We16-30 Writing Up and Dissemination

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What are we trying to achieve?

- Explicit description of Review Methods
- Transparent presentation of Data
- Trustworthiness of Authors' Analysis and Conclusions
- Starting Point for Reader's Own Observations

What is required?

- Conformity to Published Reporting Standards (e.g. PRISMA, formerly QUOROM)
- Use of Good Practice in Presentation (e.g. STARLITE for literature searches)
- Imaginative Use of Data Display

PRISMA

- Preferred Reporting Items for Systematic Reviews and Meta-Analyses.
- Evidence-based minimum set of items for reporting systematic reviews and meta-analyses.
- Aim of PRISMA Statement: to help authors improve reporting of systematic reviews and meta-analyses.
- Focus on randomized trials, but PRISMA also a basis for reporting systematic reviews of other types of research, particularly evaluations of interventions.
- May be useful for critical appraisal of published systematic reviews (not quality assessment instrument to gauge quality of a systematic review).

The PRISMA Statement

- Consists of 27-item checklist and fourphase flow diagram.
- Evolving document subject to periodic change as new evidence emerges.
- Update and expansion of now-out dated QUOROM Statement.
- Website (http://www.prisma-statement.org/) contains current definitive version of PRISMA Statement.

27-item Checklist (Items 1 & 2)

- 1. Title: Identify report as systematic review [meta-analysis, or both] (? Qualitative Systematic Review/ Qualitative Meta-Synthesis/ Qualitative Evidence Synthesis?)
- 2. Abstract: Provide structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number.

27-item Checklist – Items 3 & 4

INTRODUCTION			
Rationale	3	Describe rationale for review in context of what is already known.	
Objectives	4	Provide explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS). (?SPICE?)	

27-Item Checklist (Items 5-8, Methods)

Protocol & registration	5	Indicate if review protocol exists, if and where it can be accessed (e.g., Web address), etc.
Eligibility criteria	6	Specify study characteristics (e.g., PICOS, length of follow-up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rationale.
Information sources	7	Describe all information sources (e.g., databases with dates, contact with authors to identify additional studies) in search and date last searched.
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.

27-item Checklist (Items 9-12, Methods)

Study selection	9	State process for selecting studies (i.e., screening, eligibility, included in systematic review).
Data collection process	10	Describe method of data extraction from reports and any processes for obtaining and confirming data from investigators.
Data items	11	List and define all variables (?subject data/author data?/substantiated?) for which data were sought and assumptions and simplifications made.
Risk of bias in individual studies	12	Describe methods used for assessing risk of bias of individual studies (?Reflexivity?)

27-item Checklist

Summary measures	13	State the principal summary measures (e.g., risk ratio, difference in means).
Synthesis of results	14	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., I²) for each meta-analysis.

Analysis

Section/ topic	#	Checklist item
Risk of bias across studies	15	Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies).
Additional analyses	16	Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified.

27-item Checklist (Items 17-20, Results)

Study selection	17	Numbers of studies screened, assessed for eligibility, and included in review, with reasons for exclusions at each stage, ideally with flow diagram.	
Study characteristics	18	For each study, present characteristics for which data were extracted and provide citations.	
Risk of bias within studies	19	Present data on risk of bias of each study.	
Results of individual studies	20	For all outcomes considered provide: (a) summary data (b) effect estimates and confidence intervals, ideally with a forest plot.	

27-item Checklist (Items 21-23, Results)

Synthesis of results	21	Present results of each meta-analysis done, including confidence intervals and measures of consistency. (?reciprocal translation, line-of-argument synthesis?)
Risk of bias across studies	22	Present results of any assessment of risk of bias across studies.
Additional analysis	23	Give results of additional analyses, if done (?Disconfirming case analysis?)

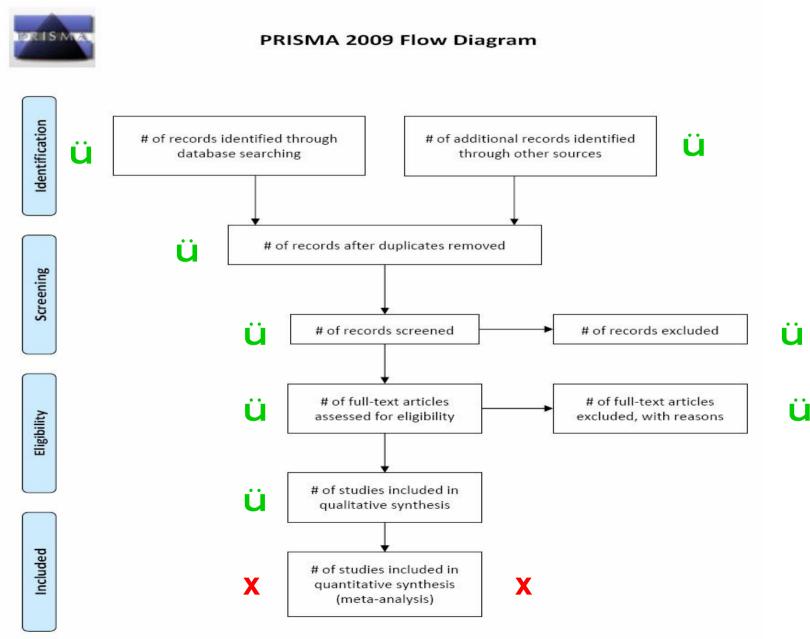
27-item Checklist (Items 24-26, Results)

Summary of evidence	24	Summarize main findings including strength of evidence for each main outcome (?theme?); consider relevance to key groups (e.g., healthcare providers, users, and policy makers).
Limitations	25	Discuss limitations at study and outcome (?theme?) level and at review-level (e.g., incomplete retrieval of identified research, reporting bias).
Conclusions	26	Provide a general interpretation of results in context of other evidence, and implications for future research

27-item Checklist (Item 27, Funding)

Funding 27 Describe sources of funding for the systematic review and other support (e.g., supply of data); role of funders for the systematic review.

Four-phase Flow Diagram



PRISMA – Explanation & Elaboration

PRISMA Explanation and Elaboration document

(http://www.plosmedicine.org/article/info:doi/10.1 371/journal.pmed.1000100) explains and illustrates principles underlying PRISMA Statement.

- To be used in conjunction with PRISMA Statement.
- Part of broader effort, to improve reporting of different types of health research, and in turn to improve quality of research used in healthcare decision-making – EQUATOR Network

Equator Network

(http://www.equator-network.org/)





What is STARLITE?

- STARLITE is a proposal for a framework for reporting the literature searching in systematic reviews and health technology assessments
- It is an acronym STAndards for Reporting LITErature searches
- But it is also a mnemonic......

STARLITE

- S Sampling Strategy
- T Type of Studies
- A Approaches
- R Range of Years (Start Date-End Date)
- L Limits
- Inclusion and Exclusions
- T Terms Used
- E Electronic Sources

Why is STARLITE needed?

- No standard for reporting of literature searching
- Considerable variation in practice
- Decisions taken in searching impact on final review
- Poor searching introduces possibility of publication bias

- Several unilateral attempts to define best practice
- Existing best practice based on effectiveness reviews/HTAs
- PRISMA has very little detail relating to literature searching

Why is STARLITE needed?

Eligibility criteria	в	Specify study characteristics (e.g., PICOS, length of follow-up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rationale.	
Information sources	7	Describe all information sources (e.g., databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched.	
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	

PRISMA items relating to literature searching

	Fully or Partially Present	Absent
Sampling Strategy	43	0
Type of Study	7	36
Approaches	28	15
Range of Years	40	3
Limits (e.g. English)	43	0
Inclusion and Exclusions	8	35
Terms Used	27	16
Electronic Sources	40	3

What is STARLITE not?

- Not yet full standard "Towards" needs tighter specification of data elements and formats
- Not yet consensual framework Phase 1 was "literary warrant", now requires Phase 2 "user warrant" and endorsement.

Good Practice?

Four purposes for data presentation

- Formative to aid conduct of review and insights from findings
- Summative as an output from the review process
- Integrative bringing together quantitative and qualitative elements (Covered in Previous Session)
- Audit to increase confidence in robustness

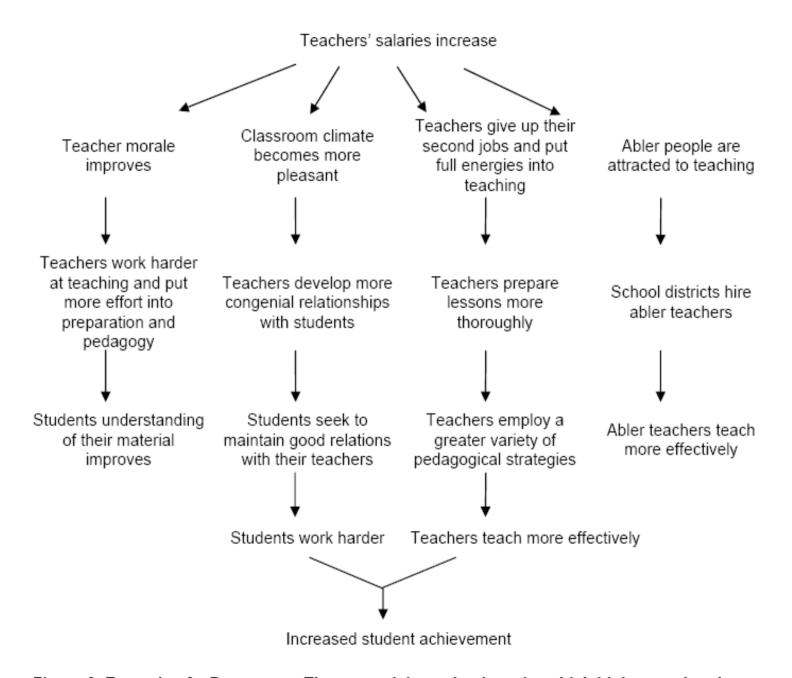
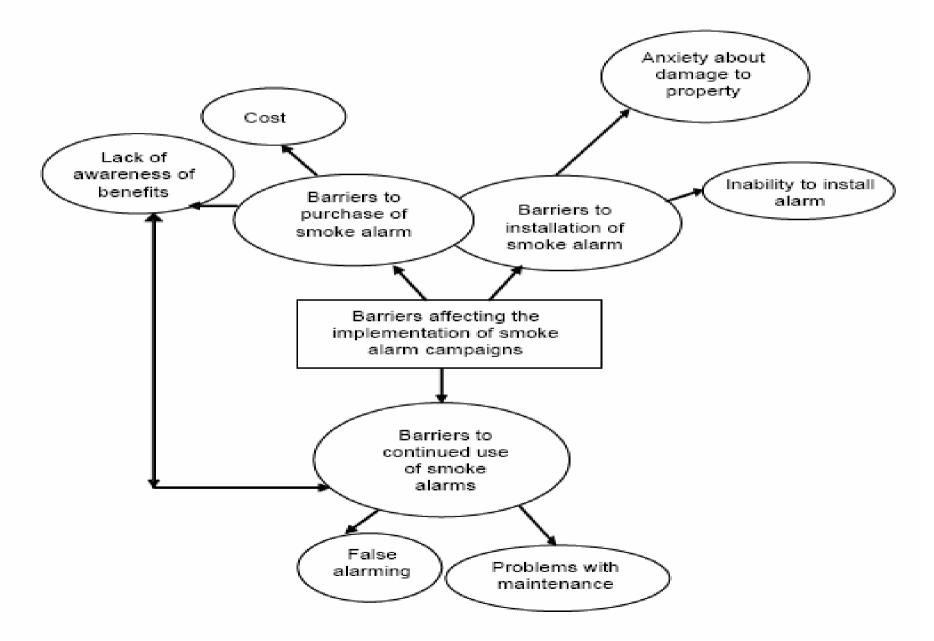
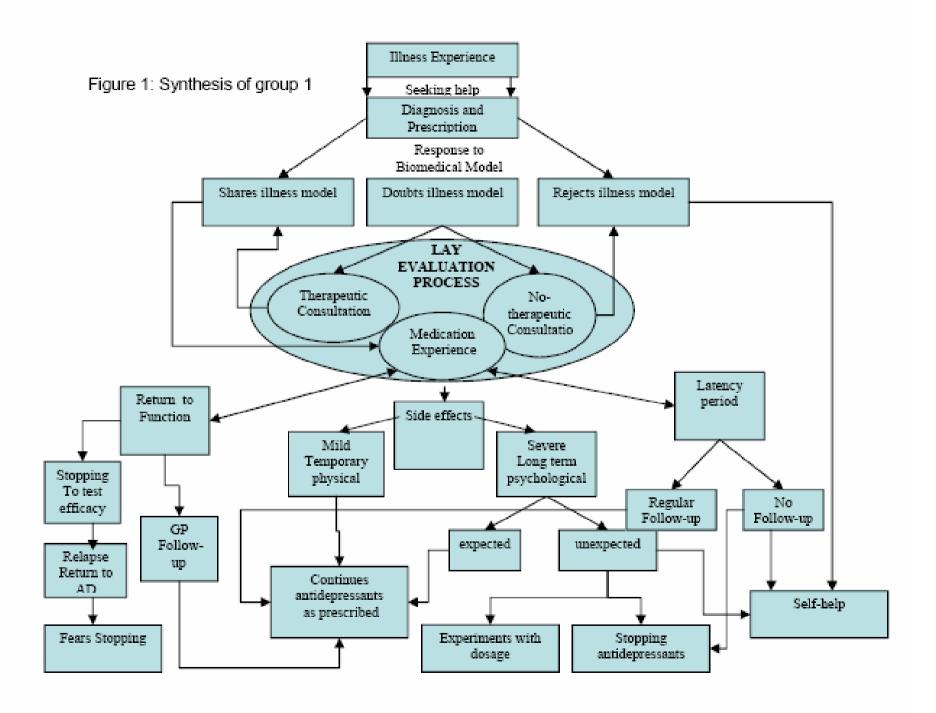


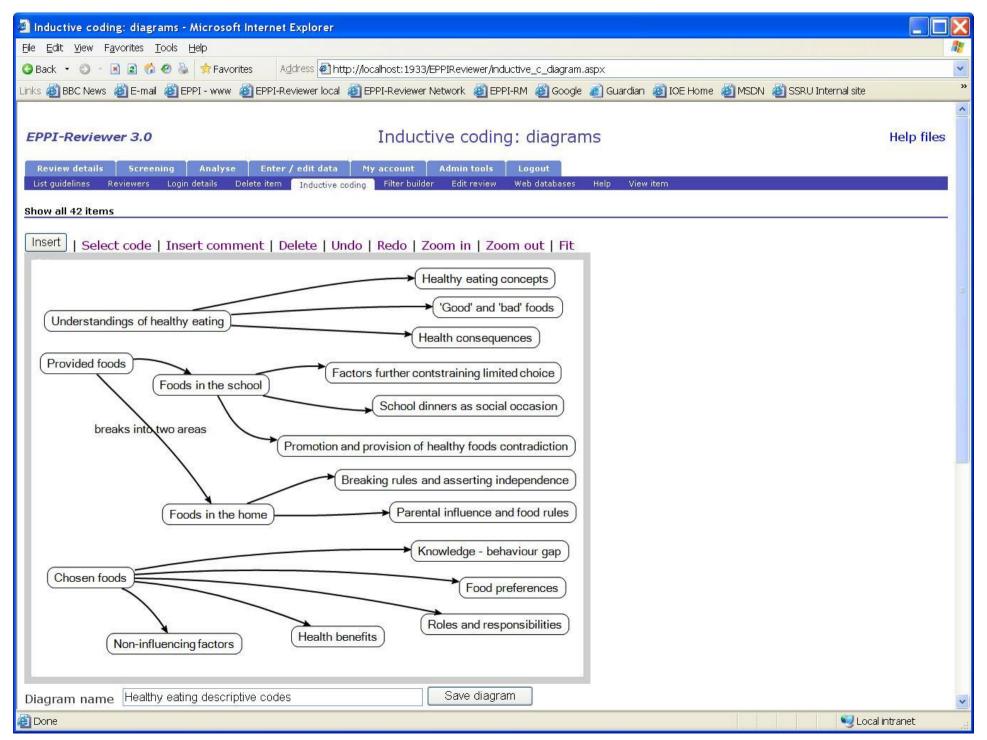
Figure 3. Example of a Programme Theory model: mechanisms by which higher teachers' pay may be linked to increased student achievement (from Weiss, 1998)

Figure 9. Examples of idea webbing

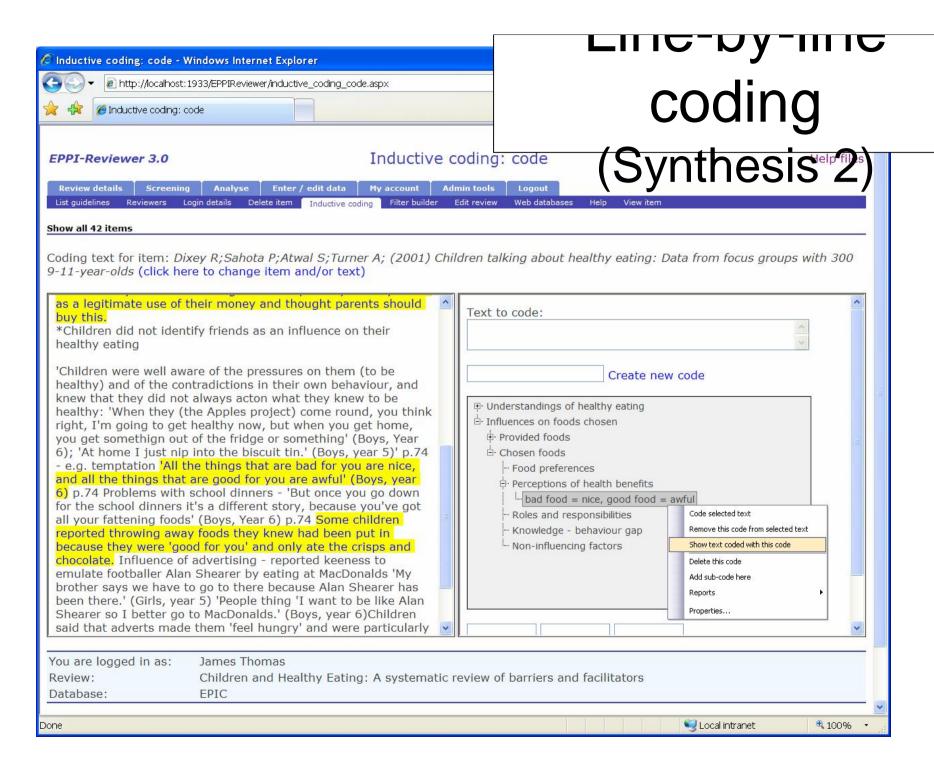
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Synthesis 2: Thematic analysis

- 1) Children don't see it as their role to be interested in health.
- 2) Children do not see future health consequences as personally relevant or credible.
- 3) Fruit, vegetables and confectionary have very different meanings for children.
- Children actively seek ways to exercise their own choices with regard to foods.
- 5) Children value eating as a social occasion.
- 6) Children recognise contradiction between what is promoted and what is provided.

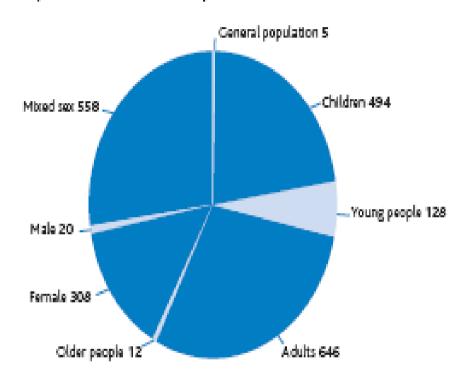
Children consider taste, not health, to be a key influence on their food choice

Food labelled as healthy may lead children to reject them ('I don't like it so it must be healthy')

Buying healthy foods

Buying healthy foods not seen as a legitimate use of their pocket money

Figure 5 Population breakdown in map



Note: Categories not mutually exclusive.

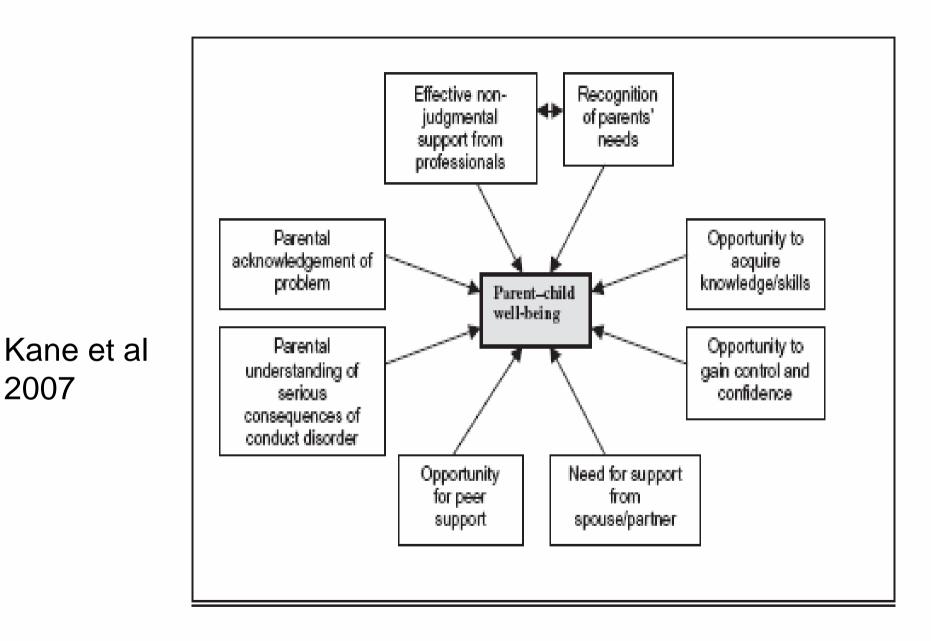


Figure 1. Line-of-argument synthesis: addressing parents' needs and promoting parent-child well-being.

2007

DES	PAIR
 Downward sub-process of despair refers to the destructive path of giving in to hopelessness 	 Upward sub-process of despair refers to the constructive path leading towards hope
CATEGORIES Stopping and being stuck in the situation (III) Losing grip and sinking into a narrowing existence (II,V,VI) Focusing on impossibilities (IV) Losing future perspective (II) Questioning the possibility of hope (II)	CATEGORIES • Fighting against sinking (VI) • Fighting to rise up with a glimmer of hope (III, V)
SUBCATEGORIES Experiencing distressing and stagnant inability (II-V) including panic (VI) Living in exhaustive agony (II) Experiencing lack of alternatives, means and resources (III, V) Being stagnant (V) Being stagnant (V) Sinking down into narrowed existence (III) described as going down (II) and being unable to take hold of anything (VI), A narrowing of the future towards the end (V) such as the narrowing of life (V), non-existence of positive factors upon which to build a future life (V), concealed dreams (V), future life having nothing to offer (V), approaching the end (V), existence of nothing after the end (V), having no grounds for life (V), and indifference in giving up and losing (III, IV) as well as acting	SUBCATEGORIES Understanding the situation (V) Fighting back constructively against sinking (V, VI) Rising up towards hope (V)

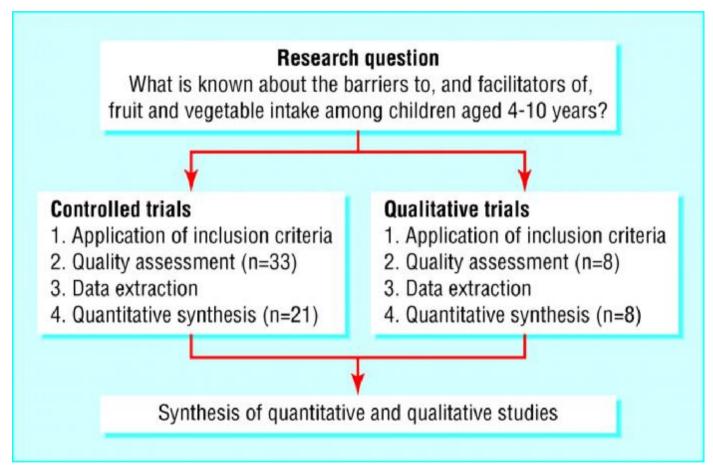
Kylma 2005

Audit - Transparency

 'Given the involvement of the researcher in the research process, the question is not whether the data are biased, but to what extent has the researcher rendered transparent the processes by which data have been collected, analysed and presented' (Popay et al, 1998, p. 348).

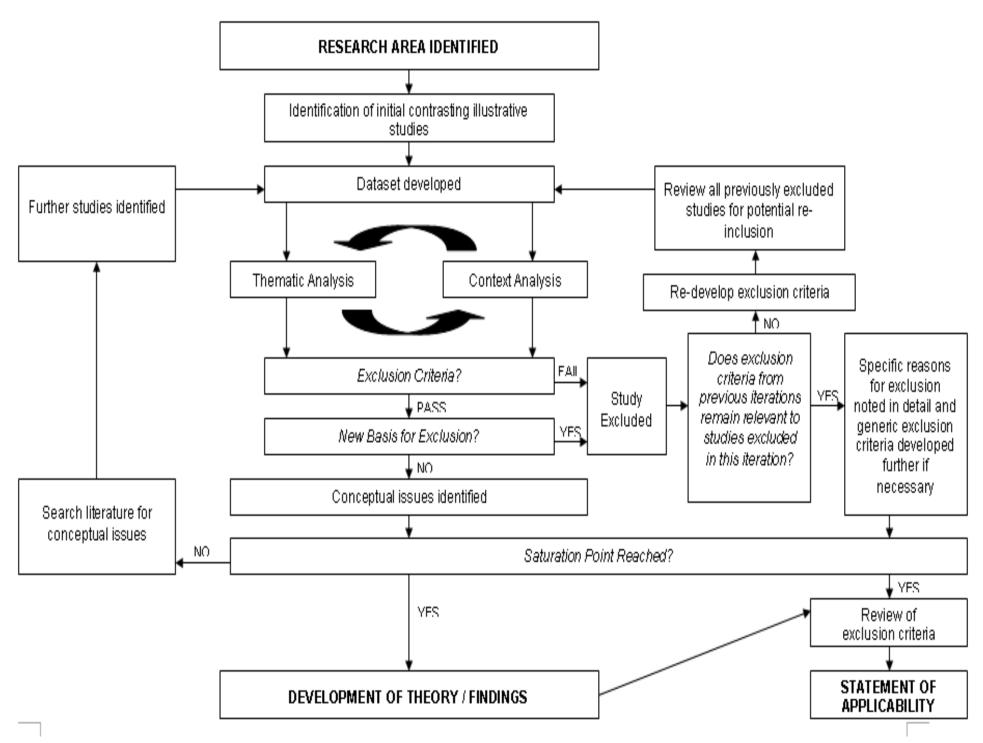
Overall Process

Fig 1 Stages of the review



Thomas, J. et al. BMJ 2004;328:1010-1012





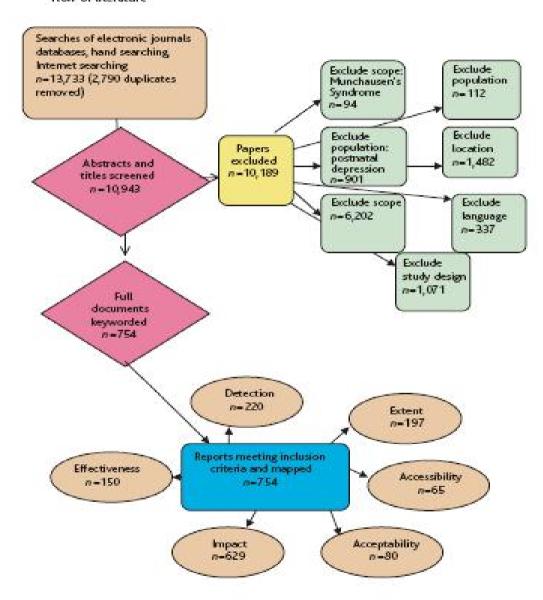
Search Process

Table 1. Final search criteria and search terms using the SPICE(S) tool

Setting	Perspective	Intervention	Comparison	Evaluation	Social science method
Depression	Patient View	Antidepressants	GP and Patient views	Anti-depressant use over time	Qualitative
Depression; Depressive disorder; Depress\$.tw.	Attitude to health; Patient satisfaction; Patient\$ adj3 view\$; Patient care; Patient Compliance; Patient acceptance of health care; Patient participation; Treatment refusal; Patient preference	Antidepressive agents; Antidepress\$.tw	Physician- patient relations	Communication; Decision making; Consultation.tw.	Qualitative research; Qualitative adj research; Grounded adj theory; Ethnograph\$; Qualitative adj studies; Interview\$; Focus groups; Nursing research tw.; mursing research/ or mursing evaluation research/ or exp mursing methodology research/ Field studies; Ethnonursing research; Field studies; Ethnonursing research; Field studies, Ethnonursing research; Field studies,

Figure 2 demonstrates the flow of literature through the systematic map.

Figure 2 Flow of literature



Source: Adapted from EPPI-Centre (2004)

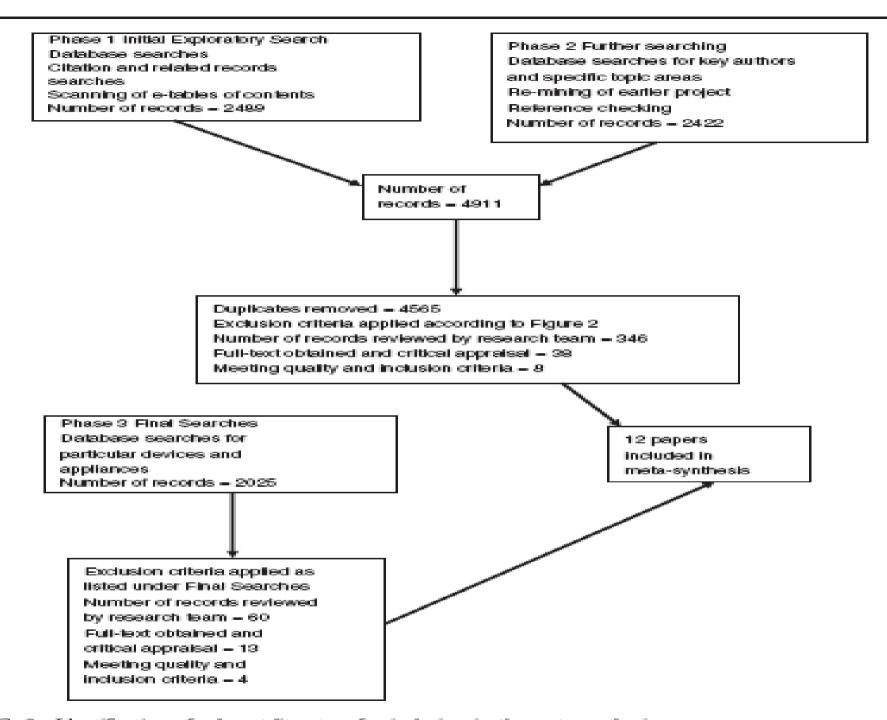


FIG. 2. Identification of relevant literature for inclusion in the meta-synthesis.

Papers identified using: CINAHL Medline. Inclusion and Sociological Abstracts ISI Web of Knowledge Databases **PubMed** Hand Searching of Key Journals **Exclusion** Papers Excluded If: Paper focuses on children Focus of paper only from carers perspective Paper focuses on medication compliance Paper focuses on health professional perspective Paper focuses on an intervention Paper uses mixed methods Paper is a review of existing literature Grey literature Studies of mental health (chronic physical conditions only). Paper presents qualitative data embedded in a randomised controlled trial Qualitative methods are being used to develop measurement tools. Paper focuses on family adaptation to Papers included it: Sufficient evidence of data trail was provided Paper included a health technology Participants were individuals with a long. physical health condition. The setting for use of health technology was the home. The research design was qualitative The study was reported in English

FIG. 1. Inclusion and exclusion criteria for synthesis of patient adaptation to health technologies.

Synthesis

Table 4 – Definition of 1^{st} , 2^{ret} and 3^{rd} order constructs, based on Noblit and Hare (1988)

		anament, contra ou recome mun anne (120		
First	Patients views, accounts			
order	and interpretations of	Interpretations of		
constructs	their experiences of	experience		
	using anti-depressants			
Second	The authors views and			
order	interpretations	T		
constructs	(expressed in terms of	Interpretations of		
	themes and concepts) of	interpretations of		
	patients views of	experience		
	antidepressant use.			
Third	The views and	マケ		
order	interpretations of the			
constructs	synthesis team,	Interpretations of		
	(expressed in terms of	interpretations of		
	themes and key	interpretations of		
	concepts)	experience		

TABLE 2. Main results from the meta-synthesis					
Synthesis of main findings	Line of argument synthesis				
Managing multiple uncertainties Heightened awareness of health deteriorating Continuous feelings of uncertainty about the future New vulnerability to technological failure Living in hope of technological advances					
Technology imposed a routine that facilitated a sense of control and certainty	Adaptation, accommodation and integration of a technology are an extension of identifying and living life with a chronic condition				
The reconstruction of identity Moral imperative to accept a technology Process of comprehension as to how technology will impact upon illness identity. Technology perceived as a signifier of illness Presumption that others will make inaccurate assumptions about the individual.					
Reconstruction of identity that retains a part of pre-illness identity	The integration of a technology or device into the user's life world can be viewed as an extension of existing 'illness work'				
The struggle to remain autonomous while allowing dependence Technology helped maintain some level of independence Devices permitted a greater sense of self-regulation Human qualities attached to the technology that aided engagement					
A new autonomy brought dependence on the technology and others					
Changes to relationships with health professionals experienced Health professional's views perceived to dominate					
Coming to zerous with living a technology-assisted life Integration involved a process of normalization New values and norms incorporated following the introduction of a technology Balance needed between illness regimen and daily life Alterations made to minimize intrusion	The introduction of a technology imposes a new time frame on the individual that must be adhered to, to meet the needs of the technology				
Usability of devices Acceptance linked to user competency and user friendliness of the device					

hygiene of the technology

Usability linked to perceived simplicity, convenience and

Table 5 – Showing translation of 2nd order constructs and their arrangement in temporal sequence

GROUPS of 2 ND	No.	der constructs and their arrangement in temporal sequence Summary definition (translation) of the 2 nd order construct	Papers that
Order	CONSTRUCTS		include the 2 nd
Constructs,			order construct
arranged in			(Figures in bold
temporal			are papers that
seguence			received at least
1.Conditions	Distressed and	Recognition that something is seriously wrong, AND that self-help is not working and experience of distress	one 'KP' rating) 1, 2, 3, 4, 10, 11,
for seeking	needing help	is beyond rational explanation.	12
help	accord acts		
•	Duty to be well	Alignment with treatment goals to return to path of productive, self regulating citizenship.	2, 6, 9, 11, 15, 16
2.Triggers for	Role strain	Recognition that emotional state was effecting the functioning of relationships and ability to fulfil roles and	2, 3, 4, 6, 10, 11
help seeking		take part in normal everyday social relationships .	
	Taking control	Feeling a loss of control and desiring to take back control	2, 3, 4, 12,
	Emotional strain	Felt guilt they had let themselves or others down. Feeling frustrated with self for 'failing' to cope, being 'weak'.	1, 2, 3, 6, 11, 12,
	Stigma	Emotional disorders are perceived as 'stigmatised'. Resisting or rejecting autidepressants (AD) is a way of	1, 2, 3, 4, 5, 6, 7,
3. Barriers to		resisting categorisation as a mentally ill person.	11, 12, 13, 16
accepting treatment	Fear of addiction	Long term use was associated with addiction so it was important to know expected treatment length. Low dosage preferred for same reasons.	1, 2, 4, 5, 8, 11, 12,
	Threat to natural	AD seen as unnatural and leading to "artificial unhappiness" that threatens "real" personality.	1, 2, 4, 11, 12
	self		
	General resistance	Does not normally take medicines, even aspirin, and keen to portray themselves in this way in order to frame	1, 3, 11, 14, 16
	to medicine taking	AD use as last resort. (Also true for patients with substance abuse history).	
4.Paradox of	AD Reduces	Emotional illness conceived as physical deficiency of serotonin, so absolves individual of personal	1, 2, 3, 4, 5, 6, 9,
biomedical	stigma	responsibility, over writing stereotype that depression results from personal weakness. Able to fulfil social	11, 12, 15,
model	AD Doubles	roles and therefore 'normalising'. Prescription of AD experienced as a 'drastic event', making the discredited (unseen illness) discreditable	1224560
	stigma	(seen), therefore doubling existing stigma associated with depression. AD created sense of normalcy (through	1, 2, 3, 4, 5, 6, 8, 9, 11, 15
	2018mm	fulfilling roles) but reduced inner sense of normalcy because taking AD not seen as 'normal'. Feared others'	9, 11, 13
		reactions.	
5.Factors	Threshold of	Desperation to feel better stronger than resistance to AD. Swallowing first pill seen as "swallowing will" and	1, 11, 13

Example of synthesising translations across illness groups

'Rejecters/sceptics' Dowell & Hudson (general medication)

Reject medication due to their values, bypassing testing process.

'Purposeful non-adherence' Johnson et al (hypertension)

A conscious decision not to take drugs, possibly following testing

'Active users' Dowell & Hudson (general medication)

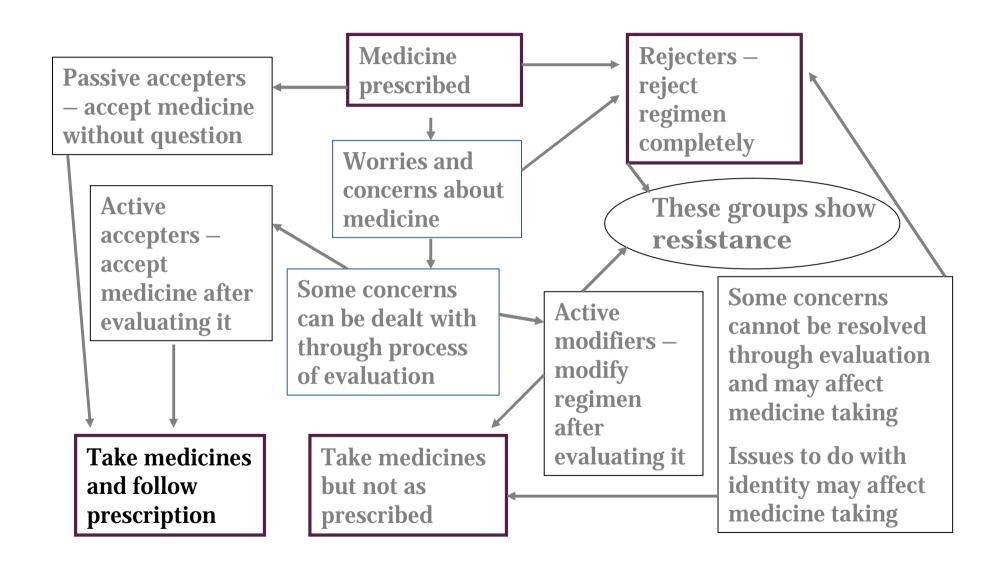
Conscious decision to modify regimen, following testing and deliberation

'Unorthodox Accounts' Britten (general medication)

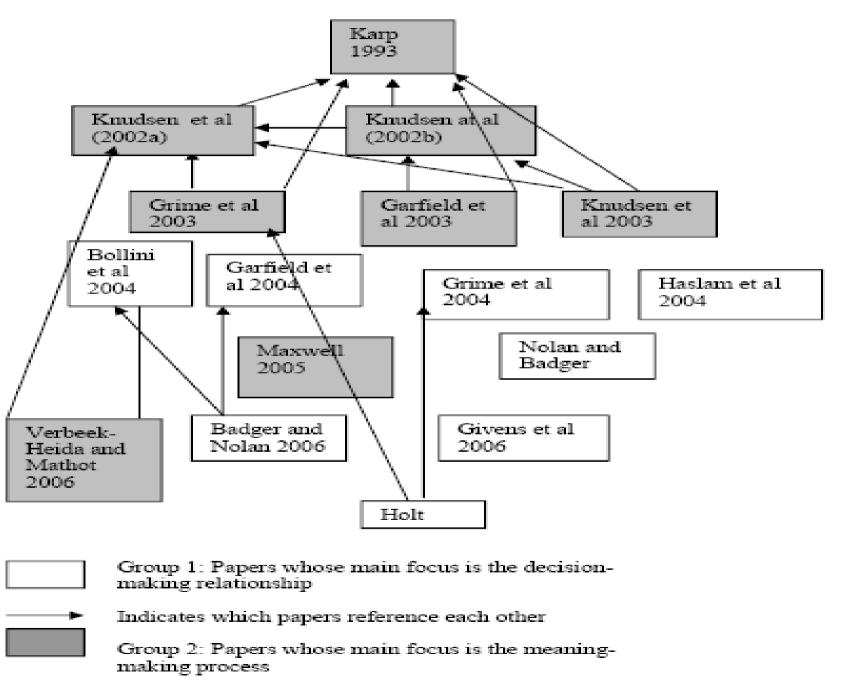
'Self-help repertoire' Lumme-Sandt et al (general medication)

'Justifiers and Excusers' (Siegel et al (HIV)

Excuses offered by those who 'admit behaviour wrong but deny responsibility'. Justifications offered by those who 'take responsibility for behaviour yet deny it has negative consequences'.



Map 1: How the articles in the synthesis reference and



Watch This Space!

- David Moher and Colleagues are currently producing Book on Reporting Standards
- Cochrane Qualitative Methods Group currently contributing Chapter on Reporting of Qualitative Research
- Discussions Ongoing about Standards for Reporting Qualitative Evidence Syntheses