

Types of missing data

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Study is completed:

- All patients measured
- Determinants/predictors/covariates/independent variables/X'n AND the outcome/dependent variable/Y
- Data in computer
- Descriptive/frequency tables MISSINGS



- MISSINGS = problems → precision (loss of statistical power) + validity (bias)
- TO PREVENT IS BETTER THAN TO CURE



- Missing data always occur (any study):
 - retrospective and prospective
 - existing (routine care) databases
 - large scale population-based
 - (even!) well organised randomised trials
- Challenge: how proper analyses with missing values?
 - unbiased effect estimates (validity)
 - precise effect estimates (precision)



- Epidemiological analyses:
 - Association determinants (predictors/ covariates/confounders) with outcome
 - Multivariable (regression) analyses
 - What happens with participant record if one variable (X or Y) is missing?
 - = complete case (CC) analyses
 - = most common
 - = always affects precision of effect estimates (not all data used)
 - = commonly causes invalidity (bias)



- Most (epidemiological) studies use complete case analysis → Ignoring (=one method)
- There are other methods to handle missings may also cause bias.
- Type and severity bias depend on method used and type of missing data.



- 3 types: MCAR, MAR, MNAR
- Missing Completely At Random (MCAR)
- Missing At Random (MAR)
- Missing Not At Random (MNAR)



1. Missing Completely At Random (MCAR)

- The probability that an observation is missing does not depend on 'anything' except chance
- Examples?
- The probability that the observation of a given variable for a certain subject is missing is constant for all subjects
 - Missingness NOT related to any other patient characteristics -> including the outcome status



- If MCAR holds, almost all analytical methods (see later) for handling missing data give unbiased results, although less precise
- Realistic?
 - Reason: missing related to other patient characteristics, including outcome (!!!)
 - = MAR = missing at random



2. Missing At Random (MAR)

most advanced methods work very well

 Probability that an observation is missing depends only on other observed values (patient characteristics, including the outcome)

Most advanced methods to handle missing values under MAR yield in principle unbiased + more precise study results → see later



3. Missing Not At Random (MNAR)

- The probability that an observation is missing depends also on unobserved values.
- E.g. probability of missing on a variable depends on the true (but unknown) value of that variable itself
- Examples
 - homosexual/heterosexual (homosexual are less likely to put down their sexual preference in a questionnaire) or higher incomes don't want to write down their income
 - income level (to estimate SES)
 - higher levels values have larger probability of being missed than lower values



- Missing data seldom (if ever) MCAR
- MNAR = problems → no general methods for properly dealing with MNAR data!

How to check likelihood of missing data being MCAR or rather MAR?

usually it's MAR - how can we find out?

Next slide = very important table in empirical research!

"most"
starts all analysis



| Table. Distribution of co-variates among subjects without and with missing values (total n=398). | | | | | | | |
|--|--------------------------------------|----|--------------------|-----------------------------|--------------------|--------------------|--|
| | one hopes that those are No missings | | $\geq 1 \text{ m}$ | | p-value | | |
| Variables | a random subcategory n=246 (62%) | | n=152 (38%) | | | | |
| Pulmonary embolism (outcome variable) | | 47 | complete CCA | 36 | at least one NA | 0.02 | |
| Dyspnoea index tests | | 80 | | 66 | | ≤0.01 | |
| Malignancy | values are associated with | 28 | | 16 | | <0.01 | |
| Surgery in previous 3 months | missingness - showed that NA | 24 | | 16 | | 0.04 | |
| Prior deep venous thrombosis | were not MCAR | 6 | | 10 | | 0.17 | |
| Wheezing | | 18 | | 11 | | 0.09 | |
| Previous pulmonary embolism | | 5 | | 12 | | 0.02 | |
| Collapse with or without loss of consciousness | | 10 | | 5 | | 0.06 | |
| Signs of deep venous thrombosis | | 11 | | 7 | | 0.15 | |
| Age (years)* | | 57 | (17) | 54 | (18) | 0.19 | |
| Positive Chest x-ray | | 43 | | 36 | | 0.13 | |
| Respiratory rate (breaths/min)* | | 22 | (7) | 18 | (6) | \<0.01 \ | |
| * Mean (sd) | | | | they are different however! | | | |

Testing for MCAR/MAR

Missing data CLEARLY not MCAR observed characteristics

because otherwise the subsets would have been equal in observed characteristics

- Analyzed subset of 246 subjects is not random subset of the original study sample (N=398) -> SELECTION bias due to missings
- Missing related to other observed characteristics to predict the missing values (Incl. outcome) = MAR
- If missingness related to observed characteristics these can be used to estimate/predict the missing values!
- Missing could still partly MNAR but also MAR. Cannot test for MNAR
 only reduce MNAR-part as much as possible by including many observed chars (increasing MAR)
 - Compare: adjustment for known confounders (MAR) versus residual confounders (MNAR)

 document known confounders as much as possible and asjust for known confounders the more conf. the less residual confounders

the more observed characteristics the less likely MNAR

- Exception: missing outcomes in RCTs previous table not enough
 - See later (Groenwold RH et al: CMAJ 2014 + AM J EPI 2012)

Thank you for your attention

