; FreBAQ= Fremantle Ba	physiotherapist; TPD ick Awareness Ques	= Two-Point Discrimination; N tionnaire	MCT = Movement Control Test; BID=

## Appendix 3: Reviewer Agreement

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological	
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating	
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)	
TPD Known- Groups Validity						
Luomajoki and Moseley (2011)	1	2	2	1	2	
	2	2	1	0	1	
	3	1	2	1	2	
	4	3	3	1	3	
	5	3	3	1	3	
	6	2	2	0	2	
	7	n.a. †	n.a.	1	n.a.	
	8	n.a.	n.a.	1	n.a.	
	9	3	3	0	3	

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	10	2	3	1	2
Stanton et al (2013)	1	2	2	1	2
	2	1	1	1	1
	3	1	1	1	1
	4	3	2	0	2
	5	3	2	0	2
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	0	1	0	1
	10	3	3	1	3

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
Nishigami et al (2015)	1	2	2	1	2
	2	1	1	1	1
	3	2	2	1	2
	4	3	1	0	1
	5	3	2	0	2
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	1	1	1	1
	10	3	3	1	3
Moseley (2008)	1	2	2	1	2
	2	2	2	1	2

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	3	0	0	1	0
	4	2	1	1	1
	5	3	2	0	2
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	1	1	1	1
	10	2	0	0	0
Wand et al (2010)	1	2	2	1	2
	2	3	1	0	1
	3	1	1	1	1
	4	1	3	0	3

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	5	3	3	1	3
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	3	1	0	1
	10	3	3	0	3
TPD Convergent Validity					
Luomajoki and Moseley (2011)	1	2	2	1	2
	2	2	1	0	1
	3	2	2	1	2
	4	3	3	1	3
Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
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	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	5	3	3	1	3
	6	2	2	1	2
	7	3	2	0	3
	8	1	1	1	1
	9	3	1	0	1
	10	2	3	0	2
Stanton et al	1	2	2	1	2
(2013)					
	2	1	1	1	1
	3	0	0	1	0
	4	3	3	1	3
	5	3	3	1	3
	6	2	2	1	2

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	7	3	3	1	3
	8	1	1	1	1
	9	0	1	0	0
	10	3	3	1	3
Wand et al	1	2	2	1	2
(2010)	2	3	1	0	1
	3	0	0	1	0
	4	1	3	0	3
	5	3	3	1	3
	6	2	2	1	2
	7	3	3	1	3
	8	1	0	0	0

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	9	3	1	0	1
	10	3	3	1	3
MCT Known- Groups Validity					
Luomajoki and Moseley (2011)	1	2	2	1	2
	2	2	1	0	1
	3	1	2	0	2
	4	3	3	1	3
	5	3	3	1	3
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	9	3	1	0	1
	10	2	3	0	2
Luomajoki et al (2008)	1	2	2	1	2
	2	1	1	1	1
	3	3	3	1	3
	4	2	2	1	2
	5	2	2	1	2
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	1	1	1	1

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	10	3	2	0	2
MCT Convergent Validity					
Luomajoki and Moseley (2011)	1	2	2	1	2
	2	2	1	0	1
	3	2	2	1	2
	4	3	3	1	3
	5	3	3	1	3
	6	2	2	1	2
	7	3	2	0	3
	8	1	1	1	1
	9	3	1	0	1

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	10	2	3	0	2
Graphesthesia Known- Groups Validity					
Wand et al (2010)	1	2	2	1	2
	2	3	1	0	1
	3	1	1	1	1
	4	1	3	0	3
	5	3	3	1	3
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	9	3	1	0	1
	10	3	3	1	3
Graphesthesia Convergent Validity					
Wand et al (2010)	1	2	2	1	2
	2	3	1	0	1
	3	2	2	1	2
	4	1	3	0	3
	5	3	3	1	3
	6	2	2	1	2
	7	3	3	1	3
	8	1	1	1	1

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	9	3	1	0	1
	10	3	3	1	3
Laterality Judgement Known- Groups Validity					
Stanton et al (2013)	1	2	2	1	2
	2	1	1	1	1
	3	0	0	1	0
	4	3	3	1	3
	5	3	3	1	3
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	8	n.a.	n.a.	1	n.a.
	9	0	1	0	0
	10	3	3	1	3
Bowering et al (2014)	1	2	3	0	3
	2	3	3	1	3
	3	3	3	1	3
	4	3	3	1	3
	5	3	3	1	3
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	3	3	1	3

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	10	3	3	1	3
Bray and Moseley (2011)	1	2	2	1	2
	2	2	1	0	1
	3	0	1	0	1
	4	3	3	1	3
	5	3	3	1	3
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	1	1	1	1
	10	3	3	1	3

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
Linder et al (2015)	1	2	2	1	2
	2	1	1	1	1
	3	1	2	0	2
	4	1	1	1	1
	5	2	2	1	2
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	3	3	1	3
	10	3	3	1	3

Laterality Judgement

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
Convergent Validity					
Stanton et al (2013)	1	2	2	1	2
	2	1	1	1	0
	3	0	0	1	0
	4	3	3	1	3
	5	3	3	1	3
	6	2	2	1	2
	7	3	3	1	3
	8	1	1	1	1
	9	0	1	0	0
	10	3	3	1	3

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
Laterality Judgement Reliability					
Linder et al (2015)	1	2	2	1	2
	2	1	1	1	1
	3	1	0	0	0
	4	3	3	1	3
	5	2	3	0	3
	6	3	3	1	3
	7	2	1	0	2
	8	3	3	1	3
	9	0	1	0	1
	10	3	1	0	1

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	11	3	2	0	3
	12	n.a.	n.a.	1	n.a.
	13	n.a.	n.a.	1	n.a.
	14	n.a.	n.a.	1	n.a.
Bray and Moseley (2011)	1	2	2	1	2
	2	2	1	0	1
	3	0	0	1	0
	4	3	3	1	3
	5	3	3	1	3
	6	3	3	1	3
	7	2	2	1	2
	8	3	3	1	3

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	9	2	2	1	2
	10	1	1	1	1
	11	2	2	1	2
	12	n.a.	n.a.	1	n.a.
	13	n.a.	n.a.	1	n.a.
	14	n.a.	n.a.	1	n.a.
Laterality Judgement Measurement Error					
Linder et al (2015)	1	2	2	1	2
	2	1	1	1	1
	3	2	0	0	0

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	4	3	3	1	3
	5	3	3	1	3
	6	3	3	1	3
	7	1	2	0	2
	8	3	3	1	3
	9	1	1	1	1
	10	1	1	1	1
	11	2	2	1	2
BIDs Known Groups Validity					
Nishigami et al (2015)	1	2	2	1	2
	2	1	1	1	1

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	3	2	2	1	2
	4	3	1	0	1
	5	3	2	0	2
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	1	1	1	1
	10	3	3	1	3
Moseley (2008)	1	2	2	1	2
	2	2	2	1	2
	3	0	0	1	0
	4	2	1	0	1
	5	3	2	0	2

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	6	2	2	1	2
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	1	1	1	1
	10	2	0	0	0
FreBaQ Known					
Groups Validity					
Wand et al (2014)	1	2	2	1	2
	2	1	1	1	1
	3	3	3	1	3
	4	1	1	1	1

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	Item	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	5	2	2	1	2
	6	2	2	1	3
	7	n.a.	n.a.	1	n.a.
	8	n.a.	n.a.	1	n.a.
	9	3	3	1	3
	10	3	3	1	3
FreBaQ Reliability					
Wand et al (2014)	1	2	2	1	2
	2	1	1	1	1
	3	0	0	1	0
	4	3	3	1	3
	5	2	2	1	2

Author (year)	COSMIN	Methodological	Methodological	Rater Agreement	Final Methodological
	ltem	Quality Rating	Quality Rating	1=agreement	Quality Rating
		Reviewer 1	Reviewer 2	0=disagreement	(consensus)
	6	3	3	1	3
	7	0	2	0	0
	8	3	3	1	3
	9	1	1	1	1
	10	3	1	0	1
	11	3	3	1	3
	12	n.a.	n.a.	1	n.a.
	13	n.a.	n.a.	1	n.a.
	14	n.a.	n.a.	1	n.a.

Legend: TPD= Two-Point Discrimination; n.a. = not applicable; MCT= Movement Control Test; BIDs=Body Image Drawing; FreBaQ=Fremantle Back Awareness Questionnaire; Reviewer 1=KE; Reviewer 2=CR; Coding methodological quality rating: 0 = poor; 1=fair; 2=good; 3=excellent

## Appendix 4: Summary of TPD measurement properties (known-groups and convergent validity)

Author (Year)	a) Instrument b) Design c) n	esign Measurement Property Result		Mean Difference (95%Cl) [p-value]	COSMIN Score
		Mean (SD) [mm]			
		CLBP	HC		
Luomajoki and Moseley (2011)	a) TPD b) Known-Groups Validity c) CLBP = 45 HC = 45	61 (13)	44 (10)	<b>17</b> * (12.14 to 21.86) [p < 0.01]	fair
Stanton et al (2013)	a) TPD b) Known-Groups Validity c) CLBP =17 HC =18	59.8 (11.7)	45.3 (5.1)	<b>14.50</b> (8.34 to 20.65) [p < 0.0001]	fair
		Related Construct			
	a) TPD b) Convergent Validity c) CLBP = 17	Laterality Reconstruction Accuracy (%): β (95%CI) -0.6 (-0.80 to -0.43)		not stated	poor
		CLBP Mean (SD) [mm]	HC Mean (SD) [mm]		
Moseley (2008)	a) TPD b) Known-Groups Validity c) CLBP = 6 HC = 10	48.83 (1.83)	47 (8)	1.83 (-5.38 to 9.037 ) [p = 0.59]	poor
Wand et al (2010)	a) TPD b) Known-Groups Validity	62.0 (21.6)	44.2 (13.7)	<b>17.80</b> (5.90 to 29.70) [p = 0.0045]	fair

Author (Year)	a) Instrument b) Design c) n	Measurement Property Result		Mean Difference (95%Cl) [p-value]	COSMIN Score				
	c) CLBP = 19 HC = 19								
Nishigami et al (2015)	<ul> <li>a) TPD (mean difference between sides)</li> <li>b) Known-Groups Validity</li> <li>c) CLBP = 42 HC = 17</li> </ul>	normal BI: 4.5 (5.5) expanded BI: 13.3 (6.8) shrink: 9.4 (7.0)	5.5 (3.8)	CLBP normal BI - HC: -1.00 (-4.27 to 2.27) [p = 0.54] CLBP expanded BI - HC: <b>7.80</b> (3.75 to 11.85) [p = 0.0005] shrink BI - HC: 3.90 (-0.23 to 8.03) [p = 0.06]	Fair				
TPD=Two-Point Discrimin HC=Healthy Controls; BI= *Bold figures indicate stat	Image: Point Discrimination; SD=Standard Deviation; 95%CI= 95%Confidence Interval; CLBP=Chronic Low Back Pain;         HC=Healthy Controls; BI= Body Image;         *Bold figures indicate statistically significant differences								

## Appendix 5: Summary of Laterality Judgement measurement properties (known-groups and convergent validity, reliability

## and measurement error)

Author (Year)	a) Instrument b) Design c) n	Measurement Property		Mean Difference (95%Cl) [p-value]	COSMIN Score
		Result Mean (SD)			
		CLBP	HC		
Linder et al (2015)	a) Laterality Judgement b) Known-Groups Validity c) CLBP = 30 HC = 30	ACC Right Trunk Rotation Mean (SD): S1: 88.7 (8.0) S2: 90.5 (9.8) S3: 90.6 (8.3) ACC Left Trunk Rotation Mean (SD): S1: 89.0 (9.1) S2: 89.1 (11.2) S3: 92.7 (7.4) RT right Trunk Rotation Mean (SD): S1: 1.89 (0.37) S2: 1.60 (0.34) S3: 1.57 (0.26) RT left Rotation Mean (SD): S1: 1.93 (0.43)	ACC Right Trunk Rotation Mean (SD): S1: 87.8 (20.2) S2: 87.9 (12.8) S3: 91.8 (6.6) ACC Left Trunk Rotation Mean (SD): S1:86.0 (10.4) S2: 89.0 (11.4) S3: 90.4 (7.9) RT right Trunk Rotation Mean (SD): S1: 2.01 (0.52) S2: 1.80 (0.53) S3:1.54 (0.34) RT left Rotation Mean (SD): S1: 2.01 (0.55)	ACC Right Trunk Rotation S1: 0.90 (-3.84 to 5.64) [p=0.90] S2: 2.600 (3.29 to 8.49) [p=0.38] S3: -1.20 (-5.08 to 2.68) [p=0.54] ACC Left Trunk Rotation: S1: 3.00 (-2.05 to 8.05) [p=0.24] S2: 0.10 (5.74 to 5.94) [p=0.97] S3: 2.30 (-1.66 to 6.26) [p=0.25]	fair

Author (Year)	a) Instrument b) Design c) n	Measurement Property		Mean Difference (95%Cl) [p-value]	COSMIN Score
		Result Mean (SD)			
		CLBP	HC		
		S2:1.64 (0.40) S3: 1.57 (0.33)	S2: 1.72 (0.47) S3: 1.55 (0.42)	RT right Trunk Rotation S1: -0.08 (-0.34  to  0.176) [p= 0.53] S2: -0.08 (0.31  to  0.15) [p= 0.50] S3: 0.02 (-0.18  to  0.22) [p= 0.84] RT left Rotation Mean : S1: -0.08 (-0.34  to  0.18) [p= 0.53] S2: -0.08 (-0.31  to  0.15) [p= 0.48] S3: 0.02 (-0.18  to  0.22) [p= 0.84]	

Author (Year)	a) Instrument b) Design c) n	Measurement Property		Mean Difference (95%Cl) [p-value]	COSMIN Score
		Result Mean (SD)			
		CLBP	HC		
Bowering et al (2014)*	<ul> <li>a) Laterality Judgement</li> <li>b) Known-Groups Validity</li> <li>c) Current back pain</li> <li>= 117 History of back pain = 462 HC = 429</li> </ul>	RT Trunk Rotation 1.89 (0.19) ACC Trunk Rotation: 76.5 (3.4)	RT Trunk Rotation 1.74 (0.07) ACC Trunk Rotation: 85.9 (0.08)	RT Trunk Rotation: <b>0.14</b> <sup>†</sup> (0.11 to 0.17) [p < 0.0001] ACC Trunk Rotation: -9.40 (-10.02 to -8.78) [p < 0.0001]	good
Bray and Moseley (2011)	a) Laterality Judgement b) Known-Groups Validity c) CLBP = 21 HC = 14	RT Trunk Rotation 2.4 (0.35) ACC Trunk Rotation bilateral pain: 53.4 (19.55) unilateral pain: 67.2 (15.27)	RT Trunk Rotation: 2.4 (0.33) ACC Trunk Rotation: 87 (19.92)	RT Trunk Rotation 0 (-0.24 to 0.24) [p=0] ACC Trunk Rotation: bilateral pain - HC: -33.60 (-47.43 to -19.77) [p < 0.0001] unilateral pain - HC: -19.80 ( -31.91 to -7.69) [p = 0.0022]	fair

Author (Year)	a) Instrument b) Design	Measurement Property		Mean Difference (95%Cl) [p-value]	COSMIN Score
	c) n	Result Mean (SD)			
		CLBP	HC		
		Related Construct			
Stanton et al (2013)	a) Laterality Judgement b) Convergent Validity c) CLBP = 17	TPD: β (95%Cl) -0.6 (-0.80 to -0.43)	1	n.a.	poor
				Result Reliability (95%CI)	
Bray and Moseley (2011)	a) Laterality Judgement b) Test-Retest Reliability: Mean Time interval (Range) [days]: 1 (1-7) c) CLBP = 10			RT Trunk Rotation: ICC <sub>2,1</sub> = 0.872 (0.731 - 0.951) ACC Trunk Rotation: ICC <sub>2,1</sub> = 0.920 (0.831 - 0.970)	poor

Author (Year)	a) Instrument b) Design c) n	Measurement Property		Mean Difference (95%Cl) [p-value]	COSMIN Score
		Result Mean (SD)			
		CLBP	HC		
				Result Reliability (95%CI)	
Linder et al (2015)	<ul> <li>a) Laterality Judgement</li> <li>b) Test-Retest Reliability Mean Time interval (Range) [days]: Session 1 to 2: 2.2 (2-5) Session 2-3: 2.4 (1-11)</li> <li>c) Session 1 and 2: CLBP = 25 Session 2 and 3: CLBP = 22</li> </ul>			Session 1 to 2: RT Trunk Rotation: ICC2,1= $0.51$ ( $0.15 - 0.75$ ) ACC Trunk Rotation: ICC2,1= $0.71$ ( $0.44 - 0.86$ ) Session 2 to 3: RT Trunk Rotation: ICC2,1= $0.91$ ( $0.79 - 0.96$ ) ACC Trunk Rotation: ICC2,1= $0.69$ ( $0.39 - 0.86$ )	poor

Author (Year)	a) Instrument b) Design c) n	Measurement Property		Mean Difference (95%Cl) [p-value]	COSMIN Score	
		Result Mean (SD)				
		CLBP	HC			
				Coefficient of Variation (95%CI):		
Linder et al (2015)	a) Laterality Judgement b) Measurement Error <sup>+</sup> c) CLBP =22			Session 1 to 2: RT Trunk Rotation: 19.6 (13.79-25.64) ACC Trunk Rotation: 6.46 (4.52-8.32) Session 2 to 3: RT Trunk Rotation: 6.23 (4.35-8.14) ACC Trunk Rotation: 6.77 (4.72-8.86)	poor	
ACC= Accu	uracy; RT= Reaction Tir	me; S1,S2, S3 =	Session 1,2,3; SD=	Standard Deviation; ICC <sub>2,1</sub> = Intraclass Cor	relation Coefficient;	
two-way ra	ndom model; 95%CI=	95%Confidence	Interval; CLBP=Chro	onic Low Back Pain; HC=Healthy Controls;	TPD= Two-Point	
Discriminat	tion; n.a.= not applicable	e;*Data from Bo	wering et al (2014) w	vere derived via DigitizeIt <sup>®</sup> and p-values we	re calculated	
online via C	online via GraphPad Quick Calcs software; <i>+Bold figures indicate statistically significant differences</i> ; <i>+</i> = minimal important change					
(MIC) data	was not provide					

## Appendix 6: Summary of Movement Control Test known-groups validity

Author	a) Instrument	Measurement Pro	perty Result		
(Year)	b) Design				
	c) n				
		Result		Mean Difference	COSMIN
		Mean (SD) [0-6]		(95%CI)	Score
			-	[p-value]	
		CLBP	HC		
Luomajoki and	a) Movement Control	3 (1.1)	1 (1.3)	2.00*	fair
Moseley (2011)	Test			(1.50 to 2.50)	
	b) Known-Groups			[p < 0.0001]	
	Validity				
	c) CLBP =45				
	HC = 45				
Luomajoki et al	a) Movement Control	CLBP	HC	1.62	fair
(2008)	Test	2.37 (1.34)	0.75 (1.03)	(1.22 to 2.02)	
	b) Known-Groups			[p < 0.0001]	
	Validity				
	c) LBP =102				
	CLBP=46				
	HC = 102				
SD= Standard Deviation	on; CLBP= chronic low ba	ack pain; HC= healthy	controls; 95%Cl	= 95% Confidence Inte	erval; [0-6] = six tests
are included in the test	t battery, the higher the se	core the worse the te	st performance;		
*Bold figures indicate s	statistically significant diffe	erences			

Appendix 7: Summary of Graphesthesia known-groups validity

Author (Year)	d) Instrument e) Design f) n	Measurement Property Result				
		Result Mean (SD) [Error Rate]		Mean Difference (95%CI) [p-value]	COSMIN Score	
		CLBP	HC			
Wand et al (2010)	<ul> <li>a) Graphesthesia</li> <li>b) Known-Groups Validity</li> <li>c) CLBP =19 HC = 19</li> </ul>	25.5 (8.0)	19.3 (6.8)	<b>6.1</b> * (1.3 to 11.0) [p = 0.01]	fair	
SD= Standard Deviation; 95%CI= 95% Confidence Interval; CLBP=Chronic low back pain; HC=Healthy Controls; *Bold figures indicate statistically significant differences						

Appendix 8: Summary of FreBaQ known-groups validity and reliability

Author (Year)	a) Instrument b) Design c) n	Measurement Property Result Mean [0-36] <i>Median (Range)</i>		Mean Difference (95%Cl) [p-value]	COSMIN Score
		CLBP	HC		
Wand et al (2014)	a) FreBAQ b) Known-Groups Validity c) CLBP =51 HC = 51	10.8 <i>11 (0-26)</i>	0.5 <i>0 (0-6)</i>	Mann-Whitney Test <b>11</b> * [p < 0.001]	fair
				Result Reliability (95%CI)	
Wand et al (2014)	a) FreBAQ b) Test-Retest Reliability (Mean Time interval: 1 week) c) CLBP =26			ICC <sub>2,1</sub> = 0.652 (0.307-0.848) (Agreement) ICC <sub>2,1</sub> = 0.667 (0.317-0.857) (Consistency)	poor
FreBAQ = Fremantle Back Awareness Questionnaire; CLBP= Chronic low back pain; HC= Healthy Controls; ICC <sub>2.1</sub> = intra class correlation coefficient (two-way random effect model with single measures); MIC=minimal important change *Bold figures indicate statistically significant differences					