Supplemental Digital Appendix 2 **Data Extraction Form**

Patient feedback review group/MR

Adapted from: BEME CODING SHEET COLLABORATION

http://www.bemecollaboration.org

♦ Citation Type:	☐ Journal article			
	□ Non-peer revi			
	□ Conf. paper / 1	_		
	☐ Official public	cation		
	□ Book			
	☐ Thesis			
	☐ Other	•••••		• • •
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Author(s) Title Publication	Year □ Electronic sea □ Personal recon □ Hand search	Volume rch nmendation	IssuePages	
Author(s) Title Publication Search Method:	Year □ Electronic sea □ Personal recon □ Hand search	Volume rch nmendation	IssuePages	
Title Publication ♦ Search Method:	☐ Electronic sea☐ Personal recon☐ Hand search☐ Other:	Volume rch nmendation	IssuePages	

2.	Aim / Goal	of the Study (cont.)
♦ Tie	d to theoretica	ll/conceptual framework □ Stated □ Not available
Spec	ify the theoreti	ical/conceptual framework used:
•••••		
2	Can dry Dool	
3.	Study Desig	Żu
♦ Tyj	pe of Study:	☐ Research study*
		□ Program description, no data
		☐ Systematic review of the literature
		☐ Literature review
		☐ Opinion or commentary
		□ Other
* spe	cify (definition	ns are provided in Appendix A):
\Box Ex	perimental des	sign
	☐ Randomi	zed controlled trial
	\square Pre-test –	•
		perimental design
	0 0	oup, no comparison
	☐ Historica	
	□ Omei	
	servational St	udies
		ly / case series
		ctional study
	☐ Cohort st	udy
□ Qı	alitative studi	es
□ M i	ixed methods:	(uses both qualitative and quantitative approaches)
4.	Context (st	udy population)
♦ Nu	mber of subjec	cts / size of group
	☐ Physician	ns
	•	

4.	Context (study population, cont.)	
♦ Cou	ntry/location of study	
♦ Lev precep	el/Stage (Please specify if the activity targets a particular group, e.g. community otors)	
□ Pos	tgraduate / residency training	
□ Pro	fessional education General practitioners or family physicians Clinical specialists / specialty: Physicians / doctors unspecified	
	dergraduate healthcare professional school	
♦ Pop	ulation selection, correction for bias:	
□ dro	ibility (random selection)p-outs	
□ inte	ention to treat analysis	
Appro	Strongly Disagree Disagree Uncertain Agree Strongly Appriateness of physician selection and analysis	rgree
□ elig	ent selection, correction for bias	
Appro	Strongly Disagree Disagree Uncertain Agree Strongly Appriateness of patient selection and analysis	ıgree

5. Intervention: Patient Feedback

Educational methods (This refers to the instructional methods used within a particular program type). Please check all that apply and describe carefully.

◆ Program type and duration of exposure to patient feedback (This refers to overall design/format of the program.)

5. Intervention: Patient Feedback (cont.)
 □ Workshop or seminar (Specify duration). □ Short course (Specify duration) □ Longitudinal program (e.g. CME) (Specify duration).
☐ Computer-based program (e.g. online; distance education)
Appropriateness of duration of the programme Strongly Disagree Disagree Uncertain Agree Strongly Agree
♦ Assessments of patient feedback
 □ Patient feedback questionnaire individual. □ Patient feedback questionnaire aggregated. □ Oral.
☐ Intermediated by others (staff, senior doctors, peers). Please specify:
☐ Patient interviews
♦ Presentation of patient feedback
 □ Experiential learning with directly presented patient feedback. □ Experiential learning with collated patient feedback. □ Coaching (Please specify): □ Tailored reports of patient feedback results. □ Other. □ Small group discussions.
♦ Preparation and additional educational methods:
 □ Case-based or problem-based learning. □ Didactic teaching (e.g. lecture). □ Role plays and simulations. □ Films, videotapes and audiotapes □ Written materials and readings. □ Other (Please specify):
Sources of bias:
 □ Adherence to educational training. □ Same intensity for all participants. □ Description of teaching protocol. □ Other.

5.	Intervention: Patient Feedback (cont.)	a	D .			a
Appro	opriateness of adherence	Strongly Disagree	Disagree	Uncertain	Agree	Strongly Agree
♦Instr	ruments					
Sourc	ees of bias:					
	truments in the study were validated ☐ Questionnaires (Please specify) ☐ Other tcomes were assessed by blinded assessors					
Appro	opriateness of questionnaires	Strongly Disagree	Disagree	Uncertain	Agree	Strongly Agree
6.	Expected Learning Outcomes of the Inte	ervention (Ch	eck al	l that a _l	oply.))
	section relates to the intended or expected lea . Please describe the specific focus of the arti	_	e – not	the imp	act o	f the
♦ By	whom were outcomes assessed					
☐ Pat ☐ Tea	ogram participants (physicians)ientsachers			• • • • • • • • • • • • • • • • • • • •		
♦ Hov	w were outcomes assessed and collected					
 □ Into □ Foo □ Liv □ Vio □ Exp 	estionnaire erview cus group ve observation deotape pert opinion ner (Please specify):					

6. Expected Learning Outcomes of the Intervention (Check all that apply. cont.)

♦ Definition	outcome measures / study area :
☐ General co ☐ Aspects or ☐ ☐	of study program/intervention (Please specify)
	lated health outcomes
□ Qu	ality of care(Please specify)
 □ Pa	tient satisfaction (Please specify)
☐ Other (Ple	ease specify)
A. Code the	level of Intervention Studied; Quantification of the Effect level of impact studied in the item and summarize / quantify the results of the at the appropriate level. Note: Include both predetermined and unintended
♦ Kirkpatricl	k hierarchy
Level 1	□ Reaction – covers participants' views on the learning experience, its organization, presentation, content, teaching methods, and aspects of the instructional organization, materials, quality of instruction (i.e. "happiness data").
Level 2a	□ Change in attitudes – outcomes here relate to changes in the attitudes or perceptions among participant groups towards teaching and learning.

7. Impa	act Level of Intervention Studied;	Quantification of the Effect (cont.)
Level 2b	acquisition of concepts, procedur acquisition of thinking/problem-	rskills – for knowledge, this relates to the res and principles; for skills this relates to the solving, psychomotor and social skills.
Level 3	willingness of learners to apply n	
Level 4a		ational practice – refers to wider changes in
	the organization, attributable to t	he educational program.
Level 4b	to improvement in student or res the educational intervention.	es' students, residents and colleagues – refers ident learning/performance as a direct result o
8. Stud	ly Quality	
	ped and equivocal presentation and conclusions (rigour)	Strongly Disagree Disagree Uncertain Agree Strongly Agree
Rigour of co	onduct (qualitative studies)	Strongly Disagree Disagree Uncertain Agree Strongly Agree
Rigour of ar	nalysis (qualitative studies)	Strongly Disagree Disagree Uncertain Agree Strongly Agree

8. Study Quality (cont.)

Optional:					
A. Please rate overall study quality					
				High	
			4		
Comment on overall quality, if applicab	ole		 		
		•••••			
B. Please describe strengths and weakned implementation and data analysis. Do questudy?					-
Strengths					

C. Comments (Please include comments regarding generalizability, educational significance, etc.). Please state the following:

Weaknesses...

No clear conclusions can be drawn
Results weak/ambiguous, but there appears to be a trend.
Conclusions can probably be based on the results.
Results are clear and very likely to be true.
Results are unequivocal.
 -

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article	Conclusions and Practice Implications for Patient Feedback (highlighted by the
• • • • • •	
10. article	Avenues for Suggested Improvements and Further Research (highlighted by the e):
• • • • • •	

Appendix A: Classification of Study Designs

♦ Experimental Designs.

Randomized Controlled Trials: Subjects are randomly assigned to the treatment or control group. Treatment of the two groups is identical, except for the intervention. Cross-over Series: Trials with self controls. Subjects are given one treatment or the control treatment. After a period of time (a "washout" period in a clinical trial), the groups are given the other treatment. This is a crossover study.

Ouasi-Experimental Designs.

Time series: This is a study of one group, over time. Multiple pre-test and post-test measures are taken. In the interrupted time series, one group is studied, multiple pre-test measures are administered over a period of time, followed by an intervention, and then multiple measures or post-tests are taken over a period of time. In the equivalent time series, a single group is studied, but the investigator alternates a treatment with a post-test measure, several times. Repeated measures: All participants in a single group participate in all experiments with each group becoming its own control. All treatments are administered in sequence to the entire group, one at a time, with a measure or post-test, following each treatment. Non-equivalent control group: Matched on key variables. Matching occurs when the investigator believes that such characteristics as age, sex, years of schooling, etc., are so important that an imbalance between the groups would affect conclusions. Both groups are matched to be similar with respect to important characteristics that may otherwise cloud or confound the conclusions.

Trials with external controls. Sometimes controls outside the study are used. These might be the results of another investigator's work, or subjects whom the investigator has treated in a different way previously. The latter are historical controls.

♦ Observational Studies.

Case study/case series: A set of case reports that describe some observations in a small number of patients (persons). These frequently lead to the generation of hypotheses investigated in the other three designs.

Cross-sectional: These are also called surveys. These look at data collected on a group of subjects at one time. They ask "what is happening now?" Surveys are generally cross sectional studies, although they can also be part of a cohort study.

Cohort or longitudinal studies: Cohorts are groups of people who have something in common and who stay together over a period of time (e.g. a medical school class). Cohort studies ask "what will happen?" and look forward in time. Surveys may be used at follow-up points in these studies.

Historical cohort studies may study events that occurred before the study occurred, but the direction of study is still forward.

Correlational studies: These studies are procedures in quantitative research in which techniques are used to describe the relationship or degree of association between or among sets of data. In these studies, there is no intervention applied.

♦ Qualitative Studies.

Grounded theory:

The common experiences of individuals are explored to build a theory.

Ethnography:

Explores the shared culture of groups of people, to understand the processes, and interactions.

Explores individual stories to describe phenomena.

♦ Mixed methods.

These studies use both qualitative and quantitative approaches.

Sources:

The coding sheet and the accompanying definitions have been adapted from:

Dawson, B. and Trapp, R.G. Basic and Clinical Biostatistics (3rd.ed.) New York: Lange Medical Books, 2001

Creswell, J. Educational Research. Planning, Conducting, and Evaluating Quantitative and Qualitative Research Upper Saddle River NJ: Merrill Prentice Hall, 2002.