

Multiple Linear Regression: Categorical Predictors

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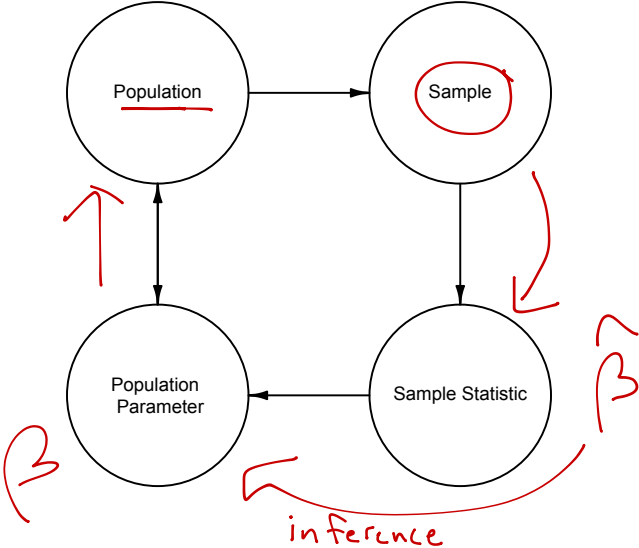
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Today's Lecture

- Sampling distribution of $\hat{\beta}$
- Confidence intervals
- Hypothesis tests for individual coefficients
- Global tests

Circle of Life



Statistical inference

- We have LSEs $\hat{\beta}_0, \hat{\beta}_1, \dots$; we want to know what this tells us about β_0, β_1, \dots
- Two basic tools are confidence intervals and hypothesis tests
 - ▶ Confidence intervals provide a plausible range of values for the parameter of interest based on the observed data
 - ▶ Hypothesis tests ask how probable are the data we gathered under a null hypothesis about the data generating distribution

$$Y_i \sim \beta_0 + \beta_1 X_i + \varepsilon_i$$

$\parallel \quad \parallel$
 $0 \quad 0$

$\underline{\sigma^2}$

Motivation

How can we draw **inference** about each of these parameters and relationships that our model is encoding?

```
mlr1 <- lm(disease ~ airqual + crowding + nutrition + smoking, d)
summary(mlr1)$coef
```

	Estimate	Std. Error	t value	Pr(> t)
## (Intercept)	11.86333	2.578819	4.600	1.316e-05
## airqual	0.25788	0.026799	9.623	1.165e-15
## crowding	1.11113	0.102037	10.889	2.404e-18
## nutrition	-0.03278	0.007954	-4.122	8.095e-05
## smoking	4.96093	1.085292	4.571	1.475e-05

t-stats

e-values

Motivation

- Can we say anything about whether the effect of airquality is “significant” after adjusting for other variables?
- Can we say whether adding airquality improves the fit of our model?
- Can we compare this model to a model with ~~only~~ crowding, nutrition and smoking?

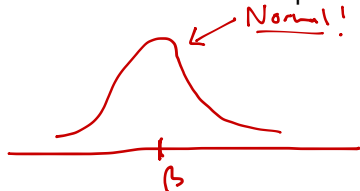
Sampling distribution

If our usual assumptions are satisfied and $\epsilon \stackrel{iid}{\sim} N[0, \sigma^2]$ then

$$\hat{\beta} \sim N \left[\beta, \sigma^2 (\mathbf{X}^T \mathbf{X})^{-1} \right]. \quad \text{- MV Normal}$$

$$\hat{\beta}_j \sim N \left[\beta_j, \sigma^2 (\mathbf{X}^T \mathbf{X})_{jj}^{-1} \right].$$

- This will be used later for inference. *jjth entry of Cov matrix*
- Even without Normal errors, asymptotic Normality of LSEs is possible under reasonable assumptions.



Sampling distribution

For real data we have to estimate σ^2 as well as β .

- Recall our estimate of the error variance is

$$\hat{\sigma}^2 = \frac{RSS}{n - p - 1} = \frac{\sum_i (y_i - \hat{y}_i)^2}{n - p - 1}$$

- With Normally distributed errors, it can be shown that

$$(n - p - 1) \frac{\hat{\sigma}^2}{\sigma^2} \sim \chi_{n-p-1}^2$$

Testing procedure

Calculate the probability of the observed data (or more extreme data) under a null hypothesis.

- Often $H_0 : \beta_1 = 0$ and $H_a : \beta_1 \neq 0$

- Set type I error rate

$\alpha = P(\text{falsely rejecting a true null hypothesis}) = .05$

- Calculate a test statistic assuming the null hypothesis is true

- Compute a p-value =

$$P(\text{As or more extreme test statistic} | H_0)$$

- Reject or fail to reject H_0

Individual coefficients

For individual coefficients

- We can use the test statistic

$$T = \frac{\hat{\beta}_j - \beta_j}{\widehat{\text{se}}(\hat{\beta}_j)} = \frac{\hat{\beta}_j - \beta_j}{\sqrt{\hat{\sigma}^2 (\mathbf{X}^T \mathbf{X})_{jj}^{-1}}} \sim t_{n-p-1}$$

- For a two-sided test of size α , we reject if

$$|T| > t_{1-\alpha/2, n-p-1}$$

- The p-value gives $P(t_{n-p-1} > T_{\text{obs}} | H_0)$

Note that t is a symmetric distribution that converges to a Normal as $n - p - 1$ increases.

Back to the example

$$\hat{\beta}_{y,x} = \frac{\hat{\beta}}{\text{sd}(x)}$$

```
summary(mlr1)
```

```
##  
## Call:  
## lm(formula = disease ~ airqual + crowding + nutrition + smoking,  
## data = dat)
```

```
## Residuals:
```

```
## Min 1Q Median 3Q Max  
## -8.130 -2.183 -0.572 1.941 13.326
```

```
## Coefficients:
```

	Estimate	Std. Error	t value	Pr(> t)
## (Intercept)	11.86333	2.57882	4.60	1.3e-05 ***
## airqual	0.25788	0.02680	9.62	1.2e-15 ***
## crowding	1.11113	0.10204	10.89	< 2e-16 ***
## nutrition	-0.03278	0.00795	-4.12	8.1e-05 ***
## smoking	4.96093	1.08529	4.57	1.5e-05 ***

$$1.2e^{-15} = 1.2 \times 10^{-15}$$

```
## ---  
## Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
## Residual standard error: 3.64 on 94 degrees of freedom
```

```
## Multiple R-squared: 0.866, Adjusted R-squared: 0.861
```

```
## F-statistic: 152 on 4 and 94 DF, p-value: <2e-16
```

Individual coefficients: CIs

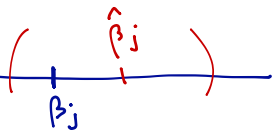
Alternatively, we can construct a confidence interval for β_j

- A confidence interval with coverage $(1 - \alpha)$ is given by

$$\hat{\beta}_j \pm \underbrace{t_{1-\alpha/2, n-p-1}}_{\sim 2} \widehat{se}(\hat{\beta}_j)$$

$1-\alpha\%$ ■ Assuming all the standard assumptions hold,
Our CI for $\hat{\beta}_j$ is (LB, UB) , if we repeated this experiment we would expect that 75% of our CIs would cover the truth.

$$(1 - \alpha) = P(LB < \beta_j < UB)$$



Back to the example

$$\frac{\alpha}{2}, 1 - \frac{\alpha}{2}$$

```
cbind(coef(mlr1), confint(mlr1))
```

##		2.5 %	97.5 %
##	(Intercept)	11.86333	6.74303 16.98364
##	airqual	0.25788	0.20467 0.31109
##	crowding	1.11113	0.90853 1.31372
##	nutrition	-0.03278	-0.04858 -0.01699
##	smoking	4.96093	2.80606 7.11580



Inference for linear combinations

Sometimes we are interested in making claims about $c^T \beta$ for some c .

- Define $H_0 : c^T \beta = c^T \beta_0$ or $H_0 : c^T \beta = 0$

- We can use the test statistic

$$T = \frac{c^T \hat{\beta} - c^T \beta}{\widehat{\text{se}}(c^T \hat{\beta})} = \frac{c^T \hat{\beta} - c^T \beta}{\sqrt{\hat{\sigma}^2 c^T (\mathbf{X}^T \mathbf{X})^{-1} c}}$$

c^T $\begin{bmatrix} \beta_0 \\ \beta_1 \\ \beta_2 \end{bmatrix}$ $H_0: \beta_0 + \beta_1 = 0$

- This test statistic is asymptotically Normally distributed
- For a two-sided test of size α , we reject if

$$|T| > z_{1-\alpha/2}$$

Inference about multiple coefficients

Our model contains multiple parameters; often we want to perform multiple tests:

$$H_{01} : \beta_1 = 0$$

$$H_{02} : \beta_2 = 0$$

$$\vdots = \vdots$$

$$H_{0k} : \beta_k = 0$$

where each test has a size of α

- For any individual test, $P(\text{reject } H_{0i} | H_{0i}) = \alpha$

Inference about multiple coefficients

What about

$$P(\text{reject at least one } H_{0i} | \text{all } H_{0i} \text{ are true}) = \alpha$$

$$Pr(\text{reject at 1 test} | H_{0i} \text{ true}) = \alpha$$

Family-wise error rate
FWER

Family-wise error rate

perform
k tests

To calculate the FWER

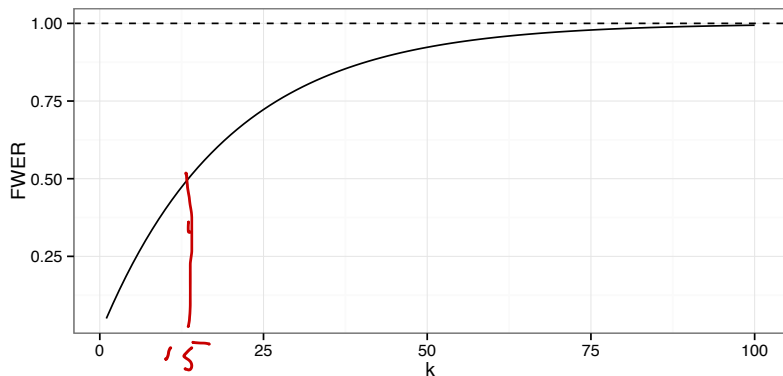
- First note $P(\text{no rejections} | \text{all } H_{0i} \text{ are true}) = \underline{(1 - \alpha)^k}$
- It follows that

$$\begin{aligned} \text{FWER} &= P(\text{at least one rejection} | \text{all } H_{0i} \text{ are true}) \\ &= 1 - (1 - \alpha)^k \end{aligned}$$

Family-wise error rate

$$\text{FWER} = 1 - (1 - \alpha)^k$$

```
alpha <- 0.05
k <- 1:100
FWER <- 1 - (1 - alpha)^k
qplot(k, FWER, geom = "line") + geom_hline(yintercept = 1, lty = 2)
```



Addressing multiple comparisons

Three general approaches

- Do nothing in a reasonable way
 - ▶ Don't trust scientifically implausible results ←
 - ▶ Don't over-emphasize ~~isolated~~ findings ←
- Correct for multiple comparisons
 - ▶ Often, use the Bonferroni correction and use $\alpha_i = \underline{\alpha/k}$ for each test
 - ▶ Thanks to the Bonferroni inequality, this gives an overall $FWER \leq \alpha$

- Use a global test

Very conservative

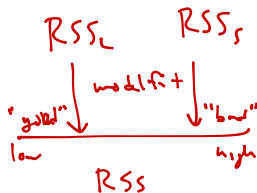
Global tests

Compare a smaller “null” model to a larger “alternative” model

- Smaller model must be nested in the larger model
- That is, the smaller model must be a special case of the larger model
- For both models, the RSS gives a general idea about how well the model is fitting
- In particular, something like

$$\frac{RSS_S - RSS_L}{RSS_L}$$

compares the relative RSS of the models



Nested models

- These models are nested:

Smaller = Regression of Y on X_1

Larger = Regression of Y on X_1, X_2, X_3, X_4

$$\beta_2 = \beta_3 = \beta_4 = 0$$

- These models are not:

Smaller = Regression of Y on X_2

Larger = Regression of Y on X_1, X_3

Global F tests

- Compute the test statistic

$$F_{obs} = \frac{(RSS_S - RSS_L)/(df_S - df_L)}{RSS_L/df_L}$$

- If H_0 (the null model) is true, then $F_{obs} \sim F_{df_S - df_L, df_L}$
- Note $df_S = n - p_S - 1$ and $df_L = n - p_L - 1$
- We reject the null hypothesis if the p-value is above α , where

$$\text{p-value} = P(F_{df_S - df_L, df_L} > F_{obs})$$

Global F tests

There are a couple of important special cases for the F test

- The null model contains the intercept only
 - ▶ When people say ANOVA, this is often what they mean (although all F tests are based on an analysis of variance)
- The null model and the alternative model differ only by one term
 - ▶ Gives a way of testing for a single coefficient
 - ▶ Turns out to be equivalent to a two-sided t -test: $t_{df_L}^2 \sim F_{1,df_L}$

Lung data: multiple coefficients simultaneously

You can test multiple coefficients simultaneously using the F test

```
mlr_null <- lm(disease ~ nutrition, data = dat)
mlr1 <- lm(disease ~ nutrition + airqual + crowding + smoking, data = dat)
anova(mlr_null, mlr1)
```

S L

```
## Analysis of Variance Table
##
## Model 1: disease ~ nutrition
## Model 2: disease ~ nutrition + airqual + crowding + smoking
##   Res.Df  RSS Df Sum of Sq  F Pr(>F)
## 1      97 9193
## 2      94 1248  3      7945 199 <2e-16 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```


Lung data: single coefficient test

The F test is equivalent to the t test when there's only one parameter of interest

```
mlr_null <- lm(disease ~ nutrition, data = dat)
mlr1 <- lm(disease ~ nutrition + airqual, data = dat)
anova(mlr_null, mlr1)

## Analysis of Variance Table
##
## Model 1: disease ~ nutrition
## Model 2: disease ~ nutrition + airqual
##   Res.Df  RSS Df Sum of Sq   F Pr(>F)
## 1      97 9193
## 2      96 5970  1      3223 51.8 1.3e-10 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

summary(mlr1)$coef

##              Estimate Std. Error t value Pr(>|t|)
## (Intercept)  37.6254    2.43946  15.42 9.946e-28
## nutrition    -0.0347    0.01692  -2.05 4.307e-02
## airqual       0.3611    0.05016   7.20 1.347e-10
```

Today's Big Ideas

- Inference for multiple linear regression models