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School of Medicine

Confidence Intervals (CIs)

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Learning Outcomes

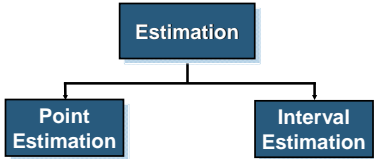
Following this session you should be able to:

- Understand the concepts and interpretation of confidence intervals;
- Explain how they are derived
- Understand how they can be used to assess precision
- Demonstrate how they are should be presented
- Use software to calculate them

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Estimation Methods

We rarely measure the whole population

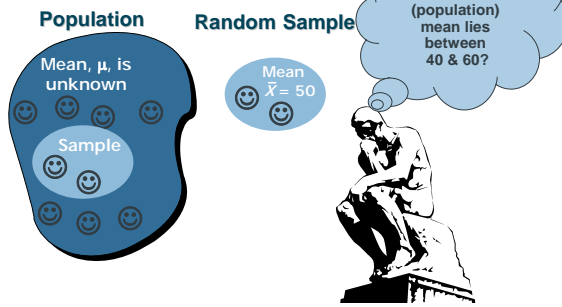


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graph TD; Estimation --> PointEstimation[Point Estimation]; Estimation --> IntervalEstimation[Interval Estimation];
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Point Estimation

- Provides Single Value
 - Based on Observations from 1 Sample
- Gives No Information about how close our value is to the unknown Population Parameter
- Example: Sample Mean $\bar{X} = 50$
Point Estimate of unknown Population Mean

Estimation Process



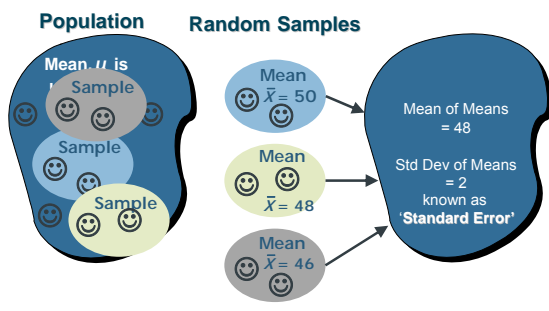
Estimation from a population

- The population is defined as the group about whom statements will be made
- If a representative sample is taken conclusions from the sample can be generalized to the wider group

Understanding Statistical notation

	Population	Sample
Mean	μ	\bar{x}
Standard Deviation	σ	S (SD Std Dev)

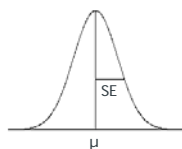
Estimation Process



Estimating the mean of a continuous variable

Repeated sampling from the population gives samples means whose frequency distribution (sampling distribution) properties are:

- The mean of this distribution would be the population mean μ
- The standard deviation of this distribution of sample means is called the **Standard Error (SE)**



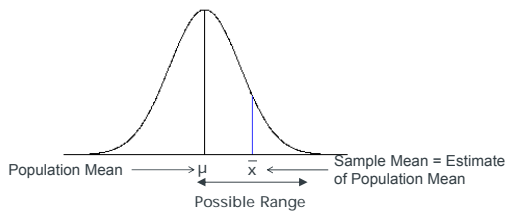
Standard Error (SE)

- The Standard Error measures how precisely the population mean is estimated by the sample mean
- SE is estimated by the sample SD divided by the square root of the number of observations

$$SE = \frac{SD}{\sqrt{n}}$$

Estimating the mean of a continuous variable

- Using the properties of the normal distribution we can estimate the range in which the unknown population mean lies

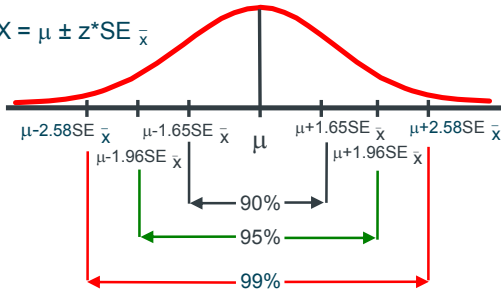


Estimating the mean of a continuous variable

- This range is called the **95%** confidence interval about the mean
 - It is calculated as:
Sample mean \pm 1.96 * Standard Error
 - All values within the confidence interval are reasonable values for the population mean that generated the observed sample
 - It gives an idea of the precision of the estimate from the sample size available

Different Degrees of Confidence

$$X = \mu \pm z * SE_{\bar{x}}$$



Different forms of Confidence Intervals

- Continuous outcome variables : Means, Medians
 - One sample
 - Two sample (difference)
- Categorical outcome variables : Proportions
 - One sample proportion
 - Two sample proportion (difference)
- Correlation
- Odds ratio (OR) & Relative risk (RR)
- Standardised Mortality ratios

Alternative Distributions

Different Confidence Interval calculations require different theoretical distributions

Means (small numbers)	t distribution
Standardised Mortality ratios	Poisson distribution
Medians	Binomial Distribution

They all need a sample estimate and a standard error

Association between CI and P values

Differences in Continuous measures or proportions

- If 95% CI includes 0 then p value will be **greater than** 0.05
- If 95% CI does not include 0 then p value will be **less than** 0.05

Ratios and Risks

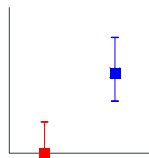
- If 95% CI includes 1 then p value will be **greater than** 0.05
- If 95% CI does not include 1 then p value will be **less than** 0.05

$$99\% = 0.01$$

CI & Hypothesis Testing

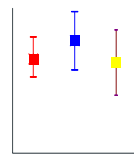
If CIs do not cross at a significance level (say 5%), then hypothesis testing is significant but the opposite is not always true?

Confidence intervals NO overlap



Can conclude that there is a real difference between the two groups

Confidence intervals overlap



Cannot draw any conclusions about difference without further information

Example 1: Interpreting a rate

- Sample of 1106 pregnancies, estimated rate of congenital abnormality was 4.2% (95% CI 3.0% to 5.3%)
- The 'true' population rate could be as low as 3.0%
- The 'true' population could be as high as 5.3%
- There is a 1 in 20 chance that our estimate is wrong and that the true population value is outside this range
- Our best estimate of congenital abnormality is 4.2%

Example 2: Interpreting a difference between two means

- Mean birthweight was measured in a sample of 15 non-smokers (3.59Kg) and 14 heavy smokers (3.20Kg)
- The difference in the mean weight was 390g (95%CI 60g to 721g)
- The 95% CI excludes 0, therefore the difference is statistically significant (P will be less than 0.05)
- Although the difference is significant, our estimate of the Percentage difference is $390 / 3400 = 11.5\%$
- Is this clinically important?

Example 3: Interpreting differences

- Length of stay in hospital
Group 1 (n= 392) Mean stay 37 days
Group 2 (n= 368) Mean stay 41 days
- Difference = 4 days (95%CI = -2 to 9) days
- 95% CI includes 0, not statistically significant ($P > 0.05$)
- The study has been unable to rule out that the true difference could be 9 days
- Lack of evidence of a difference is NOT EVIDENCE of no difference!

Example 4: Interpreting proportions

- RCT of flu vaccine
- Infection rate of placebos 80/220 (36%)
- Infection rate for subjects 20/240 (8%)
- Difference in rates 28% (95%CI 21% to 35%)

- 95% CI excludes 0, difference was significant ($P < 0.001$)
- The true difference is at least 21%, best estimate is 28%
- Vaccine clearly demonstrates protective effect
- But..... side effects, consider costs, generalisability

Confidence Intervals (CIs) or P values?

- Leading medical journals recommended both when reporting the main study results
- Use of CIs recommend by the ICMJE
- Over emphasis on the P values detracts from more useful approaches when interpreting study results

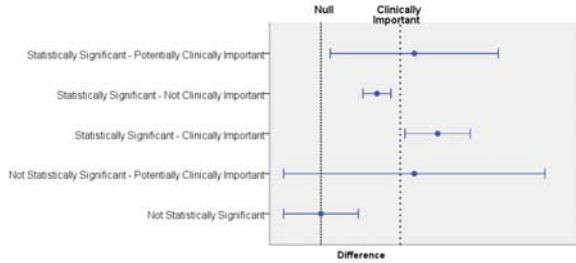
The problem with P values

- Wrong type of thinking through use of arbitrary cut off at a predefined level (5%)
- Low quality information with
 $P < 0.05$, $P > 0.05$, $P = NS$
- $P = 0.049$ is declared as **significant** and
 $P = 0.051$ as **not significant**
- Cut off leads to statistical significance being equated with clinical significance

The problem with P values continued

- A very small improvement, 1%, of one treatment compared to another may be statistically significant ($P < 0.001$)
- Only quoting P values may lead uncritical reader into thinking that treatment A was more effective than treatment B
- A clinically important effect may be non-significant because of a small sample size

Statistical and Clinical Importance



Redrawn from Machin, Campbell and Walters – Medical Statistics 2007

Confidence intervals from SPSS

Descriptives

		Statistic	Std. Error
DIABETIC	Mean	146.4500	1.8457
95% Confidence Interval for Mean	Lower Bound	142.7878	
	Upper Bound	160.1122	
5% Trimmed Mean		146.7333	
Median		146.0000	
Variance		340.654	
Std. Deviation		18.4568	
Minimum		77	
Maximum		190	
Range		113	
Interquartile Range		23.7500	
Skewness		-.422	.241
Kurtosis		1.683	.478

Confidence intervals from SPSS

Extracorporeal membrane oxygenation * 1yr survival Crosstabulation

		1yr survival		Total
		Yes	No	
Extracorporeal membrane oxygenation	Yes	63	30	93
	% within Extracorporeal membrane oxygenation	67.7%	32.3%	100.0%
	No	38	54	92
		41.3%	58.7%	100.0%
Total	Count	101	84	185
		54.6%	45.4%	100.0%

Chi-Square Tests

	Value	df	Asymp. Sig. (2-sided)	Exact Sig. (2-sided)	Exact Sig. (1-sided)
Pearson Chi-Square	13.040 ^a	1	.000		
Continuity Correction ^b	11.996		.001		
Likelihood Ratio	13.201		.000		
Fisher's Exact Test				.000	.000
Linear-by-Linear Association	12.970		.000		
N of Valid Cases	185				

a. Computed only for a 2x2 table

b. 0 cells (.0%) have expected count less than 5. The minimum expected count is 41.77.

Continuous variable: paired/unpaired

Example : Difference in systolic BP in 100 diabetic and 100 non diabetic subjects mean 146 SD 18.5 and mean 140 SD 16.8

Means and their differences:
Unpaired samples

Proportions: two unpaired sample

Example : Trial of asthma treatment which measured respiratory failure. Control group it was 63/93 and in treated it was 38/92

Proportions and their differences:
Unpaired samples

A Common Question

What is the difference between Reference Range and Confidence Interval?

Reference Ranges refer to **Individual values** and Confidence Intervals to **Estimates**

Reference Range uses **Standard Deviation**

$$\text{Mean} \pm 1.96 * \text{Std Deviation}$$

Confidence Interval uses **Standard Error**

$$\text{Mean} \pm 1.96 * \text{Std Error}$$

Quoting Confidence Intervals

- They are not required for all results
- Not required for the mean response of subjects to treatments A and B, if major outcome was the **difference** between treatments A and B
- Generally restricted to the main outcome of the study which is usually a contrast (difference) between means or proportions

Quoting Confidence Intervals

The difference between the sample mean systolic blood pressure in diabetics and non-diabetics was 6.0 mmHg, with a 95% confidence interval from 1.1 to 10.9 mmHg; the t test statistic was 2.4 with 198 degrees of freedom and an associated P value of 0.02

Mean difference was 6.0 mmHg (95%CI 1.1 to 10.9 mmHg)

Summary

- Indicate the (im)precision of sample estimates as population values
- They give a range of values for the estimated population parameter (difference)
- They depend on
 - Sample size (larger sizes give narrower CIs)
 - Variability of parameter being estimated
 - Degree of confidence required (90%, 95%, 99%)

References

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- Kirkwood B.R. & Sterne J.A.C. *Essential Medical Statistics*. 2nd Edition. Oxford: Blackwell Science Ltd 2003.
<http://www.blackwellpublishing.com/essentialmedstats/default.htm> Chapters 6 to 8 (and others).
- Altman DG, Bland JM. (1995) Absence of evidence is not evidence of absence. *BMJ* 311 485.

Questions?
