

Table 2 Additional secondary outcome measures of mixed model analysis

<i>Additional secondary outcome measures</i>								
		<i>T0 Baseline</i>	<i>T1 12 weeks</i>	<i>T2 26 weeks</i>	<i>T3 52 weeks</i>	<i>MT vs UC (T1) MD; (95%CI)</i>	<i>MT vs UC (T2) MD; (95%CI)</i>	<i>MT vs UC (T3) MD; (95%CI)</i>
Allodynia score <i>Mean (SD)</i>	<i>MT</i>	2.66 (2.23)	2.34 (2.25)	2.47 (2.26)	2.56 (2.70)	-0.22 (-1.24 to 0.80) <i>p</i> 0.67	0.98 (-0.05 to 2.01) <i>p</i> 0.06	0.81 (-0.22 to 1.85) <i>p</i> 0.12
	<i>UC</i>	3.33 (2.29)	3.15 (2.43)	2.08 (2.10)	2.42 (2.47)			
Neck pain (NPRS) <i>Mean (SD)</i>	<i>MT</i>	5.03 (2.75)	4.43 (2.58)	4.38 (2.69)	4.37 (2.74)	-0.42 (-1.70 to 0.85) <i>p</i> 0.51	0.75 (-0.56 to 2.06) <i>p</i> 0.26	0.38 (-0.94 to 1.70) <i>p</i> 0.58
	<i>UC</i>	6.55 (2.46)	5.58 (2.93)	4.62 (2.84)	5.04 (3.10)			
Muscle Endurance <i>Mean (SD)</i>	<i>MT</i>	23.14 (17.49)	32.0 (22.56)	30.45 (20.51)	31.58 (21.71)	2.77 (-8.50 to 14.05) <i>p</i> 0.63	-4.51 (-15.95 to 6.92) <i>p</i> 0.44	-1.90 (-13.50 to 9.69) <i>p</i> 0.75
	<i>UC</i>	14.45 (10.0)	24.20 (25.23)	29.48 (32.56)	28.05 (28.05)			
PPT Occipital <i>Mean (SD)</i>	<i>MT</i>	2.59 (1.10)	3.20 (1.11)	3.09 (0.96)	3.28 (1.25)	0.60 (0.21 to 0.99) <i>p</i> < 0.01	0.59 (0.19 to 0.98) <i>p</i> < 0.01	0.48 (0.09 to 0.87) <i>p</i> 0.03
	<i>UC</i>	2.65 (1.19)	2.62 (0.90)	2.60 (0.97)	2.78 (1.07)			
PPT Trapezius <i>Mean (SD)</i>	<i>MT</i>	3.31 (1.41)	3.74 (1.30)	3.57 (1.33)	4.03 (1.74)	0.29 (-0.30 to 0.89) <i>p</i> 0.33	0.45 (-0.15 to 1.05) <i>p</i> 0.14	0.22 (-0.38 to 0.82) <i>p</i> 0.47
	<i>UC</i>	3.24 (1.27)	3.35 (1.29)	3.27 (1.37)	3.81 (1.72)			
PPT anterior tibial <i>Mean (SD)</i>	<i>MT</i>	4.03 (1.64)	4.58 (1.76)	4.34 (1.60)	4.55 (1.75)	0.40 (-0.20 to 1.00) <i>p</i> 0.20	0.40 (-0.20 to 1.01) <i>p</i> 0.19	-0.06 (-0.67 to 0.54) <i>p</i> 0.84
	<i>UC</i>	4.07 (1.55)	4.10 (1.34)	4.12 (1.61)	4.67 (1.93)			

Table 2: Observed mean values (SD) at baseline and follow-up and between-group effects (group-by-time

interaction, adjusted for baseline data) for primary and secondary outcomes. MT= manual therapy; UC= usual care; *T0*= baseline; *T1*= follow-up at 12 weeks; *T2*= follow-up at 26 weeks; *T3*= follow-up at 52 weeks; SD= standard deviation; MD= mean difference; CI= confidence interval; freq.= frequency; n= number of participants; NSAIDs=non-steroidal anti-inflammatory drugs; HIT-6= headache impact test; PPTs= pressure pain thresholds; ASC-12= 12-item allodynia checklist; GPE= global perceived effect; # Very much improved score 6/6; much improved score 5/6; Not improved score 3/6.

Additional Figures:

Figure 4

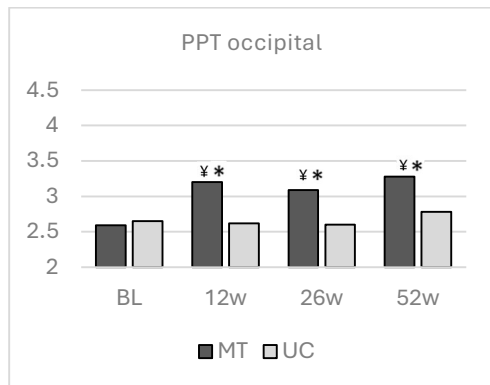


Figure 5

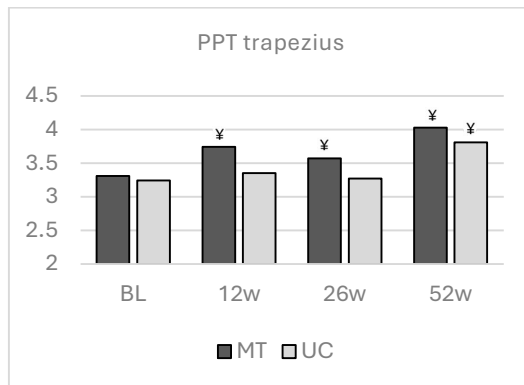


Figure 6

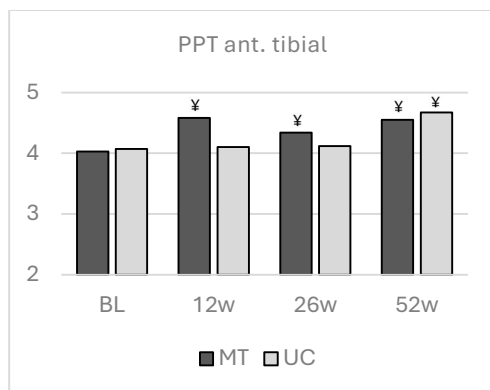


Figure 4-6: Mean PPT values of left and right; PPT= pressure pain threshold; MT= manual therapy; UC= usual care by the General Practitioner; *= significant between-group difference ($p < 0.05$); ¥= significant within-group difference ($p < 0.05$).

Figure 7

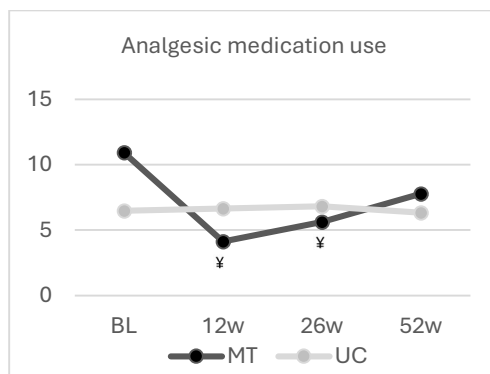


Figure 8

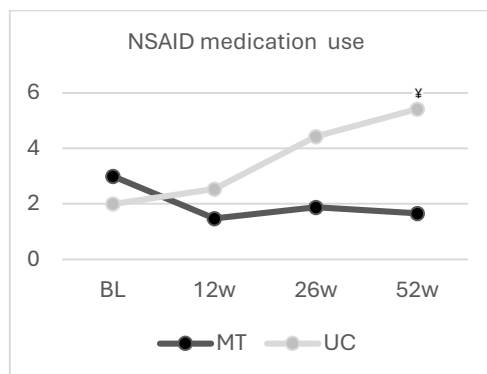


Figure 9

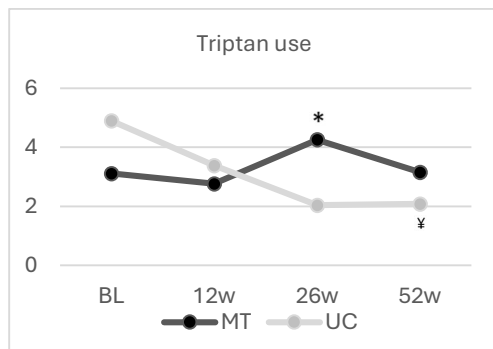


Figure 10

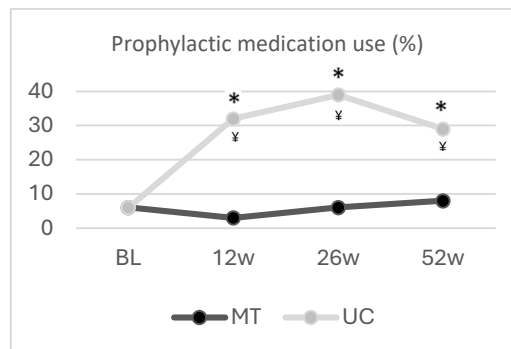


Figure 7-9: Mean medication use in number of pills per 4 weeks.

Figure 10: Proportion of participants using prophylactic medication in %.

MT= manual therapy; UC= usual care by the General Practitioner; *= significant between-group difference ($p < 0.05$); ¥= significant within-group difference ($p < 0.05$).