Part II: Generalized Linear Models

Chapter II.3

Models for Binary Response

Topics

- To be discussed...
 - Binary Regression
 - Logistic Regression
 - Soodness of Fit for Logistic Regression
 - Infinite Estimates in Logistic Regression
 - Logistic Regression with Categorical Predictors
 - Linear Logit Model for Binary Response

Models for Binary Response

■ II.3.1 Remark (ungrouped vs. grouped data)

- Data for binary responses are usually *ungrouped*, in which case each observation y_i , $i=1,\ldots,n$, is the realization of a single Bernoulli trial, i.e. $y_i \in \{0,1\}$ and is a realization of $Y_i \sim \mathcal{B}(1,\pi_i)$. The total sample size is then $n_{tot} = \sum_{i=1}^n 1 = n$.
- For grouped data, each observation y_i^* , $i=1,\ldots,n$, corresponds to a set of sampled subjects/items which have exactly the same value for all explanatory variables. Hence, they are realizations of $Y_i^* = \sim \mathcal{B}(m_i,\pi_i)$ and thus $y_i^* \in \{0,\ldots,m_i\}$. In this case, these m_i subjects corresponding to the i-th group are realizations of independent Bernoulli trials with the same success probability p_i (naturally, since p_i is modeled by the explanatory variables) and $Y_i = Y_i^*/m_i$ (s. Example II.2.5). The total sample size in this case equals $n_{tot} = \sum_{i=1}^n m_i$. Usually grouped data occur when all explanatory variables are categorical.
- A grouped data set can be converted to an ungrouped one. Unless otherwise stated explicitly, we shall consider next ungrouped data.

II.3.2 Remark

The **logistic regression model** is the most common choice for modeling a categorical response variable (usually binary or ordinal) in terms of explanatory variables, which can be continuous or/and categorical.

Onsider a binary response $Y \sim \mathcal{B}(1,p)$ with $Y \in \{0,1\}$ and success probability p = P(Y=1).

odds of success

 \rightarrow Model the dependence of p on an explanatory variable X.

$$P(Y = 1) \stackrel{?}{=} \beta_0 + \beta_1 X \qquad \qquad f(z) = \frac{1}{1 + e^{-z}} = \frac{e^z}{1 + e^z} \iff z = \beta_0 + \beta_1 X$$

$$\downarrow z = \frac{e^z}{1 + e^{-z}} = \frac{e^z}{1 + e^z} \iff z = \beta_0 + \beta_1 X$$

$$P(Y = 1) = \frac{\exp(\beta_0 + \beta_1 X)}{1 + \exp(\beta_0 + \beta_1 X)} \implies \log[P(Y = 1)] = \log\left(\frac{P(Y = 1)}{1 - P(Y = 1)}\right) = \beta_0 + \beta_1 X$$

Binary Regression

■ II.3.3 Logistic Regression as a GLM

Binary response: Y, $Y \in \{0,1\}$ — Explanatory variables: X_2, \ldots, X_p ($X_1 = 1$ Leg coefficient of the intercept)

- **1** Random component: $Y_i \sim \mathcal{B}(1, \pi_i)$ [Fig belongs to the exponential dispersion family) $\mu_i = E(Y_i) = \pi_i$ for subject $i, i = 1, \dots, n$.
- ② Systematic component (linear predictor): $\sum_{j=1}^{p} \beta_j x_{ij}$ Parameter vector: $\boldsymbol{\beta} = (\beta_1, \dots, \beta_p)' \in \mathbb{R}^p$
- 3 Link function g relating linear predictor to μ_i : $g(\mu_i) = \beta_1 + \sum_{j=2}^p \beta_j x_{ij}$ canonical link: $g(\pi_i) = \log\left(\frac{\pi_i}{1-\pi_i}\right) = \theta_i$ (the logit link is the natural parameter in exponential family representation, s. Example II.2.15)

II.3.4 Remark

Often, the explanatory variables are considered to be X_1,\ldots,X_{p-1} and the index 0 is used for the coefficient of the intercept $(X_0=1)$. In this case $\boldsymbol{\beta}=(\beta_0,\beta_1,\ldots\beta_{p-1})'\in\mathbb{R}^p$ and the linear predictor is $\sum_{j=0}^{p-1}\beta_jx_{ij}$.

76

II.3.5 Remark

Other popular link functions for the set-up in II.3.3 and the random responses Y_i , $i=1,\ldots,n$, follow (s. also Example II.2.10).

- Probit (underlying normal latent variable): $\Phi^{-1}(\pi_i) = \beta_1 + \sum_{j=2}^p \beta_j x_{ij}$
- **S** Complementary log-log: $\log[-\log(1-\pi_i)] = \beta_1 + \sum_{j=2}^p \beta_j x_{ij}$

Tolerance Distribution Justification for Binary Regression

Framework: Toxicological experiments (explanatory variable X is the dosage of a drug)

Bliss: "The sigmoid dosage-mortality curve, ..., is interpreted as a cumulative normal frequency distribution of the variation among the individuals of a population in their susceptibility to a toxic agent, which susceptibility is inversely proportional to the logarithm of the dose applied." *

II.3.6 Remark

Suppose that the binary response Y is determined by a continuous underlying explanatory variable T (tolerance) so that for $i=1,\ldots,n$:

$$Y_i = 1$$
 if $x > T_i$ and $Y_i = 0$, else.

where T_i is the tolerance of the *i*-th person to a drug dosage equal to x. Then, for fixed X = x,

$$P(Y = 1|x) = \pi(x) = P(T \le x) = G(x) = F(\beta_1 + \beta_2 x)$$
,

for some 'standard' cdf and suggests models of the form

$$F^{-1}[\pi(x)] = \beta_0 + \beta_1 x$$
,

for some $\operatorname{cdf} F$ (i.e. motivates to consider a GLM with link function g the inverse of this cdf).

^{*}C.I.Bliss (1935). The calculation of the dosage-mortality curve, Annals of Applied Biology, 22:134-167.

► II.3.7 Special Binary Regression Models

• $F(x) = \frac{e^x}{1+e^x}$ | standard logistic regression model:

$$P(Y = 1|x) = \pi(x) = P(T \le x) = F(\beta_1 + \beta_2 x) = G(x) = \frac{e^{\beta_1 + \beta_2 x}}{1 + e^{\beta_1 + \beta_2 x}}$$

2
$$F(x) = \Phi(x)$$
 probit model: $\pi(x) = \Phi(\beta_1 + \beta_2 x)$

Simple Logistic Regression: Interpretation

Linearized form of the **simple logistic regression** model (logit transform):

$$\log\left(\frac{\pi(x)}{1-\pi(x)}\right) = \beta_1 + \beta_2 x$$

II.3.8 Remark (the role of β_2)

- \bullet odds $=\frac{\pi(x)}{1-\pi(x)}=e^{\beta_1+\beta_2x}=e^{\beta_1}(e^{\beta_2})^x$; i.e. multiplicative effect
- Monotonicity

$$\beta_2 > 0$$
 : $\pi(x) \uparrow$ as $x \to \infty$

$$\beta_2 < 0$$
 : $\pi(x) \downarrow \text{ as } x \to \infty$

$$\beta_2 = 0$$
 : $\pi(x)$ constant

Thus locally, $\pi(x)$ is approximated by a line with corresponding slope 'close' to x_0 :

$$\pi(x) \approx \pi(x_0) + \beta_2 \pi(x_0) (1 - \pi(x_0)) (x - x_0)$$
.

Slope is proportional to β_2 and steepest $(\beta_2/4)$ at x-value where $\pi(x) = 0.50$; this x value is $x = -\beta_1/\beta_2$, known as *median effective level*.

II.3.9 Remark (interpretation of β_2)

For two levels of x, denoted by x_1 and x_2 ,

odds ratio =
$$\frac{\pi(x_1)/[1-\pi(x_1)]}{\pi(x_2)/[1-\pi(x_2)]} = \frac{e^{\beta_1+\beta_2x_1}}{e^{\beta_1+\beta_2x_2}} = e^{\beta_2(x_1-x_2)}$$

- For $x_1 x_2 = 1$, the odds of a success at $x = x_1$ are e^{β_2} times the odds of success at $x = x_2$, i.e., odds multiply by e^{β_2} for every 1-unit increase in x. $\beta_2=0\longleftrightarrow$ odds ratio $=1\longleftrightarrow$ no effect of x on Y

II.3.10 Remark (generalization to multiple logistic regression model)

$$\log\left(\frac{\pi(x)}{1-\pi(x)}\right) = \beta_0 + \beta_1 x_1 + \ldots + \beta_q x_q$$

In this case, $\exp(\beta_j)$ represents the odds ratio between Y and two levels of x_j that are 1-unit apart $(j=2,\ldots,p)$, adjusting for all other predictors in the model.

Logistic Regression: Estimation of β

II.3.11 Remark

The logit link is the canonical link for a binomial response and from II.2.19 and II.2.24 we have the following results that apply to grouped or ungrouped data.

Likelihood Equations for β (s. Remark II.2.21):

$$\sum_{i=1}^{n} m_i (y_i - \pi_i) x_{ij} = 0 , \quad j = 1, \dots, p ,$$

i.e. the sufficient statistics of β_i 's are equated to their expected values.

Covariance matrix of $\hat{\beta}$:

For the information matrix for GLMs with canonical link holds
$$\mathcal{I}_F = \mathcal{I}_F^{obs} = \mathbf{X}'\mathbf{W}_c\mathbf{X}$$
 with $\mathbf{W}_c = \operatorname{diag}(\mathbf{w}_1, \dots, \mathbf{w}_n)$, where $\mathbf{w}_i = \frac{b''(\vartheta_i)}{a(b\cdot i)}$. In this case (s. Example II.2.5):

 $\mathbf{w}_i = \frac{\mathsf{Var}(Y_i)}{[a(\phi:i)]^2} = \frac{\pi_i(1-\pi_i)/m_i}{1/m_i^2} = m_i\pi_i(1-\pi_i)$ and thus the estimated covariance matrix of $\hat{\boldsymbol{\beta}}$ is

$$\mathsf{Cov}(\hat{oldsymbol{eta}}) = \mathcal{I}_F^{-1} = \left(\mathbf{X}' \mathsf{diag} \left[m_i \hat{\pi}_i (1 - \hat{\pi}_i)
ight] \mathbf{X}
ight)^{-1} \,.$$

Standard iterative methods apply for solving the system of likelihood equations for the logistic regression model. Furthermore, the Newton-Raphson and the Fischer scoring algorithms coincide (s. Remark II.2.29).

► II.3.12 Example (cancer remission)

Y: cancer remission (1 = ves, 0 = no)

Data: 27 Bernoulli outcomes or 14 binomials

Predictors for cancer remission

x: labeling index (LI)

```
(for example: 2 observations with LI=8 and Y=0); same likelihood function and ML estimates either way.   

LI <- c(8,8,10,10,12,12,12,14,14,14,16,16,16,18,20,20,20, 22,22,24,26, 28,32,34,38,38,38)   

y <- c(0,0,0,0,0,0,0,0,0,0,0,0,1,1,1,0,1,0,0,1,1,0,1,1,1,0)   

logit.fit <- glm(y \sim LI, family=binomial(link=logit))
```

→ What is saved under 'logit.fit'? I names(logit.fit)

► II.3.13 Example (cancer remission) - continues

summary(logit.fit)

```
glm(formula = v \sim LI, family = binomial(link = logit))
Call:
Deviance Residuals:
                          Median
Min
                                       30 Max
-1.9448 -0.6465
                         -0.4947 0.6571 1.6971
Coefficients:
             Estimate Std. Error z value Pr(>|z|)
(Intercept) -3.77714 1.37862 -2.740 0.00615 **
       0.14486 0.05934 2.441 0.01464 *
LI
___
Signif. codes: 0 ``***'' 0.001 ``**'' 0.01 ``*''0.05 ``.'' 0.1 `` '' 1
(Dispersion parameter for binomial family taken to be 1)
Null deviance: 34.372 on 26 degrees of freedom
Residual deviance: 26.073 on 25 degrees of freedom
AIC: 30.073
Number of Fisher Scoring iterations: 4
```

► II.3.14 Example (cancer remission) - continues

ML fit of logistic regression model for $\pi = P(\text{remission})$ is

$$\log\left(\frac{\hat{\pi}}{1-\hat{\pi}}\right) = \hat{\beta}_1 + \hat{\beta}_2 x = -3.777 + 0.145x$$

Prediction equation:

$$\hat{\pi}(x) = \frac{\exp(-3.777 + 0.145x)}{1 + \exp(-3.777 + 0.145x)}$$

Thus, at $\overline{x} = 20.1$,

$$\hat{\pi}(\overline{x}) = \frac{\exp(-3.777 + 0.145 \cdot 20.1)}{1 + \exp(-3.777 + 0.145 \cdot 20.1)} = 0.296$$

The incremental rate of change at x = 20.1 is

$$\hat{\beta}_2 \hat{\pi}(\overline{x})[1 - \hat{\pi}(\overline{x})] = 0.14486(0.296)(0.704) = 0.030$$

Furthermore,
$$\hat{\pi} = 0.50 \Leftrightarrow \log\left(\frac{\hat{\pi}}{1-\hat{\pi}}\right) = 0 = \hat{\beta}_1 + \hat{\beta}_2 x \Leftrightarrow x = -\frac{\hat{\beta}_1}{\hat{\beta}_2} = 26.0$$

► II.3.15 Example (cancer remission) - continues

Interpretation of $\hat{\beta}_2 = 0.145$: for each unit change in LI, the estimated odds of remission are multiplied by $\exp(0.145) = 1.16$, i.e., 16% increase when LI \uparrow 1.

```
E.g., at x=26, \hat{\pi}=0.498 (odds = 0.990) at x=27, \hat{\pi}=0.534=0.990\cdot 1.16 (odds = 1.145) i.e., odds ratio = \frac{0.534/(1-0.534)}{0.498/(1-0.498)}=1.16
```

Simpler effect measures: Change in $\hat{\pi}$ from minimum to maximum value of x

(Here, as LI goes from 8 to 38, $\hat{\pi}$ increases from 0.07 to 0.85)

Remark: With multiple predictors, can proceed to similar interpretations for a specific predictor setting all the other predictors equal to their means, modes (or at specific values of interest/levels).

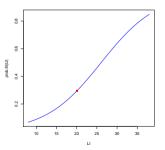
Estimated expected probabilities:

```
prob.fit <- function(x){
exp(logit.fit$coefficients[1]+ logit.fit$coefficients[2]*x)/
(1+exp(logit.fit$coefficients[1]+ logit.fit$coefficients[2]*x)) }
prob.fit(LI)  # the same as: logit.fit$fitted.values
prob.fit(mean(LI))</pre>
```

► II.3.16 Example (cancer remission) - continues

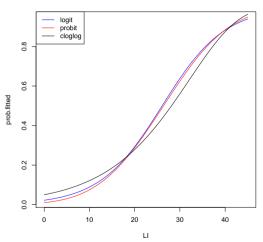
Plot of the estimated expected probabilities:

```
low <- min(LI); up <- max(LI)
plot(prob.fit, col='blue', from=low, to=up, xlab='LI', ylab='prob.fit(LI)')
x0 <- mean(LI); y0 <- prob.fit(mean(LI))
points(x0, y0, pch=15, col='red')</pre>
```



► II.3.17 Example (cancer remission) - continues





► II.3.18 Example (cancer remission) - continues

Tests of Significance and CIs

- Test of no effect $(H_0: \beta_2 = 0)$ SE = Standard Error
- $\mathbf{v} = \hat{\beta}_2/SE = \frac{0.145}{0.059} = 2.45$ ($z^2 = 5.96 \sim \mathcal{X}_1^2$, under H_0 , called **Wald** statistic) Strong evidence of a positive association between cancer remission and labeling index (p-value = 0.015).
- Confidence Interval (CI) for β_2 significant if the interval does not contain 0
- 95% Wald CI: $\hat{\beta}_2 \pm 1.96(SE) = (0.029, 0.261)$ (based on inverting the test above, e.g., the 95% CI is the set of β_2 not rejected at the 5% level in testing H_0 against $H_1: \beta_2 \neq 0$)

II.3.19 Remark

- ullet Beyond the Wald test statistic, there exist also the likelihood-ratio and the score test statistics. The three types of tests are asymptotically equivalent, when H_0 is true.
- 2 There exist other types of Cls, based on inverting the likelihood ratio and the score tests.
- 3 Methods extend to inference for multiple parameters.