Supplement Material

Application details for the single bedside-QST devices

Thermal perception/pain

Thermal parameters were investigated using four metal pieces with a size of 3x3 cm, approximately the size of the lab-QST thermode. Cold detection and cold pain thresholds were examined using metal pieces of 8°C (stored in a fridge) and 22°C (left at room temperature). Warm perception and heat pain thresholds were investigated using metal pieces heated up in a vial warmer ("Babycare," Breuer GmbH, Ulm, Germany) to 37°C and 45°C. Every metal piece was applied for approximately three seconds. First, patients were asked about the quality of the stimulus, i.e. perceived/not perceived, cold/warm (yes/no) and if perceived about the perception intensity (NRS, "0 = no perception" to "10 = strongest imaginable perception"). Afterwards, patients had to indicate whether the stimulus had also been painful (yes/no) and if painful, rate the pain intensity (NRS, "0 = no pain" to "10 = strongest imaginable pain").

Touch sensation

The dynamic mechanical detection sensitivity was examined by use of a Q-tip stroke of 5 cm first applied in the control area and directly afterwards in the most affected area. Subjects were asked to rate the perception intensity of the stimulus at the test area as compared to the control area from 0 to 20 ("0 = not perceived at test area", "<10 = less intensity at test area as compared to control", "10 = equal intensity as compared to control area", ">10 = stronger intensity at test area as compared to control", "10 = equal intensity as compared to control area", ">10 = stronger intensity at test area as compared to control").

Static mechanical detection was investigated by use of a Neuropen filament (monofilament with defined pressure of 10 g) and a 64-mN von Frey hair (Optihair2-Set; Marstock Nervtest). Both tools were applied three times in each area. Patients had to indicate the number of perceived stimuli, documented as "perceived at least two times" (yes/no) and to compare the perception intensity between both areas (i.e., 0 = same intensity in control and test area, 1 = control area more intense, and 2 = test area more intense).

Mechanical pain sensitivity

To test for pinprick hyperalgesia and hypoalgesia, a 0.7-mm CMS hair (CMS, Chicago Medical Supply, LLC) and a Neuropen with a Neurotip (disposable needle with a defined pressure of 40 g) were applied to both areas. Participants had to evaluate whether the stimulus was perceived as blunt touch or as a pinprick. If the stimulus was perceived as a pinprick, they had to rate the pain intensity (NRS, "0 = no pain" to "10 = worst imaginable pain").

Wind Up

Temporal pain summation as an indicator for central sensitization was investigated by applying the 0.7 mm CMS hair and the Neurotip at the control and the test area. Both devices were first applied once and subjects had to rate the single stimulus pain intensity (NRS, "0 = no pain" to "10 = worst imaginable pain"). Afterwards they were exposed to a series of 10 stimuli with a frequency of 1/s and had to indicate the pain intensity of the last few stimuli. Wind-up ratio was calculated as the ratio of the last few stimuli of the row of ten divided by the single stimulus pain intensity.

Dynamic mechanical allodynia

Dynamic mechanical allodynia was assessed using a Q-tip by drawing a cross twice with an angle of 90° and lines of 3-5 cm on the skin. Subjects had to indicate whether the stimulus was painful (yes/no) and if painful, rate the pain intensity (NRS, "0 = not painful" to "10 = worst imaginable pain"). Afterwards they were asked about any burning, prickling or unpleasant after sensation (yes/no) and in case of postallodynia sensation pain, the pain intensity was rated (NRS, "0 = not painful" to "10 = worst imaginable pain").

Pressure pain sensitivity

A bedside algometer (10-mL syringe sealed with a plug and felt pad with a contact area of 1 cm²) was placed above a muscle located in the control area and test area to evaluate deep somatosensory pressure pain. First the syringe was slowly compressed up to 4 mL and subjects had to indicate, whether this compression was painful (yes/no) and if painful, rate the pain intensity (NRS, "0 = no pain" to "10 = worst imaginable pain"). Afterwards the syringe was compressed again with 1mL/s and the participants had to evaluate when the pressure started being painful. This pain threshold was defined by the milliliter of compressed air in the syringe.

Vibration detection

Vibration detection threshold was investigated using a standardized tuning fork (64 Hz, 8-point scale), placed over a bony prominence located in the control and in the test area. Subjects had to state the moment when the vibration stimulus had disappeared (0 = no vibration stimulus perception, 8 = best possible vibration detection).

Parameters	ΤοοΙ	Description	Vendor/Address
Lab-QST			
CDT, WDT, TSL, PHS, CPT, HPT	Medoc TSA II system thermode	Temperature range: 0-50 °C, 1 °C/s ramp	Medoc Advance Medical System, Ramat, Yishai, Isreal
MDT	Von Frey filaments	Forces between 0.25 and 512 mN	Optihair2-Set, Marstock Nervtest, Heidelberg, Germany
MPT, MPS, WUR	Pinprick stimulators	Weighted pinpricks: forces between 128 mN to 256 mN	MRC Systems GmbH, Heidelberg, Germany
DMA	Brush, cotton wisp, cotton wool	Brush: force of 200 -400 mN	Somedic, Sweden
		Cotton wisp: force of 3 mN	
		Cotton wool tip fixed to an elastic strip: force of 100 mN	
PPT	Pressure Algometer	Probe area of 1 cm ² (probe diameter of 1.1 cm) that exerts pressure up to 20 kg/cm ² /200 N/cm ² /2000 kPa	FDN 200; Wagner Instruments, Greenwich, CT
		Increasing ramp of 50 kPa/s	
VDT	Tuning fork	128 Hz tuning fork with damper to generate a frequency of 64 Hz, 8/8 scale (Rydel-Seiffer)	
Bedside-QST			
Thermal perception/pain	Metal pieces	3x3 cm hot- rolled steel (thermal conductivity: ~52-55 W/m-K)	
	Metal cubes*	2.7x2.7x2.7 cm stainless steel (thermal conductivity: ~15 W/m-K)	e.g. Edelstahl Whisky Stein Eiswürfel Square Glacier Chiller Drink Wiederverwendbar eBay
Warm perception; Heat pain thresholds	Vial warmer	To heat up the metal pieces to 37°C and 45°C	e.g. Beurer BY 52 (95402) (<u>Beurer</u> <u>Babykost- und Fläschchenwärmer - BY</u> 52 Beurer Onlineshop (beurer-shop.de)

Supplement Table 1. Equipment details for lab- and bedside-QST parameters

Static mechanical detection	Neuropen filament	Monofilament with a defined pressure of 10 g	Neuropen Owen Mumford; https://www.owenmumford.com/en		
MPT, MPS, WUR	Neuropen with a Neurotip	Disposable needle with a defined pressure of 40 g	-		
Static mechanical detection	64 mN von Frey filament		Optihair2-Set, Marstock Nervtest, Heidelberg, Germany		
Mechanical pain sensitivity; Temporal pain summation	0.7-mm CMS hair		CMS, Chicago Medical Supply, LLC		
Dynamic mechanical detection/allodynia	Q-tip		e.g. Q-Tips Care swabs / cotton buds, pack of 3 (3 x 206 sticks).: Amazon.de: Beauty		
Pressure pain sensitivity	Bedside algometer	10-mL syringe sealed with a plug and felt pad with a contact area of 1 cm ²	BRAUN-4617100V		
VDT	Tuning fork	128 Hz tuning fork with damper to generate a frequency of 64 Hz, 8/8 scale (Rydel-Seiffer)	e.g. <u>medicalax.de - Zellamed Neurological</u> <u>tuning fork by Rydel-Seiffer Tuning Fork</u> <u>Diagnostics</u>		
*Not used in the present study but proposed as an alternative due to better practicability. CDT, cold detection threshold; CPT, cold pain threshold; DMA, dynamic mechanical allodynia; HPT, heat pain threshold; MDT, mechanical detection threshold; MPS, mechanical pain sensitivity; MPT.					

CDT, cold detection threshold; CPT, cold pain threshold; DMA, dynamic mechanical allodynia; HPT, heat pain threshold; MDT, mechanical detection threshold; MPS, mechanical pain sensitivity; MPT, mechanical pain threshold; PHS, paradoxical heat sensation; PPT, pressure pain threshold; TSL, thermal sensory limen; VDT, vibration detection threshold; WDT, warm detection threshold; WUR, wind-up ratio

Supplement Table 2. Sensitivity and specificity of dichotomous bedside-QST parameters as compared to abnormal lab-QST parameters

Lab-QST	Bedside-QST sensation	Positive	Negative	Sensitivity	Specificity	Accuracy
(normal/abnormal)	perceived (ves/no)	predictive	predictive	%	%	%
(normalianterman)		value %	value %	70		<i>,</i> ,,
ZCDT	22°C metal	61.9	76.9	59.1	78.9	71 7
Pathological: <-1.96	Perception (cold	01.0	10.0	00.1	10.0	,
	hypoesthesia)					
		85.7	69.8	27.3	97.4	71 7
Pathological: <-1.96	Perception (cold	00.7	09.0	21.5	57.4	71.7
1 athological. <-1.90	hypoesthesia)					
	az°C motol	25.0	77.2	28.6	72.0	62.2
ZVVDI Pothological: < 1.06	S/ C metal	25.0	11.5	20.0	73.9	03.3
Falliological. <-1.90	hypoosthosia)					
	AF°C motol	25.0	77.1	21.4	90.4	66.7
ZWD1 Rothological: < 1.06	AS C metal	23.0	77.1	21.4	00.4	00.7
Falliological. <-1.90	Perception (warm					
DUC	Nypoestnesia)	00.7	50.0		07.4	<u> </u>
PHS Dethelesisely 1 DUC	PHS 22°C metal Perception	66.7	59.6	8.0	97.1	60.0
Pathological: >1 PHS	(PHS)	75.0	00.7	40.0	07.4	04.7
PHS	PHS 08°C metal Perception	75.0	60.7	12.0	97.1	61.7
Pathological: >1 PHS	(PHS)					
zCPT	22°C metal	n.a.				18.3
Pathological: <-1.96	"loss" (cold hypoalgesia)					
zCPT	08°C metal	n.a.				2 <mark>1.7</mark>
Pathological: <-1.96	"loss" (cold hypoalgesia)					
zCPT	22°C metal	27.3	91.8	42.9	84.9	80.0
Pathological: >+1.96	"gain" (cold hyperalgesia)					
zCPT	08°C metal	<mark>23.1</mark>	<mark>91.5</mark>	42.9	<mark>81.1</mark>	<mark>76.7</mark>
Pathological: >+1.96	"gain" (cold hyperalgesia)					
zHPT	37°C metal	7.1	100.0	100.0	7.1	13.3
Pathological: <-1.96	"loss" (heat hypoalgesia)					
zHPT	45°C metal	11.1	100.0	100.0	42.9	46.7
Pathological: <-1.96	"loss" (heat hypoalgesia)					
zHPT	37°C metal	50.0	89.3	25.0	96.2	86.7
Pathological: >+1.96	"gain" (heat hyperalgesia)					
zHPT	45°C metal	29.2	97.2	87.5	67.3	70.0
Pathological: >+1.96	"gain" (heat hyperalgesia)					
zMDT	Q-tip	65.2	73.0	60.0	77.1	70.0
Pathological: <-1.96	Perception (mechanical					
0	hypoesthesia)					
zMDT	Neurotip	92.3	76.7	54.5	97.1	80.4
Pathological: <-1.96	Perception (mechanical		-		-	
	hypoesthesia)					
zMDT	64mN	81.8	67.3	36.0	94.3	70.0
Pathological: <-1.96	Perception (mechanical	0.110	0110	00.0	0.10	
r allological. Crico	hypoesthesia)					
7MPS	0.7mm CMS "loss"	27.3	100.0	100.0	86.0	86.7
Pathological: <-1.96	Pain (mechanical	27.0	100.0	100.0	00.0	00.7
	hypoalgesia)					
	Nouropon "loss"	25.0	100.0	100.0	92.0	02.0
ZIVIE O Dathological: < 1.06	Pain (mechanical	20.0	100.0	100.0	03.0	03.9
1 alliological. <-1.90						
		16.0	100.0	100.0	21.0	21 7
ZIVIPS		16.3	100.0	100.0	21.2	31.7
Pathological: >+1.96						
	nyperaigesia)	10.0	100.0			
ZMPS	Neuropen "gain"	18.2	100.0	100.0	25.0	35.7
Pathological: >+1.96						

	Pain (mechanical					
	hyperalgesia)					
zMPT	0.7mm CMS "loss"	63.6	98.0	87.5	92.3	91.7
Pathological: <-1.96	Pain (mechanical					
	hypoalgesia)					
zMPT	Neuropen "loss"	58.3	97.7	87.5	89.6	89.3
Pathological: <-1.96	Pain (mechanical					
	hypoalgesia)					
zMPT	0.7mm CMS "gain"	14.3	100.0	100.0	20.8	30.0
Pathological: >+1.96	Pain (mechanical					
	hyperalgesia)					
zMPT	Neuropen "gain"	15.9	100.0	100.0	24.5	33.9
Pathological: >+1.96	Pain (mechanical					
	hyperalgesia)					
zWUR	CMS WUR ratio "loss"	n.a.				1 <mark>3.0</mark>
Pathological: <-1.96	Pain (central sensitization)					
zWUR	Neuropen WUR ratio "loss"	n.a.				2.3
Pathological: <-1.96	Pain (central sensitization)					
zWUR	CMS WUR ratio "gain"	5.0	100.0	100.0	13.6	17.4
Pathological: >+1.96	Pain (central sensitization)					
zWUR	Neuropen WUR ratio "gain"	4.7	100.0	100.0	2.4	6.8
Pathological: >+1.96	Pain (central sensitization)					
logDMA	Q-tip allodynia	46.7	100.0	100.0	84.9	86.7
Pathological: >0.1	Pain					
logDMA	Q-tip postallodynia	30.0	97.5	85.7	73.6	75.0
Pathological: >0.1	sensation					
	Pain					
zPPT	Pressure algometer at 4-	16.7	97.2	100.0	62.5	65.0
Pathological: <-1.96	mL "loss"					
	Pain (deep somatosensory					
	hypoalgesia)					
zPPT	Pressure algometer at 4-	58.3	100.0	95.5	63.2	75.0
Pathological: >+1.96	mL "gain"					
	Pain (deep somatosensory					
	hyperalgesia)					

Bold parameters; parameters with a sensitivity and specificity about $\ge 60\%$ n.a., not applicable