

identify people with disease
and without first

why (definition),
relationship
how frequently
the disease is
present in each
group.

Type of Sampling
Measurements:

Case-Control Study

Nested case-control

why?
from same
same
population

select control
when each case
occurs

Source of controls

Source population:

Case selection

Control selection

Case-Crossover Study

Regression to the mean

pros:
(Epid)

- ① underlying cohort (source pop) known.
- ② less mislabeling of E
- ③ E measured before D

Adv.

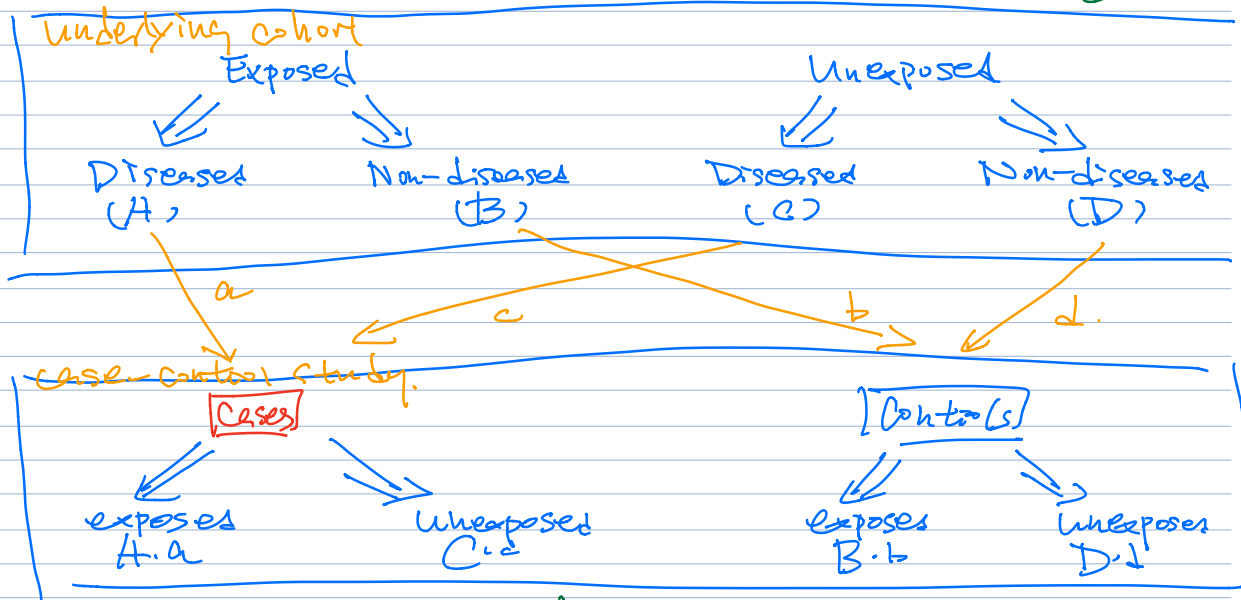
cons:

pros:

When will sampling from the cohort yield the same OR as in the full cohort?

Remember loss-to-follow-up and OR?

change type of attrition \Rightarrow type of sampling.



Fixes cohort. End of study ascertainment.

cohort OR = $\frac{A \cdot D}{B \cdot C}$. Case-control OR = cohort OR $\frac{a \cdot d}{b \cdot c}$

selection odds

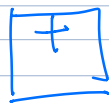
Needs balanced odds

$\frac{d}{b} > 1$ OR \uparrow

$\frac{d}{b} < 1$ OR \downarrow

Should select controls independent of exposure.

Source population Clear?
(cases)



What is the source for hospital?

- referrals
- Locals
- Internationals



- select controls from that source
- friend controls Guide. (overmatch SES).
- ① same source pop: controls are those if they got the disease would be cases in your study
 - ② Same inclusion/exclusion for case/control
 - ③ Same method for exposure assessment.

What happens if there is significant censoring in the underlying cohort?



Case-Base sampling.

Risk Ratio

$$= \frac{a}{c} = \frac{N_1}{N_0}$$

(Regardless of rare disease assumption)

Traditional:
Cumulative
Incidence
Sampling
 $OR = \frac{ad}{bc}$

Incidence-Density Sampling

Rate ratio

$$= \frac{a}{c} = \frac{PI_1}{PI_0}$$

(Regardless of rare disease assumption)

Main source of invalidity:

- Caution: Any Bias in the underlying cohort will reflect in the case-control study

- recreate wrong source pop

- create non-comparability in source.

- Exposure Ascertainment (differential misclassification)

- Reverse Causation

- cases: Surveillance Bias. Diagnostic Bias. Prevalence Bias.
Prevalence cases are survivors. risk factor may be related to survival not

↑ R_s

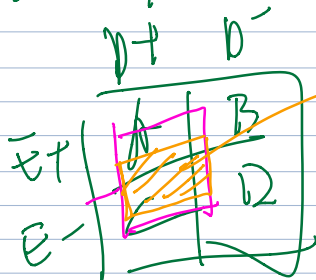
Recall bias.

Interviewer bias

Recording bias

A Good Case-Control Study depends on selecting controls from the right source population independent of exposure and with measurement of the exposure that is the same for cases and controls.

Controls should be selected from the same population - the source population - that gives rise to the study case. Exposure distribution in controls should be representative of the exposure distribution in source population.



Non-differential

highly motivated healthy controls tend to participate \Rightarrow then the frequency of exposure in control group will be representative of the frequency in the base population.