

Project/Topic of your Clinical Question: _____
Reviewer: _____ Today's Date: _____ Final Evidence Level: _____
Article Title: _____
Year: _____ First Author: _____ Journal: _____

Do the study purpose/objectives and inclusion/exclusion criteria assist in answering your clinical question?

Yes No Unknown

- Study Purpose/Objective:
- Inclusion Criteria:
- Exclusion Criteria:

If you are uncertain of your skills in evidence evaluation, please consult a local evidence expert for assistance:

CCHMC Evidence Experts: <http://groups/ce/NewEBC/EBDMHelp.htm>

Unfamiliar terms can be found in the LEGEND Glossary: <http://groups/ce/NewEBC/EBCFiles/GLOSSARY-EBDM.pdf>

SCOPE AND PURPOSE

1. Were the overall objective(s) of the recommendation specifically described? Yes No Unknown

Comments:

2. Were the health question(s) covered by the recommendation specifically described? Yes No Unknown

Comments:

3. Was the population (*patients, public, etc.*) to whom the recommendation is meant to apply specifically described? Yes No Unknown

Comments:

STAKEHOLDER INVOLVEMENT

4. Did the guideline development group include individuals from all the relevant professional groups? Yes No Unknown

Comments:

5. Were the views and preferences of the target population (*patients, public, etc.*) sought? Yes No Unknown

Comments:

6. Were the target user(s) of the guideline clearly defined? Yes No Unknown
Comments:

RIGOR OF DEVELOPMENT

7. Were systematic methods used to search for evidence? Yes No Unknown
Comments:

8. Were the criteria for selecting the evidence clearly described? Yes No Unknown
Comments:

9. Were the strengths and limitations of the body of evidence clearly described? Yes No Unknown
Comments:

10. Were the methods used for formulating the recommendations clearly described? Yes No Unknown
Comments:

11. Were the health benefits, side effects, and risks considered in formulating recommendations? Yes No Unknown
Comments:

12. Was there an explicit link between the recommendations and the supporting evidence? Yes No Unknown
Comments:

13. Was the guideline externally reviewed by experts prior to its publication? Yes No Unknown
Comments:

14. Was a procedure for updating the guideline provided? Yes No Unknown
Comments:

CLARITY AND PRESENTATION

15. Were the recommendations specific and unambiguous? Yes No Unknown
Comments:

16. Were the different options for management of the condition or health issue clearly presented? Yes No Unknown

Comments:

17. Were key recommendations easily identifiable? Yes No Unknown

Comments:

APPLICABILITY

18. Did the guideline describe facilitators and barriers to its application? Yes No Unknown

Comments:

19. Did the guideline provide advice and/or tools on how the recommendations can be put into practice? Yes No Unknown

Comments:

20. Were the potential resource implications of applying the recommendations considered? Yes No Unknown

Comments:

21. Did the guideline present monitoring and/or auditing criteria? Yes No Unknown

Comments:

EDITORIAL INDEPENDENCE

22. Was the content of the guideline free from any influence of views of the funding body? Yes No Unknown

Comments:

23. Were competing interests of guideline development group members recorded and addressed? Yes No Unknown

Comments:

24. Would you include this guideline in development of a care recommendation? Yes No Unknown

Comments:

ADDITIONAL COMMENTS OR CONCLUSIONS (“TAKE-HOME POINTS”):

QUALITY LEVEL / EVIDENCE LEVEL

- Consider each “No” answer and the degree to which this limitation is a threat to the validity of the results, then check the appropriate box to assign the level of quality for this study/article.
- Consider an “Unknown” answer to one or more questions as a similar limitation to answering “No,” if the information is not available in the article.

THE EVIDENCE LEVEL IS:

- Good Quality Guideline** [5a]
 Lesser Quality Guideline [5b]
 Not Valid, Reliable, or Applicable

Table of Evidence Levels																				
DOMAIN OF CLINICAL QUESTION	TYPE OF STUDY / STUDY DESIGN																			
	Systematic Review Meta-Analysis	Meta-Synthesis	RCT ⁺	CCT ⁺	Psychometric Study	Qualitative Study	Cohort – Prospective	Cohort – Retrospective	Case – Control	Longitudinal (Before/After, Time Series)	Cross – Sectional	Descriptive Study Epidemiology Case Series	Quality Improvement (PDSA)	Mixed Methods Study	Decision Analysis Economic Analysis Computer Simulation	Guidelines	Case Reports N-of-1 Study	Bench Study	Published Expert Opinion	Local Consensus Published Abstracts
All Domains	1a 1b											4a 4b		2/3/4 a/b	5a 5b	5a 5b	5a 5b	5a 5b	5a 5b	5

⁺ RCT = Randomized Controlled Trial; CCT = Controlled Clinical Trial

Development of this form is based on:

1. The AGREE Collaboration. Appraisal of Guidelines for Research & Evaluation (AGREE) Instrument. www.agreecollaboration.org
2. Guyatt, G., D. Rennie, et al. (2002). Users' guides to the medical literature: a manual for evidence-based clinical practice. Chicago, IL, AMA Press.
3. Fineout-Overholt and Johnston: Teaching EBP: asking searchable, answerable clinical questions. Worldviews Evid Based Nurs, 2(3): 157-60, 2005.
4. Phillips, et al: Oxford Centre for Evidence-based Medicine Levels of Evidence, 2001. Last accessed Nov 14, 2007 from <http://www.cebm.net/index.aspx?o=1025>.