Duke University School of Medicine Doctor of Physical Therapy

Background

- Low back pain (LBP) is the most common disability among adults
- 80 90% lifetime prevalence
- Leading causes: loss of work productivity and medical care dollars spent
- Traditional pharmaceuticals, treatments: exercises and/or spinal manipulation

Purpose

• Update on effectiveness of Spine Thrust Manipulation (STM) for LBP treatment as a follow-up study to the systematic review authored by Kuczynski et al. (2012)

Methods

- Systematic Review of 11 Randomized Control Trial (RCT) articles compared STM to any other treatment methods
- PRISMA guidelines used to report items
- Data pulled on July 11, 2016 from PubMed, CINAHL, and Embase following Cochrane Collaboration guidelines
- All of the studies published in English
- Articles randomly assigned to two independent authors for review
- Interrater reliability measured using Cohen's Kappa coefficient
- Risk of bias assessed using the Cochrane Risk of Bias tool
- Eligibility criteria:
 - patients with LBP
 - STM performed by a physical therapist
 - control groups not receiving STM
 - standardized outcome measures
- comparator Outcome measures and interventions reported across the entire dataset were gathered in a PICOS table

Effectiveness of Physical Therapy Administered Spinal Manipulation for the Treatment of Low Back Pain: An Updated Systematic Review of the Literature Nermin Gradascevic SPT, Meaghan Reardon SPT, Travis Rhea SPT, Hannah Colopy SPT, Chad Cook, PT, PhD, MBA, FAAOMPT Duke University School of Medicine

Analysis

- Most frequently reported outcome measure(s) identified and **Cohen's d effect size** calculated [Effect d <0 adverse ; 0.0-0.2 no effect; 0.2-0.5 small; 0.5-0.8 intermediate; \geq 0.8 large]
- Meta-analysis: not conducted due to the lack of standardized timeframe in pre- and post-treatment outcome measurements.

Results

• Total of 1,120 subjects participated in the aforementioned RCTs





Pain (NPRS)	STM n=	Mean STM (SD)	Comparator n=	Mean Comparator (SD)	P value	Effect Size (Cohen's d)	Risk of Bias Total
Bialosky et al (2009) [0-100]	12	NR	12	NR	NR	#	LOW
Bialosky et al (2014) [0-10]	28	NR	82	NR	NR	#	LOW
Castro-Sánchez et al (2016) [0-10]	31	4.9 (1.6)	31	4.6 (1.7)	0.925	-0.18	LOW
Cleland et al (2009) [0-10]	37	NR	37	NR	NR	#	LOW
Cook et al (2013) [0-10]	76	1.8 (1.8)	73	1.9 (1.5)	0.66	0.06	LOW
Fritz JM et al (2015) [0-10]	108	1.3 (1.7)	112	1.4 (1.9)	0.44	0.05	UNCLEAR
Hallegraeff et al (2009) [0-100]	31	19.0 (16.9)	33	24.8 (20.1)	0.26	0.31	LOW
Mosheni-Bandpei et al (2006) [0-100]	56	23.4 (29.4)	56	37.9 (28.3)	0.001	0.50	LOW
Perry J, et al (2015) [0-10]	25	NR	25	NR	NR	#	LOW
Venegas-Rios et al (2009) [0-100]	33	41.12 (27.25)	33	46.45 (27.64)	0.433	0.19	UNCLEAR
				Mean			
Disability (ODI)	STM n=	Mean STM (SD)	Comparator n=	Mean Comparator (SD)	P value	Effect Size (Cohen's d)	Risk of Bias Total
Disability (ODI) Bialosky et al (2014) [%]	STM n= 28	Mean STM (SD) NR	Comparator n= 82	Mean Comparator (SD) NR	P value NR	Effect Size (Cohen's d) #	Risk of Bias Total LOW
Disability (ODI) Bialosky et al (2014) [%] Castro-Sánchez et al (2016) [0-50]	STM n= 28 31	Mean STM (SD) NR 24.8 (13)	Comparator n= 82 31	Mean Comparator (SD) NR 28.1 (13.6)	P value NR 0.015	Effect Size (Cohen's d) # 0.25	Risk of Bias Total LOW LOW
Disability (ODI) Bialosky et al (2014) [%] Castro-Sánchez et al (2016) [0-50] Childs et al (2004) [%]	STM n= 28 31 70	Mean STM (SD) NR 24.8 (13) NR	Comparator n= 82 31 61	Mean Comparator (SD) NR 28.1 (13.6) NR	P value NR 0.015 NR	Effect Size (Cohen's d) # 0.25 #	Risk of Bias Total LOW LOW HIGH
Disability (ODI) Bialosky et al (2014) [%] Castro-Sánchez et al (2016) [0-50] Childs et al (2004) [%] Cleland et al (2009) [%]	STM n= 28 31 70 37	Mean STM (SD) NR 24.8 (13) NR NR	Comparator n= 82 31 61 37	Mean Comparator (SD)NR28.1 (13.6)NRNR	P value NR 0.015 NR NR	Effect Size (Cohen's d) # 0.25 # #	Risk of Bias Total LOW LOW HIGH LOW
Disability (ODI) Bialosky et al (2014) [%] Castro-Sánchez et al (2016) [0-50] Childs et al (2004) [%] Cleland et al (2009) [%] Cook et al (2013) [%]	STM n= 28 31 70 37 76	Mean STM (SD) NR 24.8 (13) NR NR 14.9 (13.9)	Comparator n= 82 31 61 37 37	Mean Comparator (SD)NR28.1 (13.6)NRNR17.2 (13.1)	P valueNR0.015NRNR0.31	Effect Size (Cohen's d) # 0.25 # 4 0.17	Risk of Bias Total LOW LOW HIGH LOW
Disability (ODI) Bialosky et al (2014) [%] Castro-Sánchez et al (2016) [0-50] Childs et al (2004) [%] Cleland et al (2009) [%] Cook et al (2013) [%] Fritz JM et al (2015) [%]	STM n= 28 31 70 37 76 108	Mean STM (SD) NR 24.8 (13) NR 14.9 (13.9) 7.0 (11.4)	Comparator n= 82 31 61 61 37 73 73 112	Mean Comparator (SD) NR 28.1 (13.6) NR 17.2 (13.1) 9.0 (11.6)	P value NR 0.015 NR 0.31 0.19	Effect Size (Cohen's d) # 0.25 # 4 0.17 0.17	Risk of Bias TotalLOWLOWHIGHLOWUNCLEAR
Disability (ODI) Bialosky et al (2014) [%] Castro-Sánchez et al (2016) [0-50] Childs et al (2004) [%] Cleland et al (2009) [%] Cook et al (2013) [%] Fritz JM et al (2015) [%] Hallegraeff et al (2009) [0-50]	STM n= 28 31 70 37 37 76 108 31	Mean STM (SD) NR 24.8 (13) NR 14.9 (13.9) 7.0 (11.4) 14 (17)	Comparator n= 82 31 61 61 37 73 73 112 33	Mean Comparator (SD) NR 28.1 (13.6) NR 17.2 (13.1) 9.0 (11.6) 14(12)	P value NR 0.015 NR 0.31 0.19 0.38	Effect Size (Cohen's d) # 0.25 # 0.25 0.17 0.17 0.17	Risk of Bias Total LOW LOW HIGH LOW LOW
Disability (ODI) Bialosky et al (2014) [%] Castro-Sánchez et al (2016) [0-50] Childs et al (2004) [%] Cleland et al (2009) [%] Cook et al (2013) [%] Fritz JM et al (2015) [%] Hallegraeff et al (2009) [0-50]	STM n= 28 31 70 37 76 108 31 31	Mean STM (SD) NR 24.8 (13) NR NR 14.9 (13.9) 7.0 (11.4) 14.9 (13.9) 12.9 (14.9)	Comparator n= 82 31 61 61 37 73 73 112 33 33	Mean Comparator (SD) NR 28.1 (13.6) NR 17.2 (13.1) 9.0 (11.6) 14(12) 22.1 (14.7)	P value NR 0.015 NR 0.31 0.31 0.19 0.38 0.001	Effect Size (Cohen's d)	Risk of Bias TotalLOWLOWHIGHLOWLOWLOWLOWLOW
Disability (ODI) Bialosky et al (2014) [%] Castro-Sánchez et al (2016) [0-50] Childs et al (2004) [%] Cleland et al (2009) [%] Cook et al (2013) [%] Fritz JM et al (2015) [%] Hallegraeff et al (2009) [0-50] Mosheni-Bandpei et al (2006) [%] Perry J, et al (2015) [%]	STM n= 28 31 70 37 76 108 31 31 56 25	Mean STM (SD) NR 24.8 (13) NR NR 14.9 (13.9) 7.0 (11.4) 12.9 (14.9) NR	Comparator n= 31 31 61 37 37 73 73 112 33 56 56	Mean Comparator (SD) NR 28.1 (13.6) NR 17.2 (13.1) 9.0 (11.6) 14(12) 22.1 (14.7) NR	P value NR 0.015 NR 0.31 0.31 0.19 0.38 0.001 NR	Effect Size (Cohen's d)	Risk of Bias TotalLOWLOWHIGHLOWLOWLOWLOWLOWLOW

- High risk
- Comparator interventions identified in PICOS: bicycle ultrasound, and non-thrust manipulation
- Resulting means and standard deviations used for intervention and control group
- The majority of the studies reported no to small effect sizes in favor of STM over the comparator

Conclusions

- Overall findings: "no" to "small" effect size in contrast to Kuczynski et al. findings
- terms of efficiency in LBP patients
- Aside from one study (Bialosky et al 2014), no true nature of their papers
- No clear evidence in clinical practice for using STM over comparator interventions

Clinical Relevance

STM vs comparator interventions: both are safe and equally effective to use for LBP treatment treatment of patients with LBP.

Acknowledgements / References

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*References are available upon request

Results

Cochrane Risk of Bias: 7 Low risk, 3 Unclear risk, & 1

cardiovascular exercise, low back extension, AROM,

Oswestry Disability Index (ODI) and **Numeric Pain** Rating Scale (NPRS) were the most commonly used outcome measures within studies identified (90.9%)

Cohen's d effect size to measure the effect between

No consistent conclusion on any meaningful differences between STM and the comparators in

control groups were used which limits the definitive

Overall, patient preference should be highly considered when selecting an intervention for the