

# Power and Sample Size

## STA 102: Introduction to Biostatistics

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The following material was used by Yue Jiang during a live lecture.

Without the accompanying oral comments, the text is incomplete as a record of the presentation.

# Why calculate sample size and power?

- ▶ To show that under certain required conditions, a hypothesis test has a good chance of showing the anticipated difference, if it really exists
- ▶ To be more confident that a null result is not simply a sample of excessive variability
- ▶ To show a funding agency that the study has a reasonable chance of reaching a useful result
- ▶ To show that necessary resources (human, animal, financial, time, etc.) will be minimized

# Example

Question: Does exercise affect body weight?

- ▶ Study design: participants will be randomized to two groups, exercise and control
- ▶ Outcome: change in weight from randomization to end of study
- ▶ Want to detect: weight change as small as one pound
- ▶ Known: SD of weight change is around 1.5 pounds, as given in exercise science literature
- ▶ Unknown: how many subjects do we need to show a difference, if one truly exists?

# Study design

Proper study design depends on a variety of factors, including

- ▶ Measurement scale of variable of interest
- ▶ Hypothesis of interest
- ▶ Size of result that is clinically meaningful
- ▶ Cost of carrying out the study
- ▶ Time needed to carry out the study

# Power

Showing adequate statistical power is usually necessary in order to get funding for research!

# Power

*Power* is the probability of rejecting the null hypothesis when it is false (i.e., of avoiding a Type II error)

$$\text{Power} = P(\text{reject } H_0 | H_0 \text{ is false})$$

and can be also thought of as the likelihood a planned study will detect a deviation from the null hypothesis if one really exists.

Power is a function of

- ▶ Sample size  $n$
- ▶ Deviation from the null one hopes to detect
- ▶ Standard deviation  $\sigma$
- ▶  $\alpha$ , the Type I error rate

$n$ , detectable difference, variance, and  $\alpha$

How do these four considerations affect power?

# Power

When we design a study, it is not enough to know we have a small probability of rejecting  $H_0$  when it is in fact true. We want to know we have a large probability of rejecting the null when it is false. Practically speaking, power less than 80% is typically considered insufficient to warrant a study.

Increasing power by tolerating more Type I errors is not acceptable. Therefore, we can increase power by

- ▶ Considering larger deviations from the null (need to think about clinical/practical importance)
- ▶ Increasing  $n$  (good, though not always affordable)

# Interactive visualization

Let's examine a [web visualization](#) and see how this all works for a one-sample z-test.

## What is this deviation?

When we calculate power, we need to know the minimum difference from the null mean  $\mu_0$  that we wish to detect. We may set up our hypotheses as

$$H_0 : \mu = \mu_0$$

$$H_1 : \mu \neq \mu_0$$

but need to specify a *minimum detectable difference*, often called  $\delta = \mu_1 - \mu_0$  such that we reject  $H_0$  with a certain power (usually 80% or 90%) when in fact  $\mu = \mu_1$ . We will need a bigger sample size when  $\delta$  is small, and fewer subjects when  $\delta$  is larger.

For example, in the weight change study, we would have  $\delta = 1$  lb.

## Sample size for two-sided one-sample test of mean

Power and sample size are interrelated. Ideally, when we plan a study, we have a prespecified idea of the minimum difference we want to detect,  $\delta = \mu_1 - \mu_0$ , the standard deviation,  $\sigma$ , and the power,  $1 - \beta$ , we'd like to have to detect it. In that case, it is simple to calculate the required sample size (here given for a one-sample test):

$$n = \left[ \frac{\sigma (z_{1-\alpha/2}^* + z_{1-\beta}^*)}{\mu_1 - \mu_0} \right]^2$$

## Sample size estimation based on CI width

Sometimes we're instead interested in estimating an effect with a given degree of precision. Suppose we wish to estimate the mean of a normal distribution with sample variance  $s^2$  and require that the two-sided  $100\% \times (1 - \alpha)$  CI for  $\mu$  be no wider than  $L$ .

How many subjects would we need to ensure that this is the case as a function of  $L$ ,  $s^2$ , and  $\alpha$ ?

Hint: use what you know about how this confidence interval would be constructed.

## Case study: ultra low dose contraception

Suppose you want to ensure a new manufacturer of birth control pills provides the correct dosage of  $0.02 \mu\text{g}$  estrogen. How many pills do you need to sample in a shipment in order to ensure 80% power to detect a difference of 10% at  $\alpha = 0.05$ , assuming  $\sigma = 0.008$ ? (10% of 0.02 is 0.002)

$$\begin{aligned} n &= \left[ \frac{\sigma (z_{1-\alpha/2}^* + z_{1-\beta}^*)}{\mu_1 - \mu_0} \right]^2 \\ &= \left[ \frac{0.008(1.96 + 0.84)}{0.002} \right]^2 \\ &= 125.44 \end{aligned}$$

# Fractional people

To be conservative, always round up for sample size calculations.

R uses a  $t^*$  instead of a  $z^*$  approximation. Do you expect the necessary sample size to be higher, lower, or the same?

## Back to the hypothetical exercise study...

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