Critical appraisal of the medical literature

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Training course in research methodology and research protocol development

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What makes a study reliable?



Nine out of ten women we asked believe that "Youth Code" makes their skin firmer and younger looking

Which design maximises the chance of getting the result the company wants?

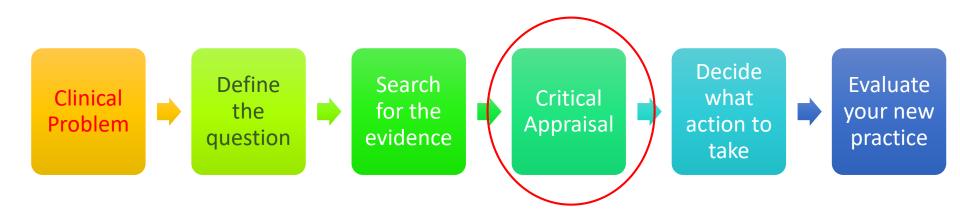
- a) Ask women buying "Youth Code" in the shops whether they agree their skin is firmer and younger looking?
- a) Ask a random sample of women to try "Youth Code" and then comment on whether they agree that their skin is firmer and younger looking?

What is critical appraisal?

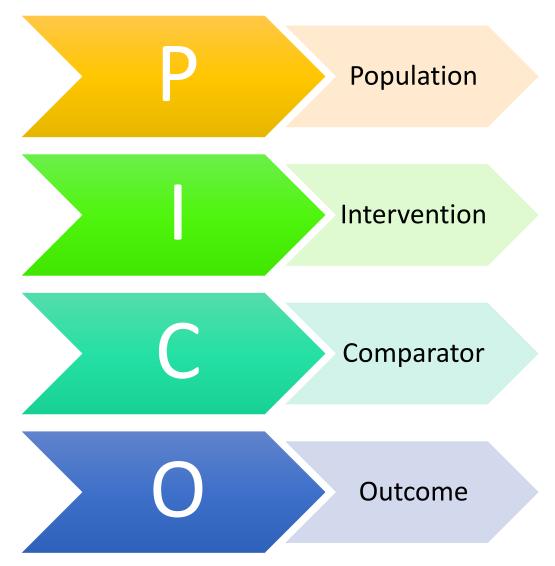
- Carefully and systematically evaluate research to assess:
 - Validity (is these findings trustworthy?)
 - Value (what do the results show?)
 - Relevance (How do these results relate to my clinical practice?)



Critical appraisal: a key component of evidence based medicine



Asking the right question



	1 Patient or Problem	Intervention (a cause, prognostic factor, treatment, etc.)	3 Comparison Intervention (if necessary)	4 Outcomes
Tips for Building	Starting with your patient, ask "How would I describe a group of patients similar to mine?" Balance precision with brevity.	Ask "Which main intervention am I considering?"Be specific.	Ask "What is the main alternative to compare with the intervention?" Again, be specific.	Ask "What can I hope to accomplish?" or "What could this exposure really affect?" Again, be specific.
Example	"In patients with heart failure from dilated cardiomyopathy who are in sinus rhythm"	" would adding anticoagulation with warfarin to standard heart failure therapy"	" when compared with standard therapy alone"	" lead to lower mortality or morbidity from thromboembolism. Is this enough to be worth the increased risk of bleeding?"

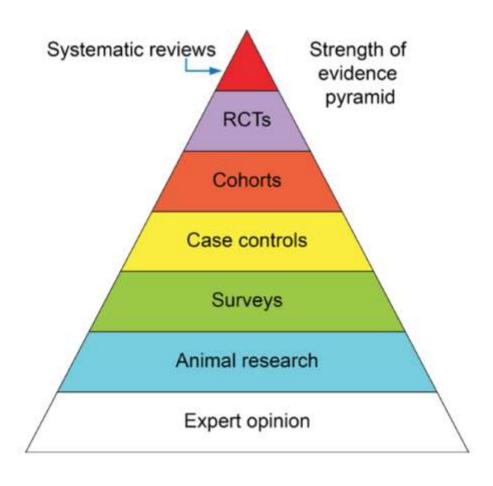
Choosing right study design

Some study designs are not appropriate to answer certain questions

All study designs are prone to different biases



Pyramid of evidence



So are RCTs the gold standard for evidence?



....depends

Slides from: K Mahtani, CEBM Oxford

Limitations of RCTs

- Excellent vs Poor RCTs quality varies
 - Impact on interpretation of result (external validity)?
- Expensive and time consuming
 - £250k £millions over 2-5 years+
- May not always be the right study design to answer that question

A RCT to examine if smoking causes lung cancer

- 30 healthy Oxford Students
- Randomise to 2 groups
 - Gp1 smokes 20 cigarettes per day every day
 - Gp2 no smoking











Types of research

- What is the best study design for answering this type of question?
 - Aetiology
 - Diagnosis
 - Prognosis
 - Harm
 - Effectiveness
 - Qualitative

How to critically appraise an article

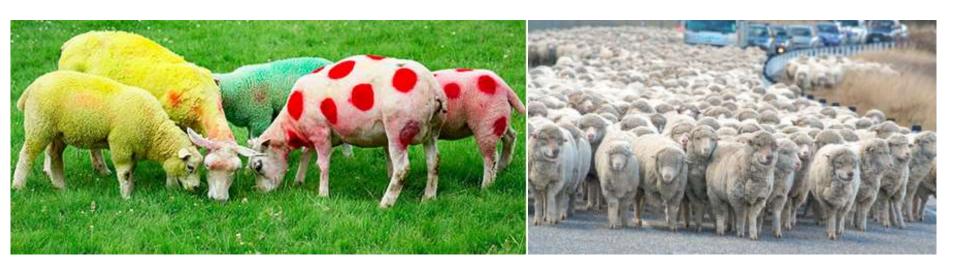
 Validity: methods to check that the biases for which that particular study design is prone have been minimised

- Results
- Clinical relevance

Validity

Internal

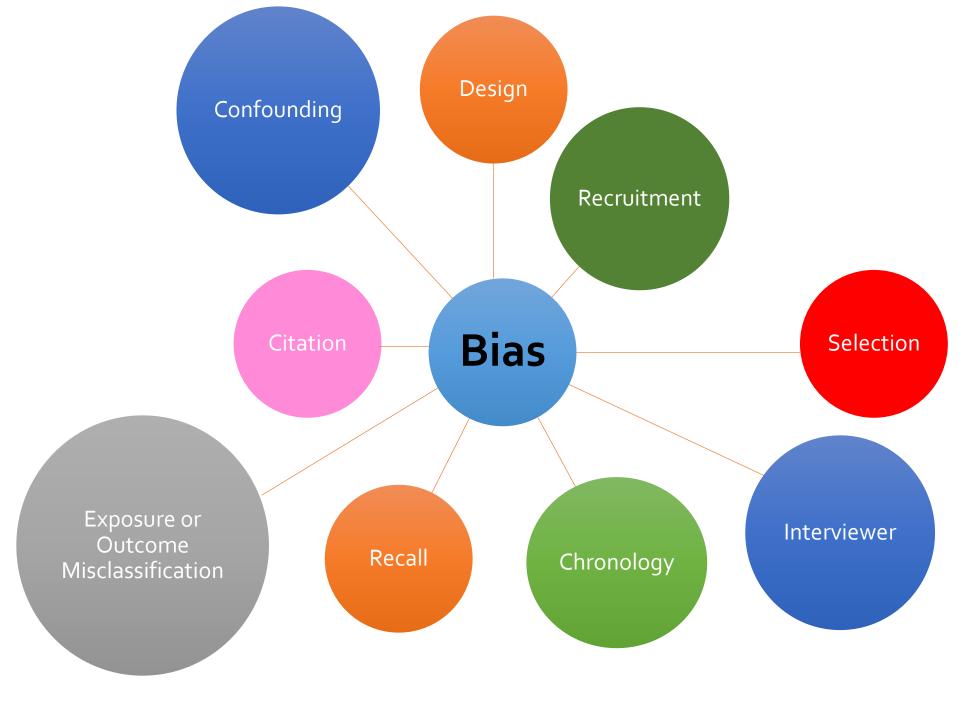
External



Bias

"the systematic deviation of the results of a study from the truth because of the way it has been conducted, analysed or reported"





Sources of bias in clinical trials

Table 1. Key sources of bias in clinical trials ²				
Selection bias	Biased allocation to comparison groups			
Performance bias	Unequal provision of care apart from treatment under evaluation			
Detection bias	Biased assessment of outcome			
Attrition bias	Biased occurrence and handling of deviations from protocol and loss to follow up			

Assessing Trials of effectiveness

Questions to ask:

- 1. Are the results of the trial valid?
- 2. What are the results?
- 3. Will the results help locally?

From: Critical Appraisal Skills Program, Oxford

https://casp-uk.net/

Checklists for clinical trials















Critical Appraisal Skills Programme (CASP)

Making sense of evidence

11 useful questions for critical appraisal of a randomised trial

1. Did the trial address a clearly focused issue?







Consider: An issue can be 'focused' In terms of

- The population studied
- The intervention given
- The comparator given
- The outcomes considered

Representative: Are the trial subjects representative of patients in this setting?

All people in study setting

Eligible participants

In trial

randomised?







Consider:

- How was this carried out, some methods may produce broken allocation concealment
- Was the allocation concealed from researchers?

Why randomise?

• Minimises measured and unmeasured confounding



Minimising allocation bias

Centralised computer randomisation the best

Other methods such as sealed envelopes doubtful

• Non randomised: date of birth, alternate patients alternate days, etc.

If answer to first two questions is no....



Detailed questions

3. Were patients, health workers and study personnel blinded?

Consider:

- · Health workers could be; clinicians, nurses etc
- · Study personnel especially outcome assessors





4. Were the groups similar at the start of the trial?







Consider: Look at

· Other factors that might affect the outcome such as age, sex, social class, these may be called baseline characteristics



Maintenance: Were the groups treated equally?

5. Aside from the experimental intervention, were the groups treated equally?



6. Were all of the patients who entered the trial properly accounted for at its conclusion?





Consider:

- Was the trial stopped early?
- Were patients analysed in the groups to which they were randomised?



From: Critical Appraisal Skills Program, Oxford

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(B) What are the results?

7. How large was the treatment effect?

Consider:

- What outcomes were measured?
- Is the primary outcome clearly specified?
- What results were found for each outcome?
- Is there evidence of selective reporting of outcomes?

8. How precise was the estimate of the treatment effect?

Consider:

- What are the confidence limits?
- Were they statistically significant?

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Intention to treat

- Once a participant is randomised, they should be analysed to the group they were assigned to
- Pros
 - Reflects "real life" e.g. non compliance
 - Unbiased estimate of true effect
 - Maintains sample size thus maintaining statistical power
- Cons
 - Noncompliance provides little data on efficacy
 - Treatment effect may be conservative
 - Dropouts/non-compliant/compliant subjects are different

What does this study tell us?

- P values (hypothesis testing):
 - Tests to exclude the null hypothesis
- Confidence intervals (estimation of effect)
 - Range of values within which the true effect is likely to lie
 - Wider the confidence interval, less precision in result
- Relative Risk
- Absolute Risk
- Odds Ratios
- Number needed to treat

(C) Will the results help locally?

9. Can the results be applied in your context?
(or to the local population?)



Consider:

- Do you have reason to believe that your population of interest is different to that in the trial
- If so, in what way?





Yes Can't tell No 10. Were all clinically important outcomes considered? Consider: Is there other information you would like to have seen? Was the need for this trial clearly described? Yes Can't tell No 11. Are the benefits worth the harms and costs? Consider:

 Even if this is not addressed by the trial, what do you think?

From: Critical Appraisal Skills Program, Oxford https://casp-uk.net/

Conclusion

 Critical appraisal helps us decide whether evidence is valid, what the results tell us and whether the study is relevant to our setting

Checklists are available to help

Don't believe everything you read in journals!