Lecture 8: Testing hypotheses continued

Lecturer: Dominik Rothenhäusler

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Abstract

These notes are mnemonics about what was covered in class. They don't replace being present or reading the book. Reading ahead in the book is very effective.

8.1 Neyman-Pearson lemma

Here we look at an optimality property of the likelihood ratio test for simple versus simple hypothesis tests. The likelihood ratio for $H_0: \theta = \theta_0$ versus $H_A: \theta = \theta_1$ is

$$\lambda = \frac{f_0(x_1, \dots, x_n)}{f_A(x_1, \dots, x_n)}.$$

For IID data we get

$$\lambda = \frac{\prod_{i=1}^{n} f(x_i; \theta_0)}{\prod_{i=1}^{n} f(x_i; \theta_1)}.$$

Because both hypotheses are simple we have an expression for both numerator and denominator. If one of them would be composite then that probability would ordinarily depend on which θ from that composite hypothesis was true.

The likelihood ratio test rejects H_0 if $\lambda \leq \lambda^*$ where λ^* is a threshold that we choose. If we use $\lambda^* = 1$ we reject H_0 if H_A has higher likelihood. In a Bayesian analysis the threshold could be adjusted to account for different prior probabilities of θ_0 and θ_1 or to account for different losses. In the Neyman-Pearson setup we choose λ^* to get a desired type I error probability.

With our chosen λ^* , we get type I error probability

$$\alpha^* = \Pr(\lambda(X) \leqslant \lambda^*; H_0)$$

and type II error probability

$$\beta^* = \Pr(\lambda(X) > \lambda^*; H_A)$$

Here we follow Rice by writing $\lambda(X)$ letting X be "all our data".

The Neyman-Pearson lemma is as follows. If any other test based on the same X has $\alpha \leqslant \alpha^*$ then that test has $\beta \geqslant \beta^*$. The power of a test is $1-\beta$ so the alternative test then has power no better than the likelihood ratio test has. Lets assume that $0 < \lambda^* < \infty$ to rule out unimportant corner cases.

Proof. We prove it for a continuously distributed X. Let the other test reject when $x \in C$ and fail to reject when $x \in \overline{C}$ (the complement of C). By the definition of the LR test, we have

$$\alpha^* = \int_{x:\lambda(x) \leq \lambda^*} f_0(x) dx$$
 and $\beta^* = \int_{x:\lambda(x) > \lambda^*} f_A(x) dx$.

Now the difference in type II error probabilities is

$$\beta - \beta^* = \int_{\bar{C}} f_A(x) \, \mathrm{d}x - \int_{\lambda > \lambda^*} f_A(x) \, \mathrm{d}x.$$

Watch the small changes in the next steps. They are standard manipulations to drive the expression towards something we know. First, we can remove $\bar{C} \cap \lambda > \lambda^*$ from both integral domains. That is where both tests fail to reject under H_A . This brings

$$\beta - \beta^* = \int_{\bar{C} \cap \lambda \leqslant \lambda^*} f_A(x) \, \mathrm{d}x - \int_{C \cap \lambda > \lambda^*} f_A(x) \, \mathrm{d}x$$

$$\geqslant \frac{1}{\lambda_*} \left(\int_{\bar{C} \cap \lambda \leqslant \lambda^*} f_0(x) \, \mathrm{d}x - \int_{C \cap \lambda > \lambda^*} f_0(x) \, \mathrm{d}x \right)$$

$$= \frac{1}{\lambda_*} \left(\int_{\bar{C} \cap \lambda \leqslant \lambda^*} f_0(x) \, \mathrm{d}x + \int_{C \cap \lambda \leqslant \lambda^*} f_0(x) \, \mathrm{d}x - \int_{C \cap \lambda \leqslant \lambda^*} f_0(x) \, \mathrm{d}x - \int_{C \cap \lambda > \lambda^*} f_0(x) \, \mathrm{d}x \right)$$

$$= \frac{1}{\lambda_*} \left(\int_{\lambda \leqslant \lambda^*} f_0(x) \, \mathrm{d}x - \int_{C} f_0(x) \, \mathrm{d}x \right)$$

$$= \frac{1}{\lambda_*} (\alpha - \alpha^*)$$

$$\geqslant 0.$$

No other test can beat the LR test on both α and β .

8.2 Example and a uniformly most powerful test

Suppose we have an exponential random variable with PDF $f(x;\theta) = (1/\theta)e^{-x/\theta}$. Consider $H_0: \theta = 1$ and $H_A: \theta = 3$. The LR test rejects if

$$\frac{e^{-x}}{(1/3)e^{-x/3}} < \lambda^*. (8.1)$$

We can rearrange that to say we reject H_0 if

$$x > x^* = -\frac{3}{2}\log(\lambda^*/3).$$

That is, whatever λ^* we choose corresponds to some x^* where we will reject H_0 if $x > x^*$. Using the exponential distribution we find that $\Pr(X > x^*; H_0) = \exp(-x^*)$. So to get a desired level α we set $\alpha = \exp(-x^*)$, that is $x^* = -\log(\alpha)$.

Knowing x^* we could if we wanted work backwards, plugging $x^* = -\log(\alpha)$ into (8.1) to find the critical value λ^* . However we already have the test in terms of x^* and that is more convenient to use.

The LR test is our most powerful α -level test for $H_0: \theta = 1$ versus $H_A: \theta = 3$ by the Neyman-Pearson lemma. It is also the most powerful α -level test for $H_0: \theta = 1$ versus $H_A: \theta = 7$. The same holds for any H_A

with a θ value larger than 1. We never used the alternative value of θ when constructing the threshold x^* . This test rejecting $H_0: \theta = 1$ in favor of $H_A: \theta = \theta_1$ rejects when $x > -\log(\alpha)$ no matter what $\theta_1 > 1$ we use. It is a **uniformly most powerful** (UMP) test because it has that property over a range of θ values.

A different test is uniformly most powerful over $H_A: \theta = \theta_1$ for $\theta_1 < \theta_0$. That test rejects for small x.

UMP tests are rare and special. We don't ordinarily have them.

8.3 A few facts about *p*-values

There is a lot of important work on p-values lately. It is worth adding a small segment about them. First, if the test statistic T(X) has a continuous distribution then the p-value is uniformly distributed on (0,1). It was constructed to have a 5% chance of being below 0.05 a 1% chance of being below 0.01 and so on.

To see formally that it is uniformly distributed, the observed value of p is $p = \Pr(T(X) \ge T(x); H_0)$. Let F be the CDF of T(X) under H_0 . Then we can write the observed p-value as p = 1 - F(T(x)). It is then the observed value of a random variable P = 1 - F(T(X)). Now

$$\Pr(P \leq \alpha; H_0) = \Pr(1 - F(T(X)) \leq \alpha; H_0)$$

$$= \Pr(F(T(X)) \geq 1 - \alpha; H_0)$$

$$= 1 - \Pr(F(T(X)) \leq 1 - \alpha; H_0)$$

$$= 1 - \Pr(T(X) \leq F^{-1}(1 - \alpha); H_0)$$

$$= 1 - F(F^{-1}(1 - \alpha))$$

$$= 1 - (1 - \alpha)$$

$$= \alpha,$$

and so $P \sim U(0,1)$.

Suppose that we test some hypothesis H_0 k times getting k independent p-values, p_j for j = 1, ..., k. This is the same hypothesis being tested k times. It might be about whether a dietary intervention reduces blood pressure. If we want to combine those k p-values into a single p-value for H_0 there are several ways to go about it. Doing this is called **meta-analysis**. It is a study of studies. We could look for a small value of

$$\min_{1 \le j \le k} p_j, \quad \max_{1 \le j \le k} p_j, \quad \text{or} \quad \prod_{1 \le j \le k} p_j$$

among many others. The latter choice was studied by Fisher and is usually favored. It has a nice distributional property. Under $H_0 - 2\log(\prod_j p_j) = -2\sum_j \log(p_j) \sim \chi_{(2k)}^2$. Under the alternative this product is even smaller than a $\chi_{(2k)}^2$ random variable. So we reject if $\prod_j p_j$ is small, or equivalently if $-2\sum_j \log(p_j)$ is larger than the $1-\alpha$ quantile of the $\chi_{(2k)}^2$ distribution. Here is a sketch of how that distribution is derived. First show directly that $-\log(U(0,1))$ has a standard exponential distribution. Then realize that is a special case of a Gamma distribution. Then note that χ^2 is a special case of the Gamma distribution. The whole reason for multiplying the logarithm by 2 is to turn the Gamma distribution into one of the χ^2 special cases.

There is a second multiple p-value issue. Suppose now that we have k different null hypotheses H_{0j} for $j=1,\ldots,k$. The example from class was based on a famous XKCD cartoon https://xkcd.com/882/. Somebody tests for a link between jelly beans and acne and finds there is not one at $\alpha=0.05$. They then run separate tests on purple, brown, pink, blue, \cdots , orange, doing p-values 20 in all. Of those, one is statistically significant at the 0.05 level (green). That one makes headlines and the others are ignored. If

all k null hypotheses are true and independent and we test them at the α level then the number of false discoveries we get is $Bin(k,\alpha)$. The expected number of false discoveries is $k \times \alpha$ and the probability of one or more false discoveries is

$$1 - (1 - \alpha)^k$$
.

This can be very close to 1 if k is large. There are many ways to contend with the multiple testing issue. The simplest is to do k tests at level α/k each. That makes the probabilty of any false discovery be at most α . To see this let $Z_j = 1$ if the j'th test brings a false discovery and 0 otherwise. Then under all k null hypotheses

$$\Pr(\text{any false discoveries}) = \Pr(\max_{1 \leqslant j \leqslant k} Z_j = 1)$$

$$= \mathbb{E}(\max_{1 \leqslant j \leqslant k} Z_j)$$

$$\leqslant \mathbb{E}(\sum_{1 \leqslant j \leqslant k} Z_j)$$

$$= \sum_{1 \leqslant j \leqslant k} \mathbb{E}(Z_j)$$

$$= \sum_{1 \leqslant j \leqslant k} \alpha/k$$

$$= \alpha$$

This approach of dividing the signifiance level α by the number k of tests being performed is known as the **Bonferroni** method.

8.4 Confidence intervals

Let X be all our data and let θ be the parameter of interest. A $100(1-\alpha)\%$ confidence interval for θ is an interval of the form [L(X), U(X)] where

$$\Pr(L(X) \leqslant \theta \leqslant U(X); \theta) = 1 - \alpha$$

holds for all θ . For an IID sample this is

$$\Pr(L(X_1,\ldots,X_n)\leqslant\theta\leqslant U(X_1,\ldots,X_n);\theta)=1-\alpha.$$

It is important to note that it is L and U that are random while θ is a fixed non-random value. So $\Pr(\cdots)$ refers to the X's inside.

We saw in class that a confidence interval can be turned into a test. If θ_0 is outside the confidence interval, reject $H_0: \theta = \theta_0$ in favor of $H_A: \theta \neq \theta_0$. Similarly, a test of $H_0: \theta = \theta_0$ can be turned into a **confidence set** made up of all the 'unrejected' θ values. We say confidence 'set' because the result might not be an interval. It would still have the coverage properties. We proved one of these equivalences in class; both are in the text.

To show you that confidence intervals exist, consider $X_i \sim N(\mu, \sigma^2)$ with μ unknown and σ somehow known. Then

$$\Pr\left(\bar{X} - 2.58 \frac{\sigma}{\sqrt{n}} \leqslant \mu \leqslant \bar{X} + 2.58 \frac{\sigma}{\sqrt{n}}\right) = 0.99,$$

so we get a confidence interval. If we don't know σ then we can plug in an estimate $\hat{\sigma}$ and (if it is a good estimate) get

$$\Pr\left(\bar{X} - 2.58 \frac{\hat{\sigma}}{\sqrt{n}} \leqslant \mu \leqslant \bar{X} + 2.58 \frac{\hat{\sigma}}{\sqrt{n}}\right) \doteq 0.99.$$

Later we will get something even more precise using the t distribution.

In the method of moments, the central limit theorem gives

$$\Pr\left(\bar{X} - 2.58 \frac{\hat{\sigma}}{\sqrt{n}} \leqslant \mathbb{E}(X) \leqslant \bar{X} + 2.58 \frac{\hat{\sigma}}{\sqrt{n}}\right) \doteq 0.99$$

if $\hat{\sigma}$ is good enough to consistently estimate $\sqrt{\operatorname{Var}(X)}$.

Using Fisher information we get

$$\Pr\left(\hat{\theta} - \frac{2.58}{\sqrt{nI(\theta_0)}} \leqslant \theta \leqslant \hat{\theta} + \frac{2.58}{\sqrt{nI(\theta_0)}}\right) \doteq 0.99.$$

If we don't know $I(\theta_0)$ we can usually plug in $I(\hat{\theta})$ (more later in the course). We can reject $H_0: \theta = \theta_0$ at the $\alpha = 0.01$ level if

$$|\hat{\theta} - \theta_0| > 2.58/\sqrt{nI(\theta_0)}.$$

8.5 Generalized likelihood ratio test

For composite hypotheses we generalize the likelihood ratio test. We reject H_0 in favor of H_A if

$$\Lambda = \frac{\max_{\theta \in H_0} L(\theta)}{\max_{\theta \in H_0 \cup H_A} L(\theta)} \leqslant \lambda^*.$$

The numerator is the likelihood under H_0 and the denominator is the likelihood under an even larger set of possibilities. So clearly $\Lambda \leq 1$. It has to be enough less than 1 for rejection to be meaningful. We will see how much less.

Suppose that X_i describe the number of copies of a gene that somebody gets. The Hardy-Weinberg model has $\Pr(X_i = 0) = \theta^2$, $\Pr(X_i = 1) = 2\theta(1 - \theta)$ and $\Pr(X_i = 2) = \theta^2$. It is a Bin(2, θ) model as if their genes were independent random Bernoulli variables one from each parent. In the Hardy-Weinberg model we might want to test $\theta = \theta_0$ for some hypothesized value of θ_0 (maybe 1/2) versus $\theta \neq \theta_0$. We might also want to test the Hardy-Weinberg model itself against the more general model

$$\Pr(X = x) = \begin{cases} \theta_0, & x = 0\\ \theta_1, & x = 1\\ 1 - \theta_0 - \theta_1, & x = 2. \end{cases}$$