Evidence Tables and Data Extraction of Randomized Trials

Canadian Chiropractic Guideline Initiative (CCGI) Centre for Disability Prevention and Rehabilitation

Systematic Review Workshop 2019







CCGI workshops

- 1. Introduction to research questions and the PICO framework
- 2. Systematic review screening of literature
- 3. Critical appraisal/risk of bias assessment of RCTs
- 4. Evidence tables and data extraction of RCTs







Outline of workshop

1. Introduction to evidence tables



- Exercise
- 2. Introduction to data extraction of RCTs to build evidence tables
 - Exercise







Learning objectives

At the end of this session, you should be able to:

- Describe the purpose of an evidence table
- Identify key characteristics of studies when reading an evidence table
- Extract data from RCTs to build evidence tables







Clinical/educational scenario





Patient

Healthcare provider





Have you used evidence tables before? If yes, how?

If not, what do you think might be involved?







Introduction to the Evidence Table



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What is an evidence table?

- A summary of the **most important information** from included studies
- Concisely summarizes evidence in a standard format
- Includes outcomes (benefits and harms) and information on the

setting and context of the study









What would you consider as key information from an RCT?







PICO framework

- Helps guide the development of clear research questions
- Helpful for questions related to treatment effectiveness

Ρ	Population of interest
I	Intervention you want to know the effectiveness of
С	Comparison – what the intervention is being compared to
0	Outcome(s) you want to learn about





PICO framework

Ρ	Population	Population of interest	E.g., Headaches, flu
I	Intervention/ exposure	Treatment or exposure level of participants	E.g., Exercise, surgery
С	Comparison	Reference group used to compare with intervention/exposure	E.g., Injections, placebo, no treatment
0	Outcome	Measure used to examine effects of intervention/exposure	E.g., Pain, quality of life





Additional details of PICO

• Population:

- Disease or condition; stage, severity
- Demographic characteristics (e.g., age, gender)

• Intervention:

- Type of intervention
- Dose, duration, timing, route, etc.

• Comparison:

- Treatment interventions
- Placebo/sham, waiting list, no intervention

• Outcome:

- Benefit or harm; mean difference, frequency, time to event, etc.
- Type: mortality, pain, quality of life, disability, etc.





What is an evidence table?



A snapshot of key information from studies.....written in a table format







Evidence table headings





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Example of evidence table

	Population	Intervention	Comparison		Outcome/time	
Authors, year	Subjects and setting, number (n) enrolled	Interventions, number (n) of subjects	Comparisons, number (n) of subjects	Follow-up	Outcomes	Key findings
Griffiths et al., 2009 [44]	Patients (≥18 y.o.) referred for outpatient physical therapy in the United Kingdom Case definition: chronic neck pain (≥3 mo) n=74	Specific exercise: up to four sessions/6 wk by physical therapists. Active range of motion, posture correction techniques, and neck stabilization/isometric exercises Advice to perform exercises at home 5–10 times daily n=37	General exercise: up to four sessions for 6 wk by physical therapists. Active range-of-motion exercises and posture correction techniques Advice to perform exercises at home 5–10 times daily n=37	6 wk, 6 mo	Primary outcome: disability (NPDS) Secondary outcomes: disability (NPQ), pain affect (NRS), severity of patient-identified worst problem (NRS), medication use (48-h recall), global improvement, cointervention, and health- related quality of life (SF- 36)	 Difference in mean change (specific exercise–general exercise) NPDS 6 wk: -0.15 (95% C -6.46 to 6.16)* NPDS 6 mo: 6.46 (95% CI -0.81 to 13.73)* No significant difference in secondary outcomes excep for medication use (genera exercise–specific exercise) Percent reporting medication use: 6 wk: 0.29 (95% CI

0.10-0.84), 6 mo: 1.16 (95% CI 0.37-3.59)









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Adapted from Fletcher et al. Clinical Epidemiology 5th Edition



Utility of an evidence table



- Summarizes large volumes of information
- May not need to read original study
- A summary of <u>multiple</u> studies
- Often a summary of <u>high quality</u> studies







Introduction to Data Extraction



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Systematic review process





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Data extraction process

- Data extraction items are listed in systematic review protocol
- One author extracts data from high quality studies to build evidence table
- Second reviewer independently checks data









Prior to data extraction

Review the critical appraisal consensus document:

- Which data are relevant for this review?
- Any limitations that modify the data to be extracted?
- Are calculations indicated/possible?
 - Calculations will not be covered in this

workshop





Examples:

Which interventions are

relevant for this review?

• Which outcomes/follow-

up periods were accepted?



Key items to be extracted





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Example for evidence table

Journal of Physiotherapy 63 (2017) 144-153



Journal of PHYSIOTHERAPY

journal homepage: www.elsevier.com/locate/jphys

Research

Early rehabilitation after lumbar disc surgery is not effective or cost-effective compared to no referral: a randomised trial and economic evaluation

Teddy Oosterhuis ^{a,b}, Raymond W Ostelo ^{a,b,c}, Johanna M van Dongen ^{a,b}, Wilco C Peul ^{d,e}, Michiel R de Boer ^{a,b}, Judith E Bosmans ^{a,b}, Carmen L Vleggeert-Lankamp ^d, Mark P Arts ^e, Maurits W van Tulder ^{a,b}





Example for evidence table

Research question of study by Oosterhuis et al:

- Is referral for early rehabilitation after lumbar disc surgery effective (and cost-effective) compared to no referral?
 - **P** Adults with herniated lumbar disc and signs of nerve root compression
 - I Early rehabilitation after lumbar disc surgery
 - **C** No referral for early rehabilitation after lumbar disc surgery
 - **O** Functional status; leg and back pain; global perceived recovery; general physical and mental health (SF12); at 3, 6, 9, 12 and 26 weeks





Column 1: Author(s), Year

Author(s), Year

Oosterhuis et al., 2017 [1]







Column 2: Subjects and Setting

Brief description of:		For study by Griffiths et al:
1.	Participants (e.g., adults 18+ y.o.)	Patients (≥18 y.o.) referred
2.	Health care setting of study	for outpatient physical
3.	Region where study took place	therapy in the United Kingdom
4.	Case definition	Case definition: chronic neck
5.	# of subjects enrolled in study	pain ($\geq 3 \text{ mo}$) n=74





Column 2: Subjects and Setting

Bri	ef description of:	For study by Oosterhuis et al:
1.	Participants (e.g., adults 18+ y.o.)	
2.	Health care setting of study	
3.	Region where study took place	
4.	Case definition	
5.	# of subjects enrolled in study	





Column 2: Subjects and Setting

Subjects and Setting; Number (n) Enrolled

Patients (18-70 y.o.) from 10 peripheral hospitals in urban or regional areas of three regions in the Netherlands.

Case definition: herniated lumbar disc confirmed by MRI and signs of nerve root compression

n=184







Column 3: Intervention

Brief description of:		For study by (Griffiths et al:
1.	Brief name of treatment arm	Specif	ic exercise: up to four
2.	Treatment frequency/duration	sess	apists. Active range of
3.	Health care provider	mot tech	ion, posture correction niques, and neck
4.	Description of what types of treatment were provided	stab exer Advice	ilization/isometric reises e to perform exercises
5.	Number of subjects in group	at h n=37	ome 5–10 times daily





Column 3: Intervention

Brief description of:		For study by Oosterhuis et al:
1.	Brief name of treatment arm	
2.	Treatment frequency/duration	
3.	Health care provider	
4.	Description of what types of treatment were provided	
5.	Number of subjects in group	





Column 4: Comparison

Brief description of:		For study by Griffiths et al:	
1.	Brief name of treatment arm	Conomi avanica, un to four	
2.	Treatment frequency/duration	sessions for 6 wk by	
3.	Health care provider	physical therapists. Active range-of-motion exercises	
4.	Description of what types of treatment were provided	and posture correction techniques Advice to perform exercises	
5.	Number of subjects in group	at home 5–10 times daily $n=37$	





Column 4: Comparison

Brief description of:		For study by Oosterhuis et al:
1.	Brief name of treatment arm	
2.	Treatment frequency/duration	
3.	Health care provider	
4.	Description of what types of	
	treatment were provided	
5.	Number of subjects in group	





Column 3-4: Interventions and Comparisons

Interventions; Number (n) of Subjects	Comparisons; Number (n) of Subjects
Referral for early rehabilitation following	No referral for early rehabilitation
<u>lumbar disc surgery:</u>	following lumbar disc surgery:
Postoperative exercise therapy in primary	Not referred for rehab after discharge
care starting the first week after	from the hospital. Participants could
received one or two individual. face-to-	consult their neurosurgeon or GP. Were
face, exercise therapy sessions of 30	requested to refrain from exercise
minutes per week. n=92	therapy or other allied health
	interventions in the 6- to 8- week study
	period. n=77





Column 5: Follow-up

Brief description of:		For stud	y by Griffiths et al:	
1.	Follow-up periods after intervention/treatment was completed		6 wk, 6 mo	





Column 5: Follow-up

Brief description of:		For study by Oosterhuis et al:
1.	Follow-up periods after intervention/treatment was completed	







Column 5: Follow-up

• List follow-up periods of the study that will be reported

Follow-up

3, 6, 9, 12, and 26 weeks following surgery







Column 6: Outcomes

Brief description of:	For study by Griffiths et al:		
Outcomes in the following format:1. Follow each outcome with outcome assessment method in brackets	Primary outcome: disability (NPDS) Secondary outcomes: disability (NPQ), pain affect (NRS), severity of		
2. Outcomes should be separated using semicolons	problem (NRS), medication use (48-h recall), global		
3. Adverse events (if assessed)	improvement, cointervention, and health- related quality of life (SF- 36)		





Column 6: Outcomes

Brief description of:	For study by Oosterhuis et al:
Outcomes in the following format:	
1. Follow each outcome with outcome assessment method in brackets	
2. Outcomes should be separated using semicolons	
3. Adverse events (if assessed)	







Column 6: Outcomes

Outcomes

Primary Outcomes:

- Functional Status (Oswestry Disability Index version 2.1a)
- Average Pain Intensity over the preceding week for leg pain and low back pain (11-point NRS)
- Global perceived effect (7-point Global Perceived Effect Scale)
- General physical and mental health (Medical Outcome Study Short Form 12)



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Column 7: Key findings

Brief description of:		For study by Griffiths et al:		
1.	Main results of study summarized by follow-up period		Difference in mean change (specific exercise–general exercise)	
2.	Effect sizes and 95% CI should be provided when possible		NPDS 6 wk: -0.15 (95% CI -6.46 to 6.16)* NPDS 6 mo: 6.46 (95% CI -0.81 to 13.73)*	
3.	We are interested in between group differences		No significant difference in secondary outcomes except for medication use (general exercise-specific exercise)	
4.	May require calculations		Percent reporting medication use: 6 wk: 0.29 (95% CI 0.10–0.84), 6 mo: 1.16 (95% CI 0.37–3.59)	





S CMCC

Column 7: Key findings

Brie	ef description of:	For study by Oosterhuis et al:
1.	Main results of study summarized by follow-up period	
2.	Effect sizes and 95% CI should be provided when possible	
3.	We are interested in between group differences	
4.	May require calculations	





Column 7: Key findings

Key findings

Mean difference (95% CI):

Functional Status (ODI, 0-100):

• Crude 1.0 (95% CI -3.7 to 5.7); adjusted 1.5 (95% CI -3.6 to 6.7)

Pain Intensity Leg (NRS, 0-10)*

• Crude –0.1 (95% CI –0.8 to 0.6); adjusted 0.1 (95% CI –0.7 to 0.8)

Pain Intensity Back (NRS, 0-10)

Crude 0.3 (95% CI –0.3 to 0.9); adjusted 0.3 (95% CI –0.3 to 0.9)







Column 7: Key findings Cont'd

Key findings

Global Perceived Effect (n (%) recovered):

• OR 1.0 (0.6 to 1.7)

General Physical Health (SF12, 0-100):

• Crude -1.1 (95% CI -8.5 to 6.3); adjusted -3.5 (95% CI -11.3 to 4.3)

General Mental Health (SF12, 0-100):

• Crude –0.9 (95% CI –6.8 to 5.0); adjusted –4.1 (95% CI –9.4 to 1.3)





Completed evidence table for Oosterhuis et al

Author(s), Year	Subjects and Setting; Number (n) Enrolled	Interventions; Number (n) of Subjects	Comparisons; Number (n) of Subjects	Follow-up	Outcomes	Key findings
Ooster- nuis et al., 2017 [1]	Patients (18-70 y.o.) from 10 peripheral hospitals in urban or regional areas of three regions in the Netherlands.Referral for early rehabilitation following lumbar disc surgery: Postoperative exercise therapy in prima starting the first received one or individual, face- face, exercise th sessions of 30 m per week. n=92Case definition: herniated lumbar disc confirmed by MRI and signs of nerve root compressionReferral for early rehabilitation following lumbar disc surgery: 	No referral for early rehabilitation following lumbar disc	3, 6, 9, 12, and 26 weeks	Primary Outcomes: -Functional Status (Oswestry Disability Index version 2.1a)	 Mean difference (95% Cl): Functional Status (ODI, 0-100): Crude 1.0 (-3.7 to 5.7); adjusted 1.5 (-3.6 to 6.7) 	
		therapy in prima starting the first after discharge. 6-8 weeks, parti received one or individual, face- face, exercise th sessions of 30 m	xt steps: Extract from all high quality studies To be checked by second reviewer			 in Intensity Leg (NRS, 0-10)* Crude -0.1 (-0.8 to 0.6); adjusted 0.1 (-0.7 to 0.8) in Intensity Back (NRS, 0-10) Crude 0.3 (-0.3 to 0.9); adjusted 0.3 (-0.3 to 0.9) obal Perceived Effect (n (%) recovered):
		refrain from exercise therapy or other allied health interventions in the 6- to 8- week study period. n=77		mental health (Medical Outcome Study Short Form 12)	 OR 1.0 (0.6 to 1.7) General Physical Health (SF12, 0-100): Crude -1.1 (-8.5 to 6.3); adjusted -3.5 (-11.3 to 4.3) General Mental Health (SF12, 0-100): Crude -0.9 (-6.8 to 5.0); adjusted -4.1 (-9.4 to 1.3) 	





Author(s), Year	Subjects and Setting; Number (n) Enrolled	Interventions; Number (n) of Subjects	Comparisons; Number (n) of Subjects	Follow- up	Outcomes	Key findings
Bronfort et	Residents	Spinal	Home exercise with	2, 4, 8,	Primary outcome:	Statistically sig. diff. in mean (SMT – HEA):
al., 2012	from	manipulative	advice (HEA) by physical	12, 26,	neck pain (NRS)	Satisfaction score: (0 to 12 weeks): 0.33 (95% CI 0.11; 0.56), (0 to 52
[2]	Minnesota	therapy (SMT) by	therapists with in-person	and 52		weeks): 0.32 (95% Cl 0.11; 0.54)
	(18-65 y.o.).	chiropractors (12	instruction (2 1-hour	weeks	Secondary outcomes:	No statistically sig. diff. between groups for mean change in neck
		weeks):	sessions with daily home		disability (NDI); global	pain, disability, medication use, physical or mental health-related
	Case	manipulation and	exercise): individualized		improvement;	quality of life or ranges of motion.
	definition:	mobilization, soft-	program of neck and		medication use	No statistically sig. diff. in mean global improvement
	acute/sub-	tissue massage,	shoulder self-		(days/week);	
	acute neck	assisted stretching,	mobilization; education		satisfaction with care;	Statistically sig. diff. in mean change (HEA – medication):
	pain grades	hot and cold packs,	and advice regarding		health-related quality	Neck pain: 26 weeks: 0.69 (95% CI 0.10; 1.28).
	I/III (2-12	and advice to stay	posture and daily		of life (SF-36); cervical	Disability: 26 weeks: 2.95 (95% CI 0.37; 5.53).
	weeks) and	active or modify	activities. (n=90)		spine range of motion	Medication use: 26 weeks: 1.49 (95% CI 0.78; 2.20),
	neck pain	activity as needed.			(CA 6000 Spine	52 weeks: 1.00 (95% CI 0.27; 1.73).
	intensity ≥	(n=91)	Medication by physician:		Motion Analyzer)	Physical SF-36: 26 weeks: 2.28 (95% CI 0.63; 3.93), 52 weeks: 2.24
	3/10.		NSAIDs, acetaminophen,			(95% CI 0.54; 3.93)
			(narcotics and, muscle			Flexion-extension: 4 weeks: 4.25 (95% CI 1.39; 7.11), 12 weeks: 3.51
	(n=272)		relaxants if necessary);			(95% CI 0.62; 6.40)
			advice to stay active or			
			modify activity. (n=91)			Statistically sig. diff. in mean (HEA – medication):
						Global improvement: (0 to 12 weeks): 0.30 (95% CI 0.01; 0.58), (0 to
						52 weeks): 0.28 (95% Cl 0.01; 0.56)
						Satisfaction score: (0 to 12 weeks): 0.36 (95% CI 0.13; 0.58), (0 to 52
						weeks): 0.38 (95% CI 0.16; 0.59)
						No statistically sig. diff. between groups for mean change in mental
			RACTIC CARE			health-related QOL.

Return to clinical/educational scenario



Patient

Evidence table can provide:

- Key information from high quality studies
- More detailed information of intervention





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Healthcare provider

Summary

- A evidence table concisely summarizes key evidence from included studies
- Includes benefits, harms, setting, and context of the study
- A standardized process is used in systematic reviews to extract key
 - information from studies (including a second check of the data)





Resources

PRISMA Statement (reporting for systematic reviews): <u>http://www.prisma-statement.org/PRISMAStatement/</u>

PRISMA-P Checklist (reporting for systematic review protocols): <u>http://www.prisma-statement.org/Extensions/Protocols.aspx</u>







Learning Objectives

At the end of this session, you should be able to:

- Describe the purpose of an evidence table
- Identify key characteristics of studies when reading an evidence table
- Extract data from RCTs to build evidence tables







Thank You



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