



CASP Checklist: 12 questions to help you make sense of a Cohort Study

**How to use this appraisal tool:** Three broad issues need to be considered when appraising a cohort study:

Are the results of the study valid? (Section A)

What are the results? (Section B)

Will the results help locally? (Section C)

The 12 questions on the following pages are designed to help you think about these issues systematically. The first two questions are screening questions and can be answered quickly. If the answer to both is "yes", it is worth proceeding with the remaining questions. There is some degree of overlap between the questions, you are asked to record a "yes", "no" or "can't tell" to most of the questions. A number of italicised prompts are given after each question. These are designed to remind you why the question is important. Record your reasons for your answers in the spaces provided.

**About:** These checklists were designed to be used as educational pedagogic tools, as part of a workshop setting, therefore we do not suggest a scoring system. The core CASP checklists (randomised controlled trial & systematic review) were based on JAMA 'Users' guides to the medical literature 1994 (adapted from Guyatt GH, Sackett DL, and Cook DJ), and piloted with health care practitioners.

For each new checklist, a group of experts were assembled to develop and pilot the checklist and the workshop format with which it would be used. Over the years overall adjustments have been made to the format, but a recent survey of checklist users reiterated that the basic format continues to be useful and appropriate.

**Referencing:** we recommend using the Harvard style citation, i.e.: *Critical Appraisal Skills Programme (2018). CASP (insert name of checklist i.e. Cohort Study) Checklist. [online] Available at: URL. Accessed: Date Accessed.* 

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Section A: Are the results of the study valid?			
Did the study address a clearly focused issue?	Yes  Can't Tell  No	HINT: A question can be 'focused' in terms of the population studied the risk factors studied is it clear whether the study tried to detect a beneficial or harmful effect the outcomes considered	
Comments:			
2. Was the cohort recruited in an acceptable way?	Yes  Can't Tell  No	HINT: Look for selection bias which might compromise the generalisability of the findings:  • was the cohort representative of a defined population  • was there something special about the cohort  • was everybody included who should have been	
Is it worth continuing?			



HINT: Look for measurement or classification bias: id they use subjective or objective measurements truly reflect what you want them to (have they been validated)  • has a reliable system been hed for detecting all the cases (for measuring disease occurrence)  • were the measurement hods similar in the different groups  • were the subjects and/or the outcome assessor blinded to exposure (does this matter)



5. (a) Have the authors identified all important confounding factors?	Yes  Can't Tell  No	HINT:  • list the ones you think might be important, and ones the author missed
Comments:		
5. (b) Have they taken account of the confounding factors in the design and/or analysis?	Yes Can't Tell No	HINT:  • look for restriction in design, and techniques e.g. modelling, stratified-, regression-, or sensitivity analysis to correct, control or adjust for confounding factors
Comments:		
6. (a) Was the follow up of subjects complete enough?	Yes  Can't Tell  No	HINT: Consider  • the good or bad effects should have had long enough to reveal themselves  • the persons that are lost to follow-up may have different outcomes than those available for assessment  • in an open or dynamic cohort, was there anything special about the outcome of the people leaving, or the exposure of the people entering the
6. (b) Was the follow up of subjects long enough?	Yes Can't Tell No	cohort



Comments:	
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Section B: What are the results?	
7. What are the results of this study?	HINT: Consider
,	<ul> <li>what are the bottom line</li> </ul>
	results
	<ul> <li>have they reported the rate or</li> </ul>
	the proportion between the
	exposed/unexposed, the
	ratio/rate difference <ul><li>how strong is the association</li></ul>
	between exposure and
	outcome (RR)
	<ul> <li>what is the absolute risk</li> </ul>
	reduction (ARR)
Comments:	
8. How precise are the results?	HINT:
	<ul> <li>look for the range of the confidence</li> </ul>
	intervals, if given
Comments:	
comments:	



9. Do you believe the results?	Yes Can't Tell No	HINT: Consider  • big effect is hard to ignore  • can it be due to bias, chance or confounding  • are the design and methods of this study sufficiently flawed to make the results unreliable  • Bradford Hills criteria (e.g. time sequence, dose-response gradient, biological plausibility, consistency)
Comments:		
Section C: Will the results help locally	?	
10. Can the results be applied to the local population?	Yes  Can't Tell  No	HINT: Consider whether  • a cohort study was the appropriate method to answer this question  • the subjects covered in this study could be sufficiently different from your population to cause concern  • your local setting is likely to differ much from that of the study  • you can quantify the local benefits and harms
Comments:		
11. Do the results of this study fit with other available evidence?	Yes Can't Tell No	
Comments:		



12. What are the implications of	Yes	HINT: Consider
this study for practice?		<ul> <li>one observational study rarely</li> </ul>
	Can't Tell	provides sufficiently robust
		evidence to recommend changes
		to clinical practice or within health
	No	
		<ul> <li>for certain questions,</li> </ul>
	l	observational studies provide the
		only evidence
		• recommendations from
		observational studies are always
		stronger when supported by other
		evidence
Comments:		