How to use "Goodness of Fit Tests?"

Paweł Polak

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STAT W4413: Nonparametric Statistics - Lecture 13

χ^2 test: example

Example (Gibbons and Chakraborti)

A quality control engineer has taken 50 samples of size 13 from a production process. The numbers of defectives for these samples are recorded below. Test the null hypothesis at significance level $\alpha=0.05$. (a) Perform the χ^2 test when the null is Binom(0.1). (b) Perform the χ^2 test when the null is Binom(p) and p is not known.

Number of defects	Number of samples
0	10
1	24
2	10
3	4
4	1
5 or more	1

Part (a)

Define $p_i = P(Number of defects = i)$. The first step is to estimate \hat{p}_i .

Number of defects	Number of samples
0	10
1	24
2	10
3	4
4	1
5 or more	1

Total number of samples = 50

Therefore,

Number of defects	р̂
0	.2
1	.48
2	.2
3	.08
4	.02
5 or more	.02

Part (a); cont'd

How to calculate π_i (the probability under null)? From Binomial distribution.

$$\pi_i = \binom{13}{i} p^i (1-p)^{13-i} = \binom{13}{i} 0.1^i (0.9)^{13-i}.$$

number of defects	\hat{p} (from observations)	π (from binomial)
0	.2	0.2542
1	0.48	0.3671
2	.2	0.2448
3	.08	0.0997
4	.02	0.0277
5 or more	.02	.0065

Now we should calculate $Q = \sum_{i=0}^{5} \frac{(\hat{p}_i - \pi_i)^2}{\pi_i} = 0.0885$.

$$nQ \stackrel{d}{\rightarrow} \chi^2(5)$$
. Therefore we have

$$> \mathsf{kappa} < - \mathsf{qchisq}(0.95,5)$$

> kappa

[1]11.0705

nQ = 4.425. Therefore we accept the null hypothesis.

Part (b)

(b) Perform the χ^2 test when the null is Binom(p) and p is not known.

We have a composite null hypothesis. The first step is to estimate p. To do so we first calculate the likelihood:

$$P(X_1 = k_1, X_2 = k_2, \dots, X_{50} = k_{50})$$

$$= \binom{n}{k_1} \binom{n}{k_2} \dots \binom{n}{k_{50}} p^{k_1 + k_2 + \dots + k_{50}} (1 - p)^{(13 - k_1) + (13 - k_2) + \dots + (13 - k_{50})}.$$

 $k_1+k_2+\ldots+k_{50}$ is the total number of defects and here it is equal to

$$24 \times 1 + 10 \times 2 + 4 \times 3 + 4 \times 1 + 5 \times 1 = 65.$$

$$\frac{d}{dp}\log P(X_1=k_1,X_2=k_2,\ldots,X_{50}=k_{50})=0\Rightarrow \frac{65}{p}=\frac{650-65}{1-p}\Rightarrow \hat{p}=0.1.$$

Therefore, our Q is the same as before.

However, $nQ \stackrel{d}{\to} \chi^2(4)$. (You lose one df because you estimate one parameter!)

Therefore, we have

- > kappa < qchisq(0.95,4)
- > kappa
- [1]9.487729 Still the null hypothesis is accepted since nQ = 4.425.

Test for Continuous CDFs

Example

When a customer arrives in a bank it takes the bank X minutes to serve him/her. One of the most well-known models for the waiting times is called M/M/1 in which we assume that the distribution of the waiting times for the customers is exponential. As a statistician you would like to check and see if this model works for the bank you are working for. For this purpose, you have collected information about the waiting time of 30 customers and you have obtained

$$x \leftarrow c(0.6, 1.6, 3.2, 3.3, 3.5, 3.8, 5.9, 6.2, \dots$$

 $6.8, 11.3, 11.9, 13.5, 15.1, 15.7, 16.2, \dots$
 $16.2, 18.5, 18.7, 20.7, 22.0, 23.1, 23.8, \dots$
 $23.9, 26.4, 27.9, 37.4, 39.6, 40.0, 60.4, 63.0)$

Based on these numbers you would like to judge wether the exponential model with parameter $\lambda=1/20$ is right or not.

qqplot

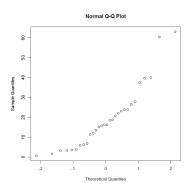
$$px < - seq(0,1,1/30)$$

y < $-qexp(px, rate = 1/20)$

qqplot(x,y)

qqplot for exponential hypothesis

qqnorm(x)



Case I: One-Sample, Simple Null

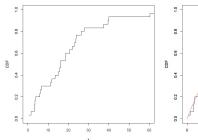
We know the exact null distribution:

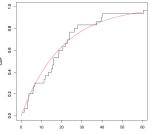
$$H_0: F(x) = 1 - e^{-x/20} \text{ for } x > 0 \quad \text{ versus } \quad H_1: F(x) \neq 1 - e^{-\frac{x}{20}}$$

Empirical CDF versus null CDF

Plotting CDF:

```
\begin{aligned} & \mathsf{plot}(\mathsf{sort}(\mathsf{x}), \ (1:30)/30, \ \mathsf{xlim} = \mathsf{c}(0,60), \ \mathsf{ylim} = \mathsf{c}(0,1), \ \mathsf{type} = "\mathsf{s}", \ \mathsf{xlab} = \\ & "\mathsf{x}", \ \mathsf{ylab} = "\mathsf{CDF}") \end{aligned} & \mathsf{par}(\mathsf{new} = \mathsf{T}) & \mathsf{z} < - \ \mathsf{seq}(0,60, \ .1) & \mathsf{plot}(\mathsf{z}, \ 1 - \ \mathsf{exp}(-\mathsf{z}/20), \ \mathsf{type} = "\mathsf{l}", \ \mathsf{xlim} = \mathsf{c}(0,60), \ \mathsf{ylim} = \mathsf{c}(0,1), \ \mathsf{xlab} = \\ & "\mathsf{x}", \ \mathsf{ylab} = "\mathsf{CDF}") \end{aligned}
```





Kolmogorov-Smirnov Test

$$H_0: F(x) = 1 - e^{-x/20} \text{ for } x > 0 \quad \text{ versus } \quad H_1: F(x) \neq 1 - e^{-\frac{x}{20}}$$

R code:

• ks.test(x,"pexp",rate=1/20)

Result:

One-sample Kolmogorov-Smirnov test data: \times D = 0.1316, p-value = 0.676 alternative hypothesis: two-sided

Our decision: p - value is very high. So, we accept the null hypothesis.

Case II: One-Sample, Composite Null

Checking Gaussianity:

$$H_0: F(x) = N(\mu, \sigma^2)$$
 versus $H_1: F(x) \neq N(\mu, \sigma^2)$

Note: (μ, σ^2) are NOT known.

Can we use ks.test(x,"pnorm",mean(x),sd(x))?

Kolmogorov-Smirnov Test for composite null

Can we use ks.test(x,"pnorm",mean(x),sd(x))?

NO.

Let's do it anyway and see what happens.

D = 0.1538, p-value = 0.4767 alternative hypothesis: two-sided

Result: if we do the test this way, p-value is high and we should accept the "Gaussianity". This result is misleading! Why?

Kolmogorov-Smirnov Test for composite null

Correct form of KS test for composite null, aka, Lilliefros test:

lillie.test(x)

Result:

```
Lilliefors (Kolmogorov-Smirnov) normality test data: x
D = 0.1538, p-value = 0.06793
```

Conclusion:

- Actual p-value is much lower than what we calculated before.
- Should we accept the null hypothesis?

Note

- Lilie test for nromality is in package "nortest"
- Lillie test for exponential is in package "exptest" (called ks.exp.test)

Discussion of KS test

the p-value was 0.06793 for the KS test. Should we accept the Gaussianity?

In such cases it is better to run other tests as well

Anderson-Darling test of normality

```
ad.test(x).
```

Result:

- Anderson-Darling normality test
- data: x
- \bullet A = 1.0224, p-value = 0.009251

We reject "Null".

ad test of normality is part of package "nortest".

ad test for exponential random variables is part of "exptest" package.

We would like to evaluate the impact of glucose treatment on the patients with Huntigotn's disease. The result for the patients that received treatment and did not receive the treatment are shown below.

We would like to see whether the treatment has had any positive or negative effect. Therefore, considering $X_i \sim F$ for the "with-treatment" samples and $Y_i \sim G$ for "no-treatment" samples we would like to test:

$$H_0: F(x) = G(x) \ \forall x \ \text{vs.} \ H_1: F(x) > G(x) \ \forall x \ \text{or} \ F(x) < G(x) \ \forall x.$$

Test this hypothesis by Wilcoxon test. Report the *p*-value.

```
x<-c(85, 89, 86, 91, 77, 93, 100, 82, 92, 86, 86)
y<-c(83, 73, 65, 65, 90, 77, 78, 97, 85, 75)
wilcox.test(x,y,alternative='two.sided',mu=0)
```

Wilcoxon rank sum test with continuity correction

```
data: x and y W = 4680, p-value < 2.2e-16 alternative hypothesis: true location shift is not equal to 0
```

Now we would like to implement permutation test with Wilcoxon statistic. Since implementing the full permutation test is time consuming we use Monte Carlo simulation with 500, 1000, 1500, 2000 Monte Carlo samples, and report the result of the permutation test with these four number of Monte Carlo samples. Then we will compare it with the result above.

```
z < -c(x,y)
zr < -rank(z)
p<-numeric(0)
number <-c (500, 1000, 1500, 2000)
m<-length(y)
n<-length(x)
stat < -abs(sum(zr[(length(x)+1):length(z)])-m*(m+1)/2-m*n/2);
for(j in 1:4){
  count<-0:
    for(i in 1:number[j]){
      test<-abs(sum(sample(zr,length(y)))-m*(m+1)/2-m*n/2);
      if(test>=stat) count<-count+1:
  p[i] <-count/number[i];</pre>
```