

"Epidemiologist know a lot about the correct way to conduct a research study but less about how to review and synthesize data from multiple studies and this, I suggest, is a principal source of the public's confusion when faced with a new result from an epidemiological study"

### Bracken MB. IJE 2001:954





## Systematic reviews of observational data







### What are observational studies?

- Data from existing database
- Cross-sectional study
- Case series
- Case-control study
- Cohort study
- Design with historical controls



# Why do we need systematic reviews of observational studies?

- Aetiological hypothesis
- Evaluation of interventions designed to prevent rare outcomes
- Evaluation if outcomes of interest are far in the future
- Evaluation of effectiveness in a community





### MAOS are common

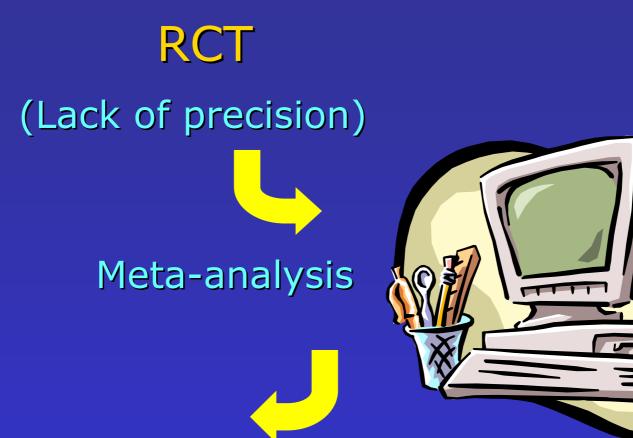
Type of article	Articles (n)
Meta-analysis of:	
Controlled trials	34
Observational studies	25
Methodological article	15
Tradicional review	15
Other	11

Source: Egger M. Systematic reviews in Health Care. Meta-analysis in context. BMJ Books. 2001

**DEPARTMENT OF REPRODUCTIVE HEALTH AND RESEARCH** 







#### More reliable estimates

**DEPARTMENT OF REPRODUCTIVE HEALTH AND RESEARCH** 





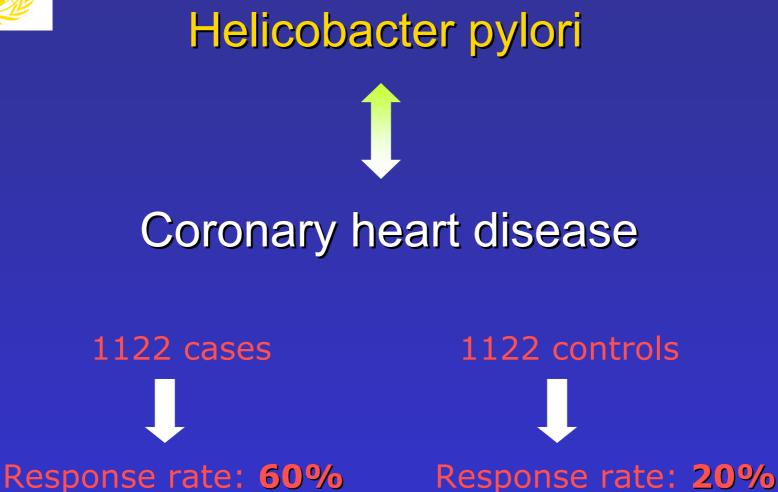
# **Observational studies** (Confounding, bias) Meta-analysis

#### More reliable estimates????

**DEPARTMENT OF REPRODUCTIVE HEALTH AND RESEARCH** 







Source: Danesh J. Helicobacter pylori infection and early onset myocardial infarction: case control and sibling pair study. BMJ 1999;319;1157.

DEPARTMENT OF REPRODUCTIVE HEALTH AND RESEARCH



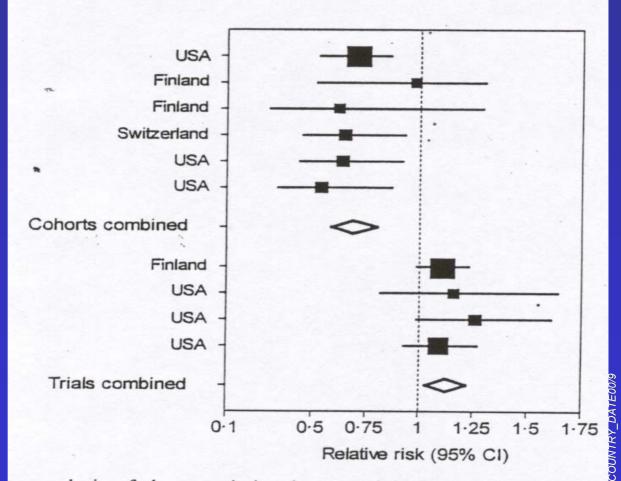
# The protective effect of beta-carotene that wasn't

#### Cohorts

Male health workers Social insurance, men Social insurance, women Male chemical workers Hyperlipidaemic men Nursing home residents

#### **Trials**

Male smokers Skin cancer patients (Ex)-smokers, asbestos workers Male physicians







There are examples of observational studies producing similar results of those from RCT

But observational studies will always have to deal with **bias** and **confounding** because the intervention was deliberately chosen and not randomly allocated





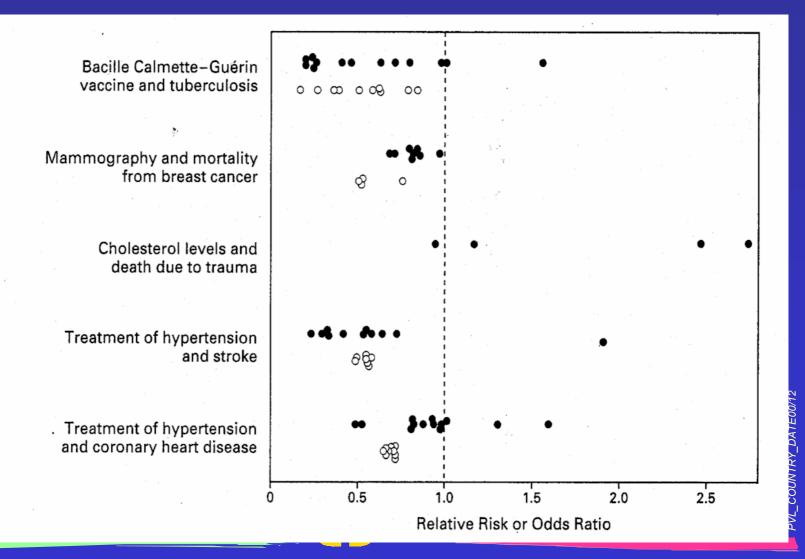
### Benson and Hartz, NEJM, 2000;342:1878-86

Freatment Evaluated	Outcome	OR ar	nd 95% Cl	
		0.10	1.00 10	.00
		First treatmen better	t Second treatmen better	t
lifedipine vs. control in patients with CAD*	Mortality			]
Observational (30–60 mg) Randomized, controlled (30–50 mg)				
ABG vs. PTCA in diabetic patients*	Mortality			
Observational Randomized, controlled			• •	
ABG vs. PTCA in patients at high risk* Observational	Mortality		•	
Randomized, controlled		F		
ABG vs. PTCA in patients at low risk* Observational	Mortality			
Randomized, controlled			•	
ABG vs. medical treatment in CASS patients	Mortality			
Observational Randomized, controlled				
ABG vs. medical treatment in Duke study patients† Observational	Mortality			
Randomized, controlled				
eta-blockers vs. control† Observational Randomized, controlled	Mortality	•		/.





### Concato et al., NEJM, 2000;342:1887-92







### This does not mean to return to narrative reviews

**DEPARTMENT OF REPRODUCTIVE HEALTH AND RESEARCH** 





### **Benefits of MAOS:**

 Systematic and explicit rules Statistical power Insight into variable interpretation Detection of discrepancies Deepness into heterogeneity Identification of gaps in knowledge





## Reporting of background should include:

- 1 Problem definition, hypothesis statement
- **2** Description of study outcome(s)
- **3** Type of exposure or intervention used
- 4 Type of study designs used
- **5** Study population



### Reporting of search should include:

- 6 Qualifications of searchers
- 7 Search strategy including time period
- 8 Effort to include all available studies
- 9 Databases and registries searched
- **10** Searching software used
- **11** Use of hand searching
- 12 List of citations located and those excluded, including justification
- 13 Methods of addressing articles not published in English
  14 Methods of handling abstracts and unpublished studies
  15 Descriptions of any contact with authors



### Reporting of methods should include:

- **16** Description of relevance/appropriateness of papers assembled for assessing the hypothesis to be tested
- **17** Rational for the selection and coding of data
- **18** Documentation about how data were classified and coded
- **19** Assessment of confounding
- 20 Assessment of study quality, including blinding of quality assessors; stratification or regression on possible predictors of study results
- **21** Assessment of heterogeneity
- **22** Description of statistical methods in sufficient detail to be replicated

**23** Provision of appropriate tables and graphics



### Reporting of results should include:

- 24 Graphic summarizing individual study estimates and overall estimate
- 25 Table giving descriptive information for each study included
- **26** Results of sensitivity testing (e.g. subgroup analysis)
- **27** Indication of statistical uncertainty of findings





28 Quantitative assessment of bias29 Justification for exclusion

**30** Assessment of quality of included studies





## Reporting of conclusions should include:

- **31** Consideration of alternative explanations for observed results
- **32** Generalization of the conclusions
- **33** Guidelines for future research
- **34** Disclosure of funding source





Quality of reviews in Epidemiology Breslow R. AJPH, 1998;88:475-7

### All 1995 issues of 7 widely read epidemiology journals were searched for reviews

### 29 reviews were found

DEPARTMENT OF REPRODUCTIVE HEALTH AND RESEARCH





### Reviews following quality guidelines

Guideline	Yes	Unable to determine	No
Search methods stated	6 (21)	1(3)	22(76)
Inclusion criteria reported	5(17)	4(14)	20(69)
Bias in selecting studies avoided	3(10)	26(90)	0(0)
Criteria for assessing validity reported	2(7)	15(52)	12(41)
Methods for combining findings reported	10(34)	6(21)	13(45)
Conclusions supported by data	24(83)	4(14)	1(3) 1(3)





### Search restriction: General medical journal, 2001

Search Procedure	19 meta- analyses	13 systematic reviews
Numerous Databases Searched (versus just MEDLINE)	13 (68%)	6 (46%)
Additional Searches Conducted (e.g., manual search of reference lists or textbooks)	17 (89%)	10 (77%)
Gray Literature Searched (e.g., manual search of conference or dissertation abstracts)	5 (26%)	4 (31%)
Contacted Experts to Find Unpublished Data	7 (37%)	2 (15%)
<b>Cochrane Databases Searched</b>	8 (42%)	4 (31%)
All Methods Employed	4 (21%)	1 (8%)

Source: Becker B, Morton S (see http://www.msri.org/calendar/talks/TalkInfo/1268/show\_talk)



### Search restriction: General medical journal, 2001

Language Restriction	19 meta- analyses	13 systematic reviews
None	6 (32%)	1 (8%)
English plus other lang.	2 (11%)	0 (0%)
English only	7 (37%)	7 (54%)
Unclear	4 (21%)	5 (38%)
Attempted to include unpublished studies	7 (37%)	5 (38%)

Source: Becker B, Morton S (see http://www.msri.org/calendar/talks/TalkInfo/1268/show\_talk)





### Other citations:

- Mulrow CD. The medical review article: state of the science. Ann Intern Med 1987, 6:233-240.
- McAlister FA, Clark HD, van Walraven C et al. The medical review article revisited: has the science improved? Ann Intern Med 1999, 131:947-951
- Bracken MB. Commentary: towards systematic reviews in epidemiology. *IJE* 2001, 30:954-957.





### Summary

- SR and MA of observational studies are as common as reviews of RCT
- Confounding and selection bias often distort the findings
- Danger in producing very precise but spurious results
- More is gained by examining heterogeneity





### WHO Systematic review of incidence/prevalence of maternal mortality and morbidity 1997-2002





### **Objectives**

- To provide a comprehensive, standardised and reliable tabulation of available data on maternal morbidity
- To provide up-to-date data for future maternal mortality estimates
- To provide case-fatality rates





### **Objectives (cont.)**

- To estimate relative risks for maternal mortality in different health care scenarios
- To calculate the proportion of maternal deaths that can be averted (preventable fraction) by eliminating or reducing leading morbidities





#### CHARACTERISTICS OF THE STUDY

- 3. Study design
  - (1) Census
  - (2) Cross-sectional
  - (3) Cohort/longitudinal
  - (4) Controlled trial
  - (5) Incidence/Prevalence survey
  - (6) Unknown
  - (7) Other, specify —
- 4. Sampling
  - (1) Random sample
    - 4a. Specify the method of randomization:

WHO CODE

WHO CODE

WHO CODE

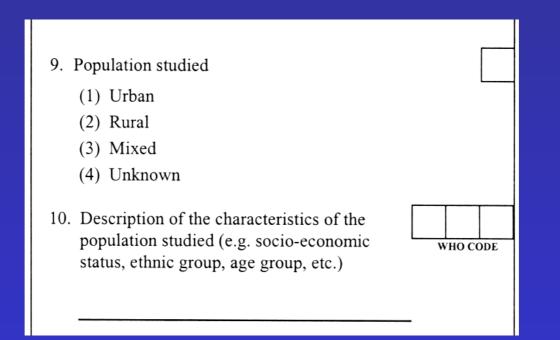
- (2) Non-random sample
  - 4b. Specify the method of sampling:
- (3) Total population (i.e. census)
- (4) Unknown

### WHO systematic review

5. Data source (1) Vital statistics/census (2) Medical record (3) Special survey/interview (4) Multiple sources (5) Clinical data collected for the study (6) Other, specify WHO CODE 6. Lowest unit of data source (1) Cluster 6a. Number of clusters (2) Individual (3) Other, specify \_\_\_\_\_ WHO CODE



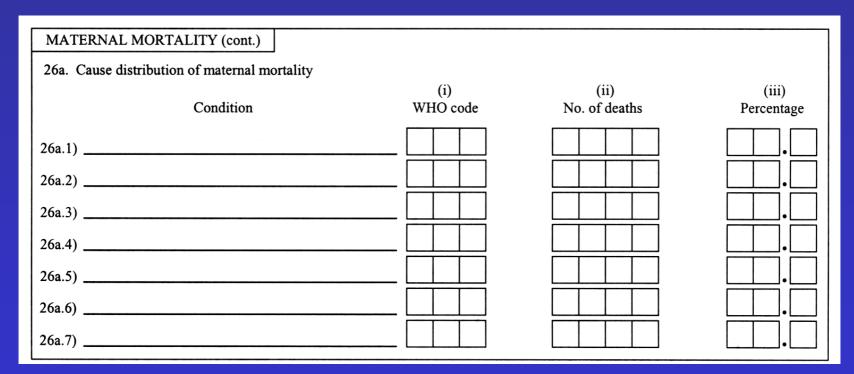
### WHO systematic review







### WHO systematic review





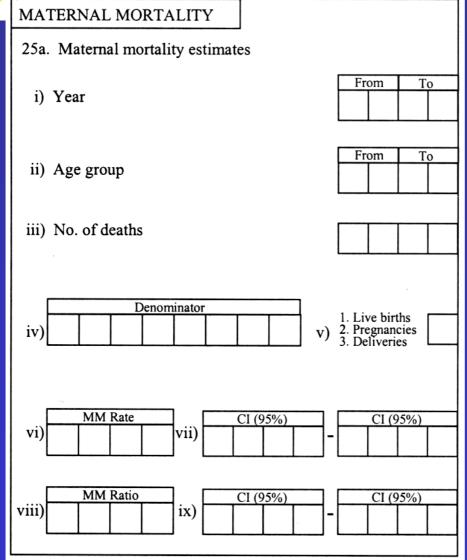


### WHO systematic review

32.	Infections	WHO code
	32a) Condition	
	32b) Does the study include a definition?	Yes No
	32c) If definition is included, please specify:	WHO code
	<ul><li>32d) Does the study explain the method of assessment of the infection?</li><li>32e) If method of assessment is explained,</li></ul>	Yes No
	please specify:	WHO code







### WHO systematic review







PVL\_COUNTRY\_DATE00/3

WilliamaHamilton RNewsYorker 2001ND RESEARCH











DEPARTMENT OF REPRODUCTIVE HEALTH AND RESEARCH

