UCLA STAT 13

Introduction to Statistical Methods for the Life and Health Sciences

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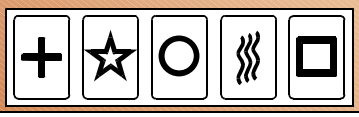
Chapter 9: Significance Testing -- Using Data to Test Hypotheses

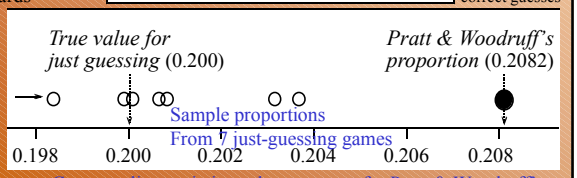
- Getting Started
- What do we test? Types of hypotheses
- Measuring the evidence against the null
- Hypothesis testing as decision making
- Why tests should be supplemented by intervals

ESP (extra sensory perception) or just guessing?

Deck of equal number of Zener/Rhine cards



n=60,000 random draws resulting in 12,489 correct guesses



Can sampling variations alone account for Pratt & Woodruff's success rate = 20.82% correct vs. 20% expected.

ESP or just guessing?

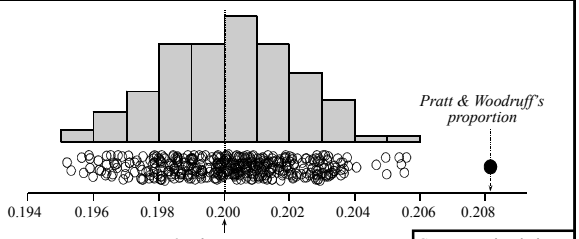


Figure 9.1.1 Sample proportions from 400 "just-guessing" experiments.

Computer simulation making 60,000 guesses with 20% chance of correct guess.

From *Chance Encounters* by C.J. Wild and G.A.F. Seber, © John Wiley & Sons, 2000.

Was Cavendish's experiment biased?

A number of famous early experiments of measuring physical constants have later been shown to be biased.

Mean density of the earth

True value = 5.517

Cavendish's data: (from previous Example 7.2.2)

5.36, 5.29, 5.58, 5.65, 5.57, 5.53, 5.62, 5.29, 5.44, 5.34, 5.79, 5.10, 5.27, 5.39, 5.42, 5.47, 5.63, 5.34, 5.46, 5.30, 5.75, 5.68, 5.85

$n = 23$, sample mean = 5.483, sample SD = 0.1904

Was Cavendish's experiment biased?

Simulate taking 400 sets of 23 measurements from N(5.517,0.1904). Plotted are the results of the sample means. Are the Cavendish values unusually diff. from true mean?

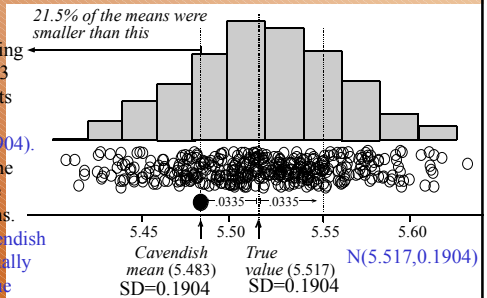


Figure 9.1.2 Sample means from 400 sets of observations from an unbiased experiment.

Cavendish: measuring distances in std errors

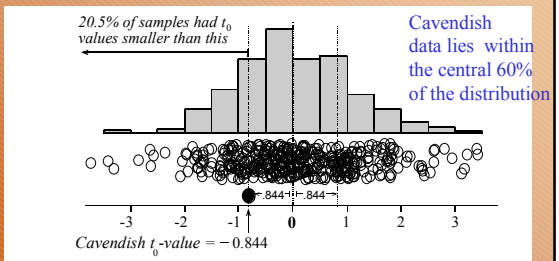


Figure 9.1.3 Sample t_0-values from 400 unbiased experiments (each t_0-value is distance between sample mean and 5.517 in std errors).

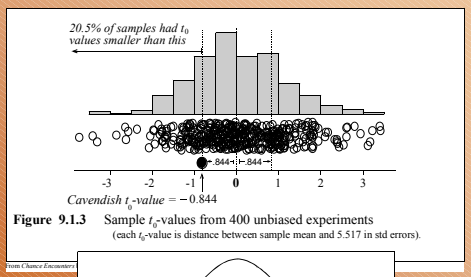


Figure 9.1.3 Sample t_0-values from 400 unbiased experiments (each t_0-value is distance between sample mean and 5.517 in std errors).

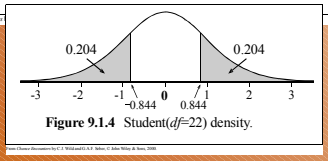


Figure 9.1.4 Student(df=22) density.

Measuring the distance between the true-value and the estimate in terms of the SE

- Intuitive criterion: Estimate is credible if it's not far away from its hypothesized true-value!
- But how far is far-away?
- Compute the distance in standard-terms:

$$T = \frac{\text{Estimator} - \text{TrueParameterValue}}{\text{SE}}$$

- Reason is that the distribution of ***T*** is known in some cases (Student's t, or N(0,1)). The estimator (obs-value) is typical/atypical if it is close to the center/tail of the distribution.

Comparing CI's and significance tests

- These are different methods for coping with the uncertainty about the true value of a parameter caused by the sampling variation in estimates.
- **Confidence interval**: A fixed level of confidence is chosen. We determine ***a range of possible values*** for the parameter that are consistent with the data (at the chosen confidence level).
- **Significance test**: *Only one possible value* for the parameter, called the hypothesized value, is tested. We determine the *strength of the evidence* (confidence) provided by the data against the proposition that the hypothesized value is the true value.

Review

- What intuitive criterion did we use to determine whether the hypothesized parameter value (p=0.2 in the ESP Example 9.1.1, and μ = 5.517 in Example 9.1.2) was credible in the light of the data? (Determine if the data-driven parameter estimate is consistent with the pattern of variation we'd expect get if hypothesis was true. If hypothesized value is correct, our estimate should not be far from its hypothesized true value.)
- Why was it that μ = 5.517 was credible in Example 9.1.2, whereas p=0.2 was not credible in Example 9.1.1? (The first estimate is consistent, and the second one is not, with the pattern of variation of the hypothesized true process.)

Review

- What do t_0-values tell us? (Our estimate is typical/atypical, consistent or inconsistent with our hypothesis.)
- What is the essential difference between the information provided by a confidence interval (CI) and by a significance test (ST)? (Both are uncertainty quantifiers. CI's use a fixed level of confidence to determine possible range of values. ST's one possible value is fixed and level of confidence is determined.)

Hypotheses

Guiding principles

We cannot **rule in** a hypothesized value for a parameter, we *can only* determine whether there is evidence *to* **rule out** a hypothesized value.

The ***null hypothesis*** tested is typically a skeptical reaction to a ***research hypothesis***

Comments

- Why can't we (**rule-in**) prove that a hypothesized value of a parameter is exactly true? (Because when constructing estimates based on data, there's always sampling and may be non-sampling errors, which are normal, and will effect the resulting estimate. Even if we do 60,000 ESP tests, as we saw earlier, repeatedly we are likely to get estimates like 0.2 and 0.200001, and 0.199999, etc. – non of which may be exactly the theoretically correct, 0.2.)
- Why use the rule-out principle? (Since, we can't use the rule-in method, we try to find compelling evidence against the observed/data-constructed estimate – to reject it.)
- Why is the null hypothesis & significance testing typically used? (H_a: skeptical reaction to a research hypothesis; ST is used to check if differences or effects seen in the data can be explained simply in terms of sampling variation!)

Comments

- How can researchers try to demonstrate that effects or differences seen in their data are real? (Reject the hypothesis that there are no effects)
- How does the alternative hypothesis typically relate to a belief, hunch, or research hypothesis that initiates a study? (H_1=H_a: specifies the type of departure from the null-hypothesis, H_0 (skeptical reaction), which we are expecting (research hypothesis itself).
- In the Cavendish's mean Earth density data, null hypothesis was $H_0 : \mu = 5.517$. We suspected bias, but not bias in any specific direction, hence $H_a: \mu != 5.517$.

Comments

- In the ESP Pratt & Woodruff data, (skeptical reaction) null hypothesis was $H_0 : \mu = 0.2$ (pure-guessing). We suspected bias, toward success rate being higher than that, hence the (research hypothesis) $H_a: \mu > 0.2$.
- Other commonly encountered situations are:
  - $H_0 : \mu_1 - \mu_2 = 0$ → $H_a : \mu_1 - \mu_2 > 0$
  - $H_0 : \mu_{rest} - \mu_{activation} = 0$ → $H_a : \mu_{rest} - \mu_{activation} != 0$

The t-test

Using $\hat{\theta}$ to test H_0: $\theta = \theta_0$ versus some alternative H_1.

STEP 1 Calculate the ***test statistic***,

$$t_0 = \frac{\hat{\theta} - \theta_0}{s\hat{e}(\hat{\theta})} = \frac{\text{estimate - hypothesized value}}{\text{standard error}}$$

[This tells us how many standard errors the estimate is above the hypothesized value (t_0 positive) or below the hypothesized value (t_0 negative).]

STEP 2 Calculate the *P*-value using the following table.

STEP 3 Interpret the *P*-value in the context of the data.

The t-test

Alternative hypothesis	Evidence against H_0: $\theta > \theta_0$ provided by	*P*-value
H_1: $\theta > \theta_0$	$\hat{\theta}$ too much bigger than θ_0 (i.e., $\hat{\theta} - \theta_0$ too large)	$P = \text{pr}(T \geq t_0)$
H_1: $\theta < \theta_0$	$\hat{\theta}$ too much smaller than θ_0 (i.e., $\hat{\theta} - \theta_0$ too negative)	$P = \text{pr}(T \leq t_0)$
H_1: $\theta \neq \theta_0$	$\hat{\theta}$ too far from θ_0 (i.e., $\lvert\hat{\theta} - \theta_0\rvert$ too large)	$P = 2\,\text{pr}(T \geq \lvert t_0 \rvert)$

where $T \sim \text{Student}(df)$

Interpretation of the p-value

TABLE 9.3.2 Interpreting the Size of a *P*-Value

Approximate size of *P*-Value		Translation
> 0.12	(12%)	**No** evidence against H_0
0.10	(10%)	**Weak** evidence against H_0
0.05	(5%)	**Some** evidence against H_0
0.01	(1%)	**Strong** evidence against H_0
0.001	(0.1%)	**Very Strong** evidence against H_0

Alternative Hypothesis	Evidence against H_0: $\theta = \theta_0$ provided by
H_1: $\theta > \theta_0$	$\hat{\theta}$ too much bigger than θ_0

Pictorial representation of the T-test
H_0: $\theta \hat{} = \theta_0$
H_1: $\theta > \theta_0$

$$t_0 = \frac{\hat{\theta} - \theta_0}{se(\hat{\theta})}$$

$\hat{\theta}$-scale → ***t*-scale (# of std errors)**

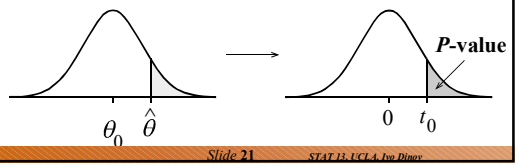


Figure 9.3.1: Testing H_0: $\theta = \theta_0$

Alternative Hypothesis	Evidence against H_0: $\theta = \theta_0$ provided by
H_1: $\theta < \theta_0$	$\hat{\theta}$ too much smaller than θ_0

Pictorial representation of the T-test
H_1: $\theta < \theta_0$

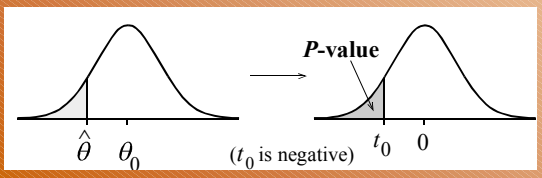


Figure 9.3.1: Testing H_0: $\theta = \theta_0$

Alternative Hypothesis	Evidence against H_0: $\theta = \theta_0$ provided by
H_1: $\theta \neq \theta_0$ **(2-sided)**	$\hat{\theta}$ too far from θ_0 (either direction)

Pictorial representation of the T-test
H_1: $\theta \mathrel{!}= \theta_0$

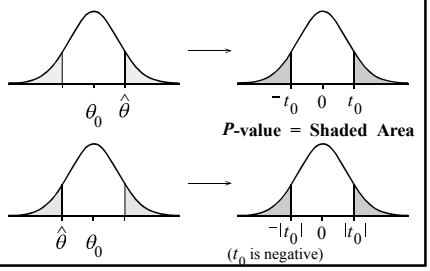
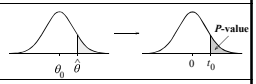
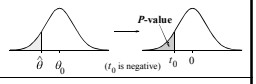
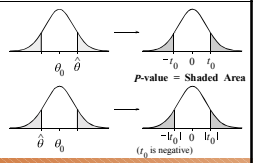


Figure 9.3.1: Testing H_0: $\theta = \theta_0$ (Pictorial version of Table 9.1.1)

$$t_0 = \frac{\hat{\theta} - \theta_0}{se(\hat{\theta})}$$

Alternative Hypothesis	Evidence against H_0: $\theta = \theta_0$ provided by	$\hat{\theta}$-scale → *t*-scale (# of std errors)
H_1: $\theta > \theta_0$	$\hat{\theta}$ too much bigger than θ_0	
H_1: $\theta < \theta_0$	$\hat{\theta}$ too much smaller than θ_0	
H_1: $\theta \neq \theta_0$ (2-sided)	$\hat{\theta}$ too far from θ_0 (either direction)	

P-values from *t*-tests

- The ***P-value*** is the probability that, *if the hypothesis was true*, sampling variation would produce an estimate that is further away from the hypothesized value than our data-estimate.
- The ***P-value*** measures the strength of the evidence against H_0.
- The ***smaller*** the *P-value*, the ***stronger*** the evidence against $\boldsymbol{H_0}$.

(The second and third points are true for significance tests generally, and not just for *t*-tests.)

Review

- What does the *t*-statistic tell us?

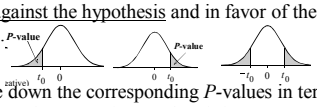
  The T-statistics, $t_0 = \frac{\hat{\theta} - \theta_0}{se(\hat{\theta})}$ tells us (in std. units) if the observed value/estimate is typical/consistent and can be explained by the variation in the sampling distribution.

- When do we use a 2-tailed rather than a 1-tailed test?

  We use two-sided/two-tailed test, unless there is a prior (knowledge available before data was collected) or a strong reason to believe that the result should go in one particular direction (← μ →).

Review

- What were the 3 types of alternative hypothesis involving the parameter θ and the hypothesized value θ_0? Write them down!
- Let's go through and construct our own *t-Test* Table.
  - For each alternative, think through what would constitute evidence against the hypothesis and in favor of the alternative.



  - Then write down the corresponding *P*-values in terms of t_0 and represent these *P*-values on hand-drawn curves (cf. Fig. 9.3.1). [P=Pr(T>=t_0), P=Pr(T<=t_0), P=2Pr(T>=|t_0|) .]

Review

- What does the *P*-value measure? (If H_0 was true, sampling variation alone would produce an estimate farther then the hypothesized value.)
- What do very small *P*-values tell us? What do large *P*-values tell us? (strength of evidence against H_0 .)
- Pair the phrases: "the ↑|↓ ..... the *P*-value, the ↑|↓ ..... the evidence for/against ..... the null hypothesis."
- Do large values of t_0 correspond to large or small *P*-values? Why?
- What is the relationship between the Student (*df*) distribution and Normal(0,1) distribution? (identical as $n \to \infty$)

Is a second child gender influenced by the gender of the first child, in families with >1 kid?

TABLE 9.3.4 First and Second Births by Sex

		Second Child		
		Male	Female	Total
First Child	Male	3,202	2,776	5,978
	Female	2,620	2,792	5,412
	Total	5,822	5,568	11,390

- Research hypothesis needs to be formulated first before collecting/looking/interpreting the data that will be used to address it. Mothers whose 1st child is a girl are more likely to have a girl, as a second child, compared to mothers with boys as 1st children.
- Data: 20 yrs of birth records of 1 Hospital in Auckland, NZ.

Analysis of the birth-gender data – data summary

	Second Child	
Group	Number of births	Number of girls
1 (Previous child was girl)	5412	2792 (approx. 51.6%)
2 (Previous child was boy)	5978	2776 (approx. 46.4%)

- Let p_1=true proportion of girls in mothers with girl as first child, p_2=true proportion of girls in mothers with boy as first child. Parameter of interest is $p_1 - p_2$.
- H_0: $p_1 - p_2 = 0$ (skeptical reaction). H_a: $p_1 - p_2 > 0$ (research hypothesis)

Hypothesis testing as decision making

TABLE 9.4.1 Decision Making

	Actual situation	
Decision made	H_0 is true	H_0 is false
Accept H_0 as true	OK	Type II error
Reject H_0 as false	Type I error	OK

- Sample sizes: n_1=5412, n_2=5978, Sample proportions (estimates) $\hat{p}_1 = 2792/5412 \approx 0.5159$, $\hat{p}_2 = 2776/5978 \approx 0.4644$,
- H_0: $p_1 - p_2 = 0$ (skeptical reaction). H_a: $p_1 - p_2 > 0$ (research hypothesis)

Analysis of the birth-gender data

- Samples are large enough to use Normal-approx. Since the two proportions come from totally diff. mothers they are independent → use formula 8.5.5.a

$$t_0 = \frac{\text{Estimate} - \text{HypothesizedValue}}{SE} = 5.49986 = \frac{\hat{p}_1 - \hat{p}_2 - 0}{SE\left(\hat{p}_1 - \hat{p}_2\right)} = \frac{\hat{p}_1 - \hat{p}_2}{\sqrt{\frac{\hat{p}_1(1-\hat{p}_1)}{n_1} + \frac{\hat{p}_2(1-\hat{p}_2)}{n_2}}} =$$

$$P - value = \Pr(T \geq t_0) = 1.9 \times 10^{-8}$$

Analysis of the birth-gender data

- We have strong evidence to reject the H_0, and hence conclude mothers with first child a girl a more likely to have a girl as a second child.
- How much more likely? A 95% CI:

CI (p_1- p_2) =[0.033; 0.070]. And computed by:

$$\text{estimate} \pm z \times \text{SE} = \hat{p}_1 - \hat{p}_2 \pm 1.96 \times SE\left(\hat{p}_1 - \hat{p}_2\right) =$$

$$\hat{p}_1 - \hat{p}_2 \pm 1.96 \times \sqrt{\frac{\hat{p}_1(1-\hat{p}_1)}{n_1} + \frac{\hat{p}_2(1-\hat{p}_2)}{n_2}} =$$

$$0.0515 \pm 1.96 \times 0.0093677 = [3\%; 7\%]$$

Review

- What is the difference between fixed-level hypothesis testing and the *P*-value approach?
  - Fisher's P-value approach: P-value is considered in context
    - P-value is considered as evidence against H_0
    - The smaller the P-value → the stronger the evidence against H_0
  - Neyman-Pearson approach: fixed-level hypothesis testing
    - For a fixed significance level α, the hypothesis testing procedure is called testing at α -level.
    - When H_0 is true, P-values <= α occur purely by chance only α % of the time (e.g., testing at significance level α =5%).

Review

- For fixed-level hypothesis tests, why are low significance levels chosen? (Large levels would imply we falsely reject the skeptical null hypothesis too often and commit Type I error! On the contrary, if our significance level is too low, preventing to reject H_0, we may end up being unlikely to reject important false hypotheses – Type II error.)
- If you wanted to perform a fixed-level hypothesis test, for what values of the *P*-value would you "reject the null hypothesis at the 1% level" ? (P<0.01)
- Give another name for the significance level. (Probability of making Type I error)

Review

- If 120 researchers each independently investigated a it true/ hypothesis, how many researchers would you expect to obtain a result that was significant at the 5% level (just by chance)? (Type I, false-positive; 120*5%=6)
- What was the other type of error described? What was it called? When is the idea useful? (Type II, false-negative)
- Power of statistical test = 1-β, where

β =P(Type II error) = P(Accepting Ho as true, when its truly false)

Review

- Why is the expression "accept the null hypothesis" dangerous? (Data can not really provide all the evidence that a hypothesis is true, however, it can provide support that it is false. That's why better lingo is "we can't reject H_0")
- What is meant by the word ***non-significant*** in many research literatures? (P-value > fixed-level of significance)
- In fixed-level testing, what is a Type I error? What is a Type II error? (Type I, false-positive, reject H_0 as false, when it's true in reality; Type II, false-negative, accepting H_0 as true, when its truly false)

Tests and confidence intervals

A ***two-sided*** test of H_0: $\theta = \theta_0$ is ***significant*** at the 5% level **if and only if** θ_0 lies ***outside*** a 95% confidence interval for θ.

A ***two-sided*** test of H_0: $\theta = \theta_0$ gives a result that is significant at the 5% level **if** the P-value=2Pr(T >=$|t_0|$) < 0.05. Where t_0 =(estimate-Hypoth'dValue)/SE(θ) ➔ $t_0 = (\hat{\theta} - \theta_0)/SE(\theta)$. Let **t** be a **threshold** chosen so that Pr(T>= **t**) = 0.025. Now $|t_0|$ tells us now many SE's $\hat{\theta}$ *and* θ are apart (without direction in their diff.) If $|t_0|$> **t**, then θ_0 is more than **t** SE's away from $\hat{\theta}$ and hence lies outside the 95% CI for θ.

"Significance"

- ***Statistical significance*** relates to the strength of the evidence of ***existence*** of an effect.
- The ***practical significance*** of an effect depends on its size – how large is the effect.
- A small *P*-value provides evidence that the effect ***exists*** but says ***nothing*** at all about the ***size*** of the effect.
- To estimate the ***size*** of an effect (its practical significance), ***compute a confidence interval.***

"Significance" cont.

A non-significant test does not imply that the null hypothesis is true (or that we accept H_0).

It simply means we do not have (this data does not provide) the evidence to reject the skeptical reaction, H_0 .

To prevent people from misinterpreting your report: ***Never quote a P-value*** about the existence of an effect ***without*** also ***providing a confidence interval*** estimating the size of the effect.

Review

- What is the relationship between a 95% confidence interval for a parameter θ and the results of a two-sided test of H_0: $\theta = \theta_0$? (θ_0 *is inside the 95% CI(θ)* , ➔ *P-value for the test is >0.025,. Conversely, the test is significan, at 5%-level,* ➔ θ_0 *is outside the 95% CI(θ)).*
- If you read, "research shows that ..... θ is significantly bigger than ..... θ_0 .", what is a likely explanation? (there is evidence that a real effect exists to make the two values different).
- If you read, "research says that ..... drug makes no difference to ..... disease treatment", what is a likely explanation? (the data does not have the evidence to reject the skeptical reaction, H_0, or no effects).

Review

- Is a "significant difference" necessarily large or practically important? Why? (No, significant difference indicates the existence of an effect, practical importance depends on the effect-size.)
- What is the difference between statistical significance and practical significance? (stat-significance relates to the strength of the evidence that a real effect exists (e.g., that true difference is not exact;y 0); practical significance indicates how important the observed difference is in practice, how large is the effect.)
- What does a *P*-value tell us about the size of an effect? (P-value says whether the effect is significant, but says nothing about its size.)
- What tool do we use to gauge the size of an effect? (CI(parameter) provides clues to the size of the effect.)

Review

- If we read that a difference between two proportions is *non-significant*, what does this tell us? What does it not tells us? (Do not have evidence proportions are different, based on this data. Doesn't mean accept H_0).
- What is the closest you can get to showing that a hypothesized value is true and how could you go about it? (Suppose, H_0: $\theta = \theta_0$, and our test is not-significant. To show $\theta = \theta_0$ we need to show that all values in the CI(θ_0) are essentially equal to θ_0 , this is a practical subjective matter decision, not a statistical one.)

General ideas of "test statistic" and "*p*-value"

A ***test statistic*** is a measure of discrepancy between what we see in data and what we would expect to see if H_0 was true.

The ***P-value*** is the probability, calculated assuming that the null hypothesis is true, that sampling variation alone would produce data which is more discrepant than our data set.

Chapter 9 Summary

Significance Tests vs. Confidence Intervals

- The main use of significance testing is to check whether apparent differences or effects seen in data can be explained away simply in terms of sampling variation. The essential **difference between confidence intervals and significance tests** is as follows:
  - *Confidence interval* : A range of possible values for the parameter are determined that are consistent with the data at a specified confidence level.
  - *Significance test* : Only one possible value for the parameter, called the hypothesized value, is tested. We determine the strength of the evidence provided by the data against the proposition that the hypothesized value is the true value.

Hypotheses

- The ***null hypothesis***, denoted by H_0, is the (skeptical reaction) hypothesis tested by the statistical test.
- *Principle guiding the formulation of null hypotheses*: We cannot rule a hypothesized value in; we can only determine whether there is enough evidence to rule it out. Why is that?
- ***Research (alternative) hypotheses*** lay out the conjectures that the research is designed to investigate and, if the researchers hunches prove correct, establish as being true.

Example: Is there racial profiling or are there confounding explanatory effects?!?

- The book by Best (*Damned Lies and Statistics: Untangling Numbers from the Media, Politicians and Activists*, Joel Best) shows how we can test for racial bias in police arrests. Suppose we find that among 100 white and 100 black youths, 10 and 17, respectively, have experienced arrest. This may **look plainly discriminatory**. But suppose we then find that of the 80 middle-class white youths 4 have been arrested, and of the 50 middle-class black youths 2 arrested, whereas the corresponding numbers of lower-class white and black youths arrested are, respectively, 6 of 20 and 15 of 50. These arrest rates correspond to 5 per 100 for white and 4 per 100 for black middle-class youths, and 30 per 100 for both white and black lower-class youths. Now, better analyzed, the data suggest **effects of social class, not race as such**.

Hypotheses cont.

- The ***null hypothesis*** tested is typically a skeptical reaction to the research hypothesis.
- The most commonly tested null hypotheses are of the "it makes no difference" variety.
- Researchers try to demonstrate the existence of real treatment or group differences by showing that the idea that there are no real differences is implausible.
- The ***alternative hypothesis*** , denoted by H_1, specifies the type of departure from the null hypothesis, H_0, that we expect to detect.

Hypotheses cont.

- The ***alternative hypothesis***, typically corresponds to the research hypothesis.
- We use ***one-sided alternatives*** (using either : H_1: $\theta > \theta_0$ or H_1: $\theta < \theta_0$) when the research hypothesis specifies the direction of the effect, or more generally, when the investigators had good grounds for believing the true value of θ was on one particular side of θ_0 before the study began. Otherwise a ***two-sided alternative***, $H_1 : \theta \neq \theta_0$, is used.

P-values

- Differences or effects seen in data that are easily explainable in terms of sampling variation do not provide convincing evidence that real differences or effects exist.
- The ***P*-value** is the probability that, if the hypothesis was true, sampling variation would produce an estimate that is further away from the hypothesized value than the estimate we got from our data.
- The *P*-value measures the strength of the evidence against H_0.

P-values cont.

- The *smaller* the *P*-value, the stronger the evidence against H_0.
- A large *P*-value provides no evidence against the null hypothesis.
- A large *P*-value does *not* imply that the null hypothesis is true.
- A small *P*-value provides evidence that the effect exists but says *nothing* at all about the *size* of the effect.
- To estimate the **size** of an effect, *compute a confidence interval.*

P-values cont.

- Never quote a *P*-value about the existence of an effect without also providing a confidence interval estimating the size of the effect.
- Suggestions for ***verbal translation of P-values*** are given in Table 9.3.2.
- ***Computation of P-values*** : Computation of P-values for situations in which the sampling distribution of $(\hat{\theta} - \theta_0)/se(\hat{\theta})$, is well approximated by a Student(*df*) distribution or a Normal(0,1) distribution is laid out in Table 9.3.1.
- The *t*-test statistic tells us how many standard errors the estimate is from the hypothesized value.

P-values

- Examples given in this chapter concerned means and differences between means, proportions and differences between proportions.
- In general, a test statistic is a measure of discrepancy between what we see in the data and what we would have expected to see if H_0 was true.

Significance

- If, whenever we obtain a *P*-value less than or equal to 5%, we make a decision to reject the null hypothesis, this procedure is called ***testing at the 5% level of significance***.
  - The significance level of such a test is 5%.
- If the *P*-value $\leq \alpha$, the effect is said to be significant at the α-level.
- If you always test at the 5% level, you will reject one true null hypothesis in 20 over the long run.

Significance cont.

- A two-sided test of $H_0 : \theta = \theta_0$ is significant at the 5% level if and only if θ_0 lies outside a 95% confidence interval for θ.
- In reports on research, the word "significant" used alone often means "significant at the 5% level" (i.e. P-value ≤ 0.05). "Non-significant", "does not differ significantly" and even "is no different" often mean P-value > 0.05.
- A non-significant result does not imply that H_0 is true.

Significance cont.

- A Type I error (false-positive) is made when one concludes that a true null hypothesis is false.
- The significance level is the probability of making a Type I error.
- ***Statistical significance*** relates to having evidence of the ***existence*** of an effect.
- The ***practical significance*** of an effect depends on its ***size***.