



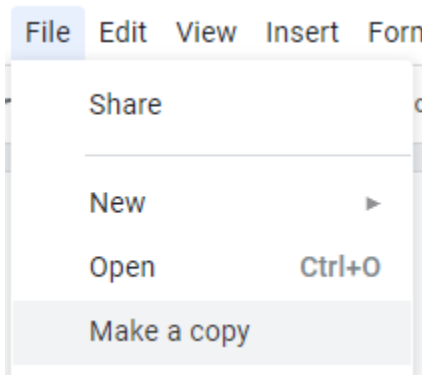
NERDCAT-SRMA

A clinician's guide to appraising systematic reviews and meta-analyses

This checklist is intended to be complemented by the NERDCAT Pressbook which can be found [here](#). Each section heading in this checklist will also link to the corresponding NERDCAT section (see both the [Generalizability](#) and [Systematic reviews and meta-analyses](#) sections). The Pressbook provides insight into how to answer each question, why each question is important, as well as illustrative examples of the concepts involved.

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Article title:

CLINICAL QUESTION ADDRESSED BY THIS REVIEW

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P Age, condition, setting, etc.	
I Drug, dose, duration, etc.	
C	
O Clinical outcomes assessed, etc.	
Trial Inclusion and Exclusion Criteria	

(1) GENERALIZABILITY – DO THESE RESULTS (NOT) APPLY TO MY PATIENTS?

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How does my practice setting differ from that in the trials?	
How do my patients differ from those included in the trial?	
How do the trial interventions differ from those available in my practice?	
Are the trial outcomes clinically important?	
Do the included trials reflect my patients' risk of adverse events? What differences exist?	
Was each element of PICO (i.e. patient, intervention, comparator, and outcome) sufficiently reported to assess generalizability?	
Do the differences above impede the generalizability of the study findings to my practice?	

(10) SEARCH

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Databases of published literature Were a reasonable number of relevant databases searched?	<u>Question</u>	Yes	No
	MEDLINE/PubMed		
	CENTRAL		
	EMBASE		
	Other relevant topic-specific databases:		
Timeframe When was the search conducted? Is it likely there have been subsequent publications that may alter the results?	Search timeframe:		
Grey literature Was a sufficient effort made to find unpublished studies (or unreported results of published studies)?	<u>Clinical trial registries</u>		
	<u>Searched?</u>	Yes	No
	WHO International Clinical Trials Registry (includes clinicaltrials.gov & many more)		
	ClinicalTrials.gov		
	International Federation of Pharmaceutical Manufacturers and Associations [IFPMA] Clinical Trials Portal (pharmaceutical industry-sponsored trials)		
	Others:		
	<u>Regulatory body websites</u>		
	<u>Searched?</u>	Yes	No
	drugs@FDA		
	EMA		
	Others:		
Poster presentations, conference proceedings, or abstracts:			
Additional measures for comprehensiveness Were sources of additional published/unpublished data sought out?	<u>Question</u>	Yes	No
	Handsearch of bibliographies?		
	Contacted study authors?		
	Language restrictions (e.g. English-only)?		
	Others:		

(11) RESULTS OF THE SYSTEMATIC REVIEW (i.e. which trials were included?)

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Eligible trials <ul style="list-style-type: none">• Do all inclusions & exclusions of trials make sense?• Are you aware of any relevant studies that were not identified/included in this review?	Inclusions/exclusions adequately described (e.g. PRISMA flowchart)?	
	Irrational eligibility criteria?	
	Missing trial(s)?	
Risk of bias within trials (trial internal validity) <ul style="list-style-type: none">• Did reviewers adequately assess individual trials for risk of bias?• Was each component reported separately, or summarized with a composite quality score?	RCTs: Cochrane RoB 1 or 2 used?	
	Observational Studies: ROBINS-I used?	
	Other risk of bias assessment used?	
	“Quality control”: Do you agree with the risk of bias assessment reported for the largest weighted trial?	
Clinical and methodological heterogeneity <p>Are there any differences between studies that should preclude meta-analysis?</p>		

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[illegible]