Generalized Linear Models and their Applications STAT 431/STAT 831* Fall 2021 (1219)†

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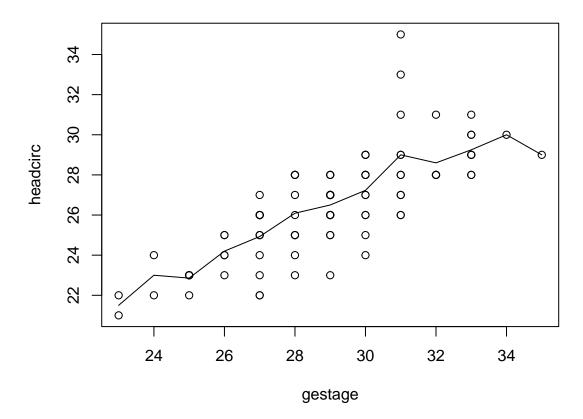
Topic 1a: Review of Linear Regression

Example: Low birthweight infants study¹

A study was conducted at two teaching hospitals in Boston, Massachusetts, where the head circumference, gestational age and some other variables are recorded for 100 low birth weight infants.

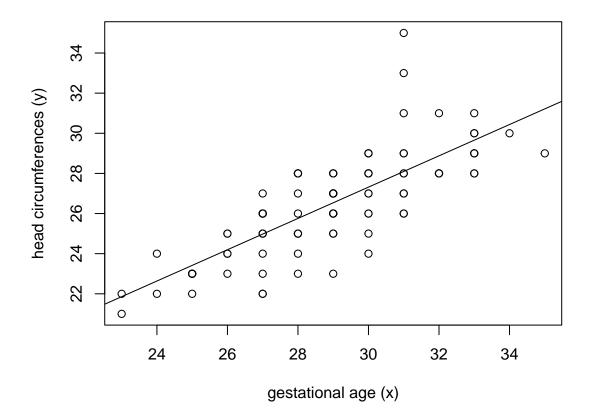
Question: what is the relationship between gestational age & head circumference?

A Scatterplot of the Data



We wish to model the relationship between gestational age and head circumference using a straight line!

¹Principles of Biostatistics 2nd Edition by Marcello Pagano, Kimberlee Gauvreau.



THE MODEL FITTING PROCESS

- (1) Model Specification: select a probability distribution for the response variable and a linear equation linking the response to the explanatory variables.
- 2 Estimation: finding the equation (the parameters of the model).
- 3 Model checking: how well does the model fit the data?
- (4) Inference: interpret the fitted model, calculate confidence intervals, conduct hypothesis tests.

1 Model Specification

Notation

For each subject i = 1, ..., n we have:

- Y_i = random variable representing the response, and
- $\boldsymbol{x}_i = (1, x_{i1}, \dots, x_{ip})^{\top}$ a vector of explanatory variables.

Specification for Multiple Linear Regression

• Linear regression equation:

$$Y_i = \beta_0 + \beta_1 x_{i1} + \dots + \beta_p x_{ip} + \varepsilon_i$$
 where $\varepsilon_i \stackrel{\text{iid}}{\sim} \mathcal{N}(0, \sigma^2)$.

• Equivalently, Y_i 's are independent $\mathcal{N}(\mu_i, \sigma^2)$ random variables or

$$\mu_i = E[Y_i] = \beta_0 + \beta_1 x_{i1} + \dots + \beta_p x_{ip}.$$

• For convenience, we often write linear regression models in matrix form as

$$Y = X\beta + \varepsilon$$
,

where

$$\boldsymbol{Y} = \begin{bmatrix} Y_1 \\ Y_2 \\ \vdots \\ Y_n \end{bmatrix}, \quad \boldsymbol{X} = \begin{bmatrix} 1 & x_{11} & \cdots & x_{1p} \\ 1 & x_{21} & \cdots & x_{2p} \\ \vdots & \vdots & \ddots & \vdots \\ 1 & x_{n1} & \cdots & x_{np} \end{bmatrix}, \quad \boldsymbol{\beta} = \begin{bmatrix} \beta_0 \\ \beta_1 \\ \vdots \\ \beta_p \end{bmatrix}, \quad \boldsymbol{\varepsilon} = \begin{bmatrix} \varepsilon_1 \\ \varepsilon_2 \\ \vdots \\ \varepsilon_n \end{bmatrix}$$

and

$$\boldsymbol{\varepsilon} \sim \text{MVN}(\boldsymbol{0}, \sigma^2 \boldsymbol{I}).$$

2 ESTIMATION

Least Squares Method

We wish to minimize a loss function:

$$S(\boldsymbol{\beta}) = \sum_{i=1}^{n} (y_i - \hat{y}_i)^2$$

$$= \sum_{i=1}^{n} (y_i - (\beta_0 + \beta_1 x_{i1} + \dots + \beta_p x_{ip}))^2$$

$$= (\boldsymbol{Y} - \boldsymbol{X}\boldsymbol{\beta})^{\top} (\boldsymbol{Y} - \boldsymbol{X}\boldsymbol{\beta}).$$

The least squares estimators (LSE) are the solutions to the equations:

$$\frac{\partial S}{\partial \boldsymbol{\beta}} = \frac{\partial}{\partial \boldsymbol{\beta}} (\boldsymbol{Y} - \boldsymbol{X} \boldsymbol{\beta})^{\top} (\boldsymbol{Y} - \boldsymbol{X} \boldsymbol{\beta}) = 0.$$

Maximum Likelihood Method

The probability density function for Y_i is:

$$f(y_i) = \frac{1}{\sqrt{2\pi\sigma^2}} \exp\left\{-\frac{1}{2\sigma^2} (y_i - (\beta_0 + \beta_1 x_{i1} + \dots + \beta_p x_{ip}))^2\right\}.$$

The log-likelihood function is therefore:

$$\ell(\boldsymbol{\beta}, \sigma^2) = \log \left(\prod_{i=1}^n f(y_i) \right)$$

$$= \sum_{i=1}^n \left(-\frac{1}{2} \log(2\pi\sigma^2) - \frac{1}{2\sigma^2} \left(y_i - (\beta_0 + \beta_1 x_{i1} + \dots + \beta_p x_{ip}) \right)^2 \right)$$

$$= -\frac{n}{2} \log(2\sigma^2) - \frac{1}{2\sigma^2} (\boldsymbol{Y} - \boldsymbol{X} \boldsymbol{\beta})^\top (\boldsymbol{Y} - \boldsymbol{X} \boldsymbol{\beta}).$$

The maximum likelihood estimators (MLE) of β are obtained by solving:

$$\frac{\partial \ell}{\partial \boldsymbol{\beta}} = \frac{\partial}{\partial \boldsymbol{\beta}} \left[-\frac{1}{2\sigma^2} (\boldsymbol{Y} - \boldsymbol{X}\boldsymbol{\beta})^\top (\boldsymbol{Y} - \boldsymbol{X}\boldsymbol{\beta}) \right] = 0.$$

• Parameter Estimates: For linear regression LSE and MLE of β are the same

$$\hat{\boldsymbol{\beta}} = (\boldsymbol{X}^{\top} \boldsymbol{X})^{-1} \boldsymbol{X}^{\top} \boldsymbol{Y}.$$

- Fitted values: $\hat{Y} = X\hat{\beta}$.
- Residuals: $\hat{r}_i = (y_i \hat{y}_i)$.
- Variance estimates:
 - An unbiased estimate of σ^2 is:

$$\hat{\sigma}^2 = \frac{1}{n - (p+1)} \sum_{i=1}^{n} \hat{r}_i^2.$$

- An estimate of the variance of $\hat{\beta}$ is:

$$\widehat{\mathbf{V}}(\hat{\boldsymbol{\beta}}) = \hat{\sigma}^2 (\boldsymbol{X}^{\top} \boldsymbol{X})^{-1}.$$

Low Birthweight Infant Data Example

- For n=100 infants, we have observed $Y_i=$ head circumference and $x_i=$ gestational age for baby i, $i=1,\ldots,100.$
- Consider a simple linear regression model:

$$Y_i = \beta_0 + \beta_1 x_i + \varepsilon_i.$$

• We can fit the model and obtain LSE/MSE using the lm() function in R.

- What is the interpretation of regression parameters β_0 and β_1 ?
 - β_0 (intercept): expected headcirc for a baby of a gestational age zero (x=0).
 - β_1 (slope): expected change in headcirc associated with a one unit increase in gestational age.

3 Model Checking

Standardized Residuals:

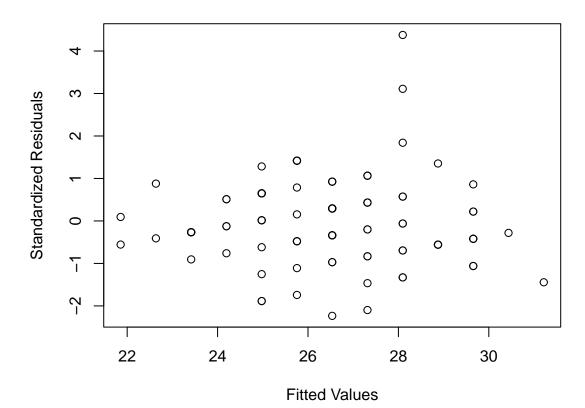
$$d_i = \frac{r_i}{\sqrt{\hat{\sigma}^2(1 - h_{ii})}},$$

where h_{ii} is the (i, i) element of $\mathbf{H} = (\mathbf{X}^{\top} \mathbf{X})^{-1} \mathbf{X}^{\top}$. By asymptotic theory, if the model provides a good fit to the data then we should expect that:

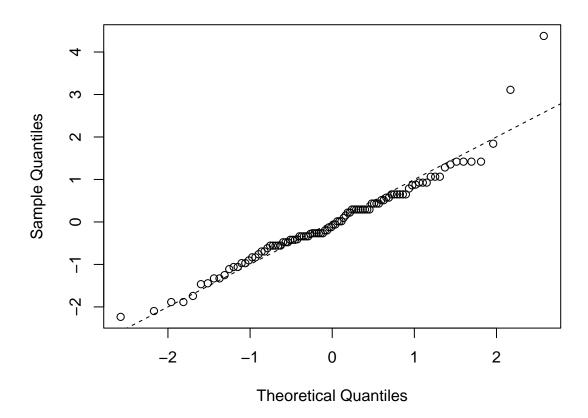
$$d_i \stackrel{\text{iid}}{\sim} \mathcal{N}(0,1).$$

We visually check this by examining residual plots such as:

- Standardized residuals versus the fitted values.
- Standardized residuals versus the explanatory variable(s).
- Normal probability plot (QQ plot) of the standardized residuals.



Normal Q-Q Plot



4 Inference

• Under suitable assumptions, the fitted regression parameters are asymptotically normally distributed:

$$\begin{split} \hat{\boldsymbol{\beta}} &\sim \text{MVN}(\boldsymbol{\beta}, \sigma^2(\boldsymbol{X}^{\top}\boldsymbol{X})^{-1}), \\ \hat{\beta}_j &\sim \mathcal{N}(\beta_j, \sigma^2 v_{jj}), \qquad \text{where } v_{jj} = \left[(\boldsymbol{X}^{\top}\boldsymbol{X})^{-1} \right]_{(j,j)}. \end{split}$$

- Since σ^2 is generally unknown, we replace it with the unbiased estimate $\hat{\sigma}^2$, and obtain $\operatorname{se}(\hat{\beta}_j) = \sqrt{\hat{\sigma}^2 v_{jj}}$.
- The inference is then based on the *t*-distribution result:

$$\frac{\hat{\beta}_j - \beta_j}{\operatorname{se}(\hat{\beta}_j)} \sim t_{n-p-1}.$$

Low Birthweight Infant Data Example

Is there a significant (linear) relationship between head circumference and gestational age?
 We wish to test H₀: β₁ = 0 vs H_A: β₁ ≠ 0.

$$t = \frac{\hat{\beta}_1 - (0)}{\operatorname{se}(\hat{\beta}_1)} \sim t_{98},$$

if H_0 is true, and we reject H_0 if $|t| > t_{98,0.975} = 1.985$. Here we have $t = 0.78/0.063 = 12.37 \gg 1.985$, so we reject H_0 .

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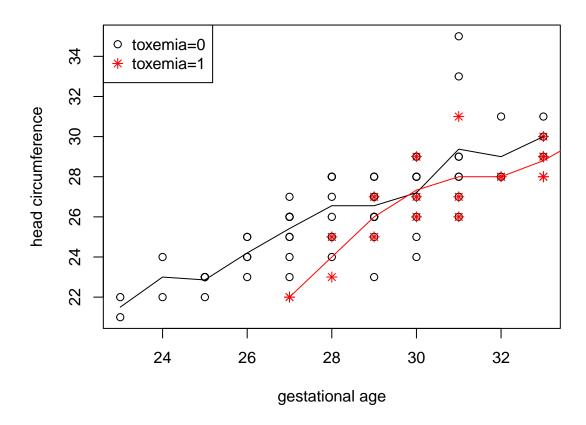
• What is the 95% confidence interval for the expected increase in head circumference when the gestational age of a baby increases by 1 week?

A 95 % CI for
$$\beta_1$$
:
$$\hat{\beta}_1 \pm t_{98,0.975} \operatorname{se}(\hat{\beta}_1) = 0.78 \pm 1.985 (0.063) = (0.665, 0.905).$$

LINEAR MODELS WITH MULTIPLE PREDICTORS

Low Birthweight Infant Data Example

• *Toxemia*, a pregnancy complication characterized by high blood pressure and signs of damage to liver and kidneys, may also have an impact on the development of babies.



• Does toxemia, after adjustment for gestational age, also affect the head circumference?

```
fit <- lm(headcirc ~ gestage + factor(toxemia), data = lowbwt)
summary(fit)

Call:
lm(formula = headcirc ~ gestage + factor(toxemia), data = lowbwt)

Residuals:
    Min    1Q    Median    3Q    Max</pre>
```

What is the interpretation of β_2 ?

 $\hat{\beta}_3 = -1.41233$. After adjustment of gestational age, the babies whose mothers had toxemia have smaller (by 1.41 cm) than those whose mothers did not. This difference is significant (test H_0 : $\beta_2 = 0$, p-value = 0.0076 < 0.05).

• Is the rate of increase of head circumference with gestational age the same for infants whose mothers with toxemia as those whose mother without it?

$$Y_i = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2} + \beta_3 x_{i1} x_{i2} + \varepsilon_i.$$

```
fit <- lm(headcirc ~ gestage * factor(toxemia), data = lowbwt)</pre>
summary(fit)
Call:
lm(formula = headcirc ~ gestage * factor(toxemia), data = lowbwt)
Residuals:
   Min
            1Q Median
                        3Q
-3.8366 -0.8366 -0.0928 0.7910 6.4341
Coefficients:
                      Estimate Std. Error t value Pr(>|t|)
(Intercept)
                       1.76291 2.10225 0.839 0.404
                        0.86461 0.07390 11.700 <2e-16 ***
gestage
                       -2.81503 4.98515 -0.565
                                                  0.574
factor(toxemia)1
gestage:factor(toxemia)1 0.04617 0.16352 0.282 0.778
Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
Residual standard error: 1.515 on 96 degrees of freedom
Multiple R-squared: 0.6531, Adjusted R-squared: 0.6422
F-statistic: 60.23 on 3 and 96 DF, p-value: < 2.2e-16
```

What is the interpretation of β_3 ?

 β_3 is the differences in slopes between the two groups (toxemia=1 vs toxemia=0). We want to test H_0 : $\beta_3 = 0$, t = 0.282, p-value = 0.778 > 0.05. No evidence to reject H_0 .

LIMITATIONS OF LINEAR REGRESSION

Linear regression models can be very useful but may not be appropriate to use when response Y is not continuous and can not be assumed to be normally distributed, e.g.,

- Binary data (Y = 0 or Y = 1),
- Count data (Y = 0, 1, 2, 3, ...).

Generalized Linear Models (GLM) extend the linear regression framework to address the above issue.

- Suitable for continuous and discrete data.
- Normal/Gaussian linear regression is a special case of GLM.
- Inference based on maximum likelihood methods (review next class 431 Appendix, Stat 330 notes).

WEEK 2 13th to 17th September

Topic 1b: Review of Likelihood Methods

DISTRIBUTIONS WITH A SINGLE PARAMETER

Setup

- Suppose Y is a random variable with probability density (or mass) function $f(y;\theta)$, where $\theta \in \Omega$ is a continuous parameter.
- The true value of θ is unknown.
- We wish to make inferences about θ (i.e., we may want to estimate θ , calculate a 95 % CI or carry out tests of hypotheses regarding θ).

LIKELIHOOD FUNCTION

• The Likelihood function is any function which is proportional to the probability of observing the data one actually obtained, i.e.,

$$L(\theta; y) = cf(y; \theta) = c P(Y = y; \theta),$$

where c is a proportionality constant that does not depend on θ .

- $L(\theta; y)$ contains all the information regarding θ from the data.
- $L(\theta;y)$ ranks the various parameter values in terms of their consistency with the data.
- Since $L(\theta; y)$ is defined in terms of the random variable y, it is itself a random variable.

MAXIMUM LIKELIHOOD ESTIMATOR

- For the purposes of estimation we typically want to find θ value that makes the observed data the most likely (hence the term maximum likelihood).
- The maximum likelihood estimator (MLE) of θ is

$$\hat{\theta} = \arg\max_{\theta} L(\theta; y).$$

• Estimation becomes a simple optimization problem!

• It is often easier to work with the logarithm of the likelihood function, i.e., the log-likelihood function

$$\ell(\theta; y) = \log(L(\theta; y)).$$

- Equivalently, since the $\log(\cdot)$ function is monotonic, the value of θ that maximizes $L(\theta; y)$ also maximizes the log-likelihood $\ell(\theta; y)$.
- For simplicity, we drop the y and use $L(\theta) = L(\theta; y)$ and $\ell(\theta) = \ell(\theta; y)$.

A LIST OF IMPORTANT FUNCTIONS

- Log-likelihood function: $\ell(\theta) = \log(L(\theta))$.
- Score function: $S(\theta) = \frac{\partial \ell(\theta)}{\partial \theta} = \ell'(\theta)$.
- Information function: $I(\theta) = -\frac{\partial^2 \ell(\theta)}{\partial \theta^2} = -\ell''(\theta)$.
- Fisher information function: $\mathcal{I}(\theta) = \mathbb{E}[I(\theta)]$.
- Relative likelihood function: $R(\theta) = L(\theta)/L(\hat{\theta})$.
- Log relative likelihood function: $r(\theta) = \log(L(\theta)/L(\hat{\theta})) = \ell(\theta) \ell(\hat{\theta})$.

MAXIMUM LIKELIHOOD ESTIMATION

- Want θ that maximizes $\ell(\theta)$, or equivalently solves $S(\theta) = 0$.
- Sometimes $S(\theta) = 0$ can be solved explicitly (easy in this case), but often we must solve iteratively.
- Check that the solution corresponds to a maxima of $\ell(\theta)$ by verifying the value of the second derivative at $\hat{\theta}$ is negative, or

$$I(\hat{\theta}) = -\ell''(\hat{\theta}) > 0.$$

• Invariance property of MLEs: if $g(\theta)$ is any function of the parameter θ , then the MLE of $g(\theta)$ is $g(\hat{\theta})$.

If $\hat{\theta}$ is the MLE of θ , then $e^{\hat{\theta}}$ is the MLE of e^{θ} .

EXAMPLE: BINOMIAL DISTRIBUTION

Example: Binomial Distribution

- A study was conducted to examine the risk for hormone use in healthy postmenopausal women.
- Suppose a group of *n* women received a combined hormone therapy, and were monitored for the development of breast cancer during 8.5 years follow-up.
- Let

$$Y_i = \begin{cases} 1 & \text{, if woman } i \text{ developed breast cancer,} \\ 0 & \text{, otherwise,} \end{cases}$$

for $i = 1, \ldots, n$.

• Suppose $Y_i \stackrel{\text{iid}}{\sim} \text{BERN}(\pi)$ where $\pi = P(Y_i = 1)$, then the total number of woman developed breast cancer is:

$$Y = \sum_{i=1}^{n} Y_i \sim BIN(n, \pi).$$

- We wish to find the MLE of unknown parameter π (probability of cancer).
- Likelihood function:

$$L(\pi; y) = c P(Y = y; \pi) = \pi^{y} (1 - \pi)^{n-y},$$

where we take $c = 1/\binom{n}{v}$ to simplify the likelihood.

• Log-likelihood function:

$$\ell(\pi) = y \log(\pi) + (n - y) \log(1 - \pi).$$

• Score function:

$$S(\pi) = \frac{y}{\pi} - \frac{n-y}{1-\pi}.$$

• Maximum Likelihood Estimator:

$$S(\pi) = 0 \implies \hat{\pi} = \frac{\sum_{i=1}^{n} y_i}{n} = \bar{y}.$$

• Second derivative test using information function:

$$I(\pi) = -\ell''(\pi) = \frac{y}{\pi^2} + \frac{n-y}{(1-\pi)^2} > 0 \ \forall \pi \in (0,1).$$

Example: Hormone Therapy Data

- A group of n=8506 postmenopausal women aged 50-79 received EPT and Y=166 developed invasive breast cancer during the follow-up.
- Assume $Y \sim \text{BIN}(n, \pi)$ with unknown parameter π .
- The maximum likelihood estimate of π is:

$$\hat{\pi} = \bar{y} = \frac{y}{n} = \frac{166}{8506} = 0.0195.$$

EXAMPLE: POISSON DISTRIBUTION

Suppose y_1, \ldots, y_n is an iid sample from a Poisson distribution with probability mass function:

$$f(y; \lambda) = P(Y = y; \lambda) = \frac{\lambda^y e^{-\lambda}}{y!}, \ \lambda > 0, \ y = 0, 1, 2, \dots$$

• Likelihood function:

$$L(\lambda; y_1, \dots, y_n) = \prod_{i=1}^n f(y_i; \lambda) = \frac{\lambda^{\sum y_i} e^{-n\lambda}}{\prod_i y_i!}.$$

• Log-likelihood function:

$$\ell(\lambda) = \left(\sum_{i} y_{i}\right) \log(\lambda) - n\lambda - \sum_{i=1}^{n} \log(y_{i}!).$$

• Score function:

$$S(\lambda) = \frac{\sum_{i} y_i}{\lambda} - n = 0 \implies \hat{\lambda} = \frac{\sum_{i=1}^{n} y_i}{n} = \bar{y}.$$

NEWTON RAPHSON ALGORITHM FOR FINDING MLE

- Sometimes, solving $S(\theta)=0$ can be challenging and closed form solutions may not be obtained, iterative method need to be used to find the MLE.
- Recall Taylor Series expansion of a differentiable function f(x) about a point a:

$$f(x) = f(a) + \frac{f'(a)}{1!}(x-a) + \frac{f''(a)}{2!}(x-a)^2 + \cdots$$

- Now suppose we wish to find $\hat{\theta}$, the root of $S(\theta) = 0$ and $\theta^{(0)}$ is a guess that is "close" to $\hat{\theta}$.
- Consider the Taylor series expansion of $S(\theta)$ about $\theta^{(0)}$:

$$S(\theta) = S(\theta^{(0)}) + \frac{S'(\theta^{(0)})}{1!} (\theta - \theta^{(0)}) + \frac{S''(\theta^{(0)})}{2!} (\theta - \theta^{(0)})^2 + \cdots$$

• For $|\theta - \theta^{(0)}|$ very small, the second and higher order terms can be dropped to a good approximation:

$$S(\theta) \simeq S(\theta^{(0)}) + S'(\theta^{(0)})(\theta - \theta^{(0)}).$$

 $S(\theta) \simeq S(\theta^{(0)}) - I(\theta^{(0)})(\theta - \theta^{(0)}).$

• Then at $\theta = \hat{\theta}$,

$$\begin{split} S(\hat{\theta}) &\simeq S(\theta^{(0)}) - I(\theta^{(0)})(\hat{\theta} - \theta^{(0)}) \\ I(\theta^{(0)})(\hat{\theta} - \theta^{(0)}) &\simeq S(\theta^{(0)}) \\ (\hat{\theta} - \theta^{(0)}) &\simeq I^{-1}(\theta^{(0)})S(\theta^{(0)}) \\ \hat{\theta} &\simeq \theta^{(0)} + I^{-1}(\theta^{(0)})S(\theta^{(0)}). \end{split}$$

• This suggests a revised guess for $\hat{\theta}$ is:

$$\theta^{(1)} = \theta^{(0)} + I^{-1}(\theta^{(0)})S(\theta^{(0)})$$

Newton Raphson Algorithm for finding the MLE

- Begin with an initial estimate $\theta^{(0)}$.
- Iteratively obtain updated estimate by using:

$$\theta^{(i+1)} = \theta^{(i)} + I^{-1}(\theta^{(i)})S(\theta^{(i)}).$$

- Iteration continues until $\theta^{(i+1)} \simeq \theta^{(i)}$ within a specified tolerance.
- Then set $\hat{\theta} = \theta^{(i+1)}$, check that $I(\hat{\theta}) > 0$.

Inference for Scalar Parameters θ

- So far we have discussed estimation of $\hat{\theta}$, next we want to conduct inference about θ , i.e., carry out hypothesis tests and construct confidence intervals of θ .
- Likelihood inference relies on the following asymptotic distribution results:

Useful asymptotic distributional results

- (log) Likelihood ratio statistic: $-2\log(R(\theta)) = -2r(\theta) \sim \chi_{(1)}^2$.
- Score statistic: $(S(\theta))^2/I(\theta) \sim \chi_{(1)}^2$.
- Wald statistic: $(\hat{\theta} \theta)^2 I(\hat{\theta}) \sim \chi^2_{(1)}$ or $(\hat{\theta} \theta) \sqrt{I(\hat{\theta})} \sim \mathcal{N}(0, 1)$ since $Z \sim \mathcal{N}(0, 1) \implies Z^2 \sim \chi^2_1$.

CONFIDENCE INTERVAL (CI)

Suppose we want a $100(1-\alpha)$ % confidence interval for θ .

• The Likelihood ratio (LR) based pivotal gives a confidence interval:

$$\{\theta: -2r(\theta) < \chi_1^2(1-\alpha)\},\$$

where $\chi_1^2(1-\alpha)$ is the upper α percentage point of the χ_1^2 distribution.

• The Wald-based pivotal gives an interval:

$$\Big\{\theta: (\hat{\theta}-\theta)^2 I(\hat{\theta}) < \chi_1^2(1-\alpha)\Big\},$$

or equivalently

$$\hat{\theta} \pm Z_{1-\alpha/2} \big(I(\hat{\theta}) \big)^{-1/2},$$

where $Z_{1-\alpha/2}$ is the upper $\alpha/2$ percentage point of the standard normal.

EXAMPLE: HORMONE THERAPY DATA

Likelihood Ratio based 95 % CI: $\{\theta: -2r(\theta) < \chi_1^2(0.95)\}$ where $r(\theta) = \ell(\theta) - \ell(\hat{\theta})$.

• For the Binomial distribution: $\hat{\theta} = y/n$, and

$$r(\theta) = \left(y\log(\theta) + (n-y)\log(1-\theta)\right) - \left(y\log\left(\frac{y}{n}\right) + (n-y)\log\left(1-\frac{y}{n}\right)\right).$$

• To find the root of $-2r(\theta) = \chi_1^2(0.95)$:

• The likelihood ratio based 95% CI is (0.017, 0.023).

Wald based 95 % CI: $\hat{\theta} \pm Z_{0.975} (I(\hat{\theta}))^{-1/2}$.

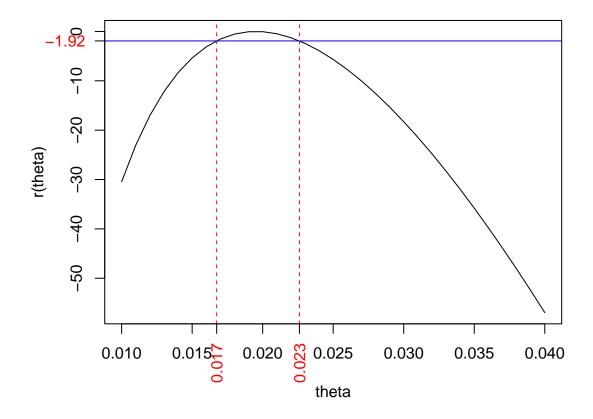
• For Binomial distribution $\hat{\theta} = y/n$ and

$$I(\hat{\theta}) = \frac{y}{\hat{\theta}^2} + \frac{n-y}{(1-\hat{\theta})^2} = n^2 \left(\frac{1}{y} + \frac{1}{n-y}\right).$$

• So we solve:

$$\hat{\theta} \pm 1.96 (I(\hat{\theta}))^{-1/2} = 0.0195 \pm 1.96 (0.0015)$$
$$= (0.017, 0.022).$$

• The Wald based 95% CI is: (0.017, 0.022).



HYPOTHESES TEST

Suppose we are interested in testing hypotheses:

$$H_0$$
: $\theta = \theta_0$ vs H_A : $\theta \neq \theta_0$.

• Likelihood ratio (LR) test: p-value = $P(\chi_1^2 > -2r(\theta_0))$.

- Score test: p-value = $P(\chi_1^2 > (S(\theta))^2 / I(\theta_0))$.
- Wald test:

$$p$$
-value = $P\left(\chi_1^2 > (\hat{\theta} - \theta_0)^2 I(\hat{\theta})\right)$, or p -value = $P\left(|Z| > |\hat{\theta} - \theta_0|\sqrt{I(\hat{\theta})}\right)$.

EXAMPLE: HORMONE THERAPY DATA

Suppose we wish to test if women received EPT would have a risk of breast cancer same as that of the general population, say about 1.5%.

$$H_0$$
: $\theta = 0.015$ vs H_A : $\theta \neq 0.015$.

• Likelihood Ratio based test:

$$r(\theta_0 = 0.015) = \left(y\log(0.015) + (n-y)\log(1-0.15)\right) - \left(y\log\left(\frac{y}{n}\right) + (n-y)\log\left(1-\frac{y}{n}\right)\right)$$

= -5.3637.

Thus, the p-value for the test is given by:

$$p = P(\chi_{(1)}^2 > -2r(0.015)) = P(\chi_{(1)}^2 > 10.7274) = 0.001.$$

Therefore, we *reject* H_0 and conclude that the risk of breast cancer for women received EPT is significantly different from 1.5%.

Notes on Asymptotic Inference

- Asymptotic results: approximation improves as sample size increases.
- Results are exact for a Normal linear model if θ is the mean parameter and σ^2 is known.
- LR approach:
 - Need to evaluate (log) likelihood at two locations.
 - Not always a closed from solution for a CI.
 - Usually the best approach.
- · Score approach:
 - Usually the least powerful test.
 - Don't actually need to find MLE to use.
- Wald's approach:
 - Always get a closed form solution for a CI.
 - May not behave well for skewed likelihoods (transform?).
- All three are asymptotically equivalent!

LIKELIHOOD METHODS FOR PARAMETER VECTORS

Suppose $\theta \in \Omega$ is a continuous $p \times 1$ parameter vector indexing a probability density (or mass) function $f(y; \theta)$. The likelihood and log-likelihood functions are defined as before, but

• $S(\theta) = \frac{\partial \ell(\theta)}{\partial \theta}$ is the $p \times 1$ Score vector, i.e.,

$$oldsymbol{S}(oldsymbol{ heta}) = egin{bmatrix} rac{\partial \ell(heta)}{\partial heta_1} \ dots \ rac{\partial \ell(heta)}{\partial heta_p} \end{bmatrix}.$$

• $I(\theta)=-rac{\partial^2\ell(\theta)}{\partial \theta^{\top}\,\partial \theta}$ is the p imes p Information matrix, i.e.,

$$m{I}(m{ heta}) = egin{bmatrix} -rac{\partial^2\ell(heta)}{\partial heta_1^2} & -rac{\partial^2\ell(heta)}{\partial heta_1\,\partial heta_2} & \cdots & rac{\partial^2\ell(heta)}{\partial heta_1\,\partial heta_p} \ -rac{\partial^2\ell(heta)}{\partial heta_2^2} & \cdots & rac{\partial^2\ell(heta)}{\partial heta_1\,\partial heta_p} \ & \ddots & rac{\partial^2\ell(heta)}{\partial heta_p^2} \end{bmatrix}.$$

• The Newton Raphson algorithm applies as before, but with vectors and matrices as follows:

$$\boldsymbol{\theta}^{(i+1)} = \boldsymbol{\theta}^{(i)} + \boldsymbol{I}^{-1}(\boldsymbol{\theta}^{(i)})\boldsymbol{S}(\boldsymbol{\theta}^{(i)}).$$

- Again, we apply iteratively until we obtain convergence, but now check to see if $I(\hat{\theta})$ is a positive definite matrix.
- Analogues to the LR, Score and Wald results apply based on partitioning the Information matrix by
 θ = (α, β)^T, where α is a p × 1 vector of nuisance parameters and β is a q × 1 vector of parameters of
 interest:

$$I = I(lpha,eta) = egin{pmatrix} I_{lphalpha}(lpha,eta) & I_{lphaeta}(lpha,eta) \ I_{etalpha}(lpha,eta) & I_{etaeta}(lpha,eta) \end{pmatrix},$$

where $I_{\alpha\alpha}(\alpha,\beta) = -\frac{\partial^2 \ell}{\partial \alpha \partial \alpha^{\top}}$ is $p \times p$, $I_{\alpha\beta}(\alpha,\beta) = -\frac{\partial^2 \ell}{\partial \alpha \partial \beta^{\top}}$ is $p \times q$, $I_{\beta\alpha}(\alpha,\beta) = -\frac{\partial^2 \ell}{\partial \beta \partial \alpha^{\top}}$ is $q \times p$, and $I_{\beta\beta}(\alpha,\beta) = -\frac{\partial^2 \ell}{\partial \beta \partial \beta^{\top}}$ is $q \times q$.

Topic 2a: Formulation of Generalized Linear Models

THE EXPONENTIAL FAMILY

Definition (Exponential Family)

Consider a random variable Y with probability density (or mass) function $f(y; \theta, \phi)$, we say that the distribution is a member of the exponential family if we can write

$$f(y; \theta, \phi) = \exp\left\{\frac{y\theta - b(\theta)}{a(\phi)} + c(y; \phi)\right\},$$

for some functions $a(\cdot)$, $b(\cdot)$, and $c(\cdot)$.

- The parameter θ is called the canonical parameter, and it is unknown.
- The parameter ϕ is called the scale/dispersion parameter, is constant, and assumed to be known.

Many well known distributions (continuous/discrete) can be shown to be a member of the exponential family.

EXAMPLES

• Poisson Distribution: $Y \sim POI(\lambda)$,

$$f(y; \lambda) = \frac{\lambda^y e^{-\lambda}}{y!}, \ \lambda > 0, \ y = 0, 1, \dots$$

Show that Poisson is a member of exponential family and identify the canonical parameter and the functions $a(\cdot)$, $b(\cdot)$, and $c(\cdot)$.

Solution.
$$f(y;\lambda)=\exp\Bigl\{\log(f(y;\lambda))\Bigr\}=\exp\Bigl\{\frac{y\log(\lambda)-\lambda}{1}-\log(y!)\Bigr\}$$
. Therefore,
$$\theta=\log(\lambda) \qquad \text{(canonical/natural parameter)},$$

$$b(\theta)=\lambda=e^{\theta},$$

$$\phi=1,$$

$$a(\phi)=1,$$

$$c(y;\phi)=-\log(y!).$$

• Normal Distribution: $Y \sim \mathcal{N}(\mu, \sigma^2)$ and σ^2 known,

$$f(y; \theta, \phi) = \frac{1}{\sqrt{2\pi\sigma^2}} \exp\left\{-\frac{(y-\mu)^2}{2\sigma^2}\right\}.$$

Show that this Normal distribution is a member of the exponential family. **Solution.**

$$f(y; \mu, \sigma^2) = \exp\left\{-\frac{y^2 - 2\mu y + \mu^2}{\sigma^2} - \frac{1}{2}\log(2\pi\sigma^2)\right\}$$
$$= \exp\left\{\frac{y\mu - \mu^2/2}{\sigma^2} - \frac{y^2}{2\sigma^2} - \frac{1}{2}\log(2\pi\sigma^2)\right\}.$$

Therefore,

$$\theta = \mu,$$

$$\phi = \sigma^2,$$

$$a(\phi) = \phi = \sigma^2,$$

$$b(\theta) = \frac{\mu^2}{2} = \frac{\theta^2}{2},$$

$$c(y; \phi) = -\frac{y^2}{2\sigma^2} - \frac{1}{2}\log(2\pi\sigma^2).$$

PROPERTIES OF EXPONENTIAL FAMILY

Consider a single observation y from the exponential family.

$$\begin{split} L(\theta,\phi;y) &= f(y;\theta,\phi) = \exp\biggl\{\frac{y\theta - b(\theta)}{a(\phi)} + c(y;\phi)\biggr\}. \\ \ell(\theta,\phi;y) &= \log\bigl(f(y;\theta,\phi)\bigr) = \frac{y\theta - b(\theta)}{a(\phi)} + c(y;\phi). \\ S(\theta) &= \frac{\partial \ell}{\partial \theta} = \frac{y - b'(\theta)}{a(\phi)}. \\ I(\theta) &= -\frac{\partial^2 \ell}{\partial \theta^2} = \frac{b''(\theta)}{a(\phi)}. \\ \mathcal{I}(\theta) &= \mathrm{E}\left[-\frac{\partial^2 \ell}{\partial \theta^2}\right] = I(\theta). \end{split}$$

SOME GENERAL RESULTS FOR SCORE AND INFORMATION

Result # 1

The expectation of the score function is zero.

$$\mathrm{E}\big[S(\theta)\big] = 0.$$

Proof:

$$\int f(y;\theta,\phi) \, dy = 1$$

$$\frac{\partial}{\partial \theta} \int f(y;\theta,\phi) \, dy = 0$$

$$\int \frac{\partial}{\partial \theta} f(y;\theta,\phi) \, dy = 0$$

$$\int \left(\frac{\partial}{\partial \theta} \log(f(y;\theta,\phi))\right) f(y;\theta,\phi) \, dy = 0$$

$$\int S(\theta) f(y;\theta,\phi) \, dy = 0$$

$$\text{E}[S(\theta)] = 0$$
(1)

Result # 2

The expectation of the score function squared is the expected information.

$$E[S(\theta; y)^2] = E[I(\theta; y)]$$

Proof: Differentiate (1) again,

$$\int \left(\frac{\partial}{\partial \theta} \log(f(y;\theta,\phi))\right) f(y;\theta,\phi) \, \mathrm{d}y = 0$$

$$\int \left(\frac{\partial^2}{\partial \theta^2} \log(f(y;\theta,\phi))\right) f(y;\theta,\phi) \, \mathrm{d}y + \int \left(\frac{\partial}{\partial \theta} \log(f(y;\theta,\phi))\right) \frac{\partial}{\partial \theta} f(y;\theta,\phi) \, \mathrm{d}y = 0$$

$$\int \frac{\partial^2}{\partial \theta^2} \log(f(y;\theta,\phi)) f(y;\theta,\phi) \, \mathrm{d}y + \int \left(\frac{\partial}{\partial \theta} f(y;\theta,\phi)\right)^2 f(y;\theta,\phi) \, \mathrm{d}y = 0$$

$$\int -I(\theta) f(y;\theta,\phi) \, \mathrm{d}y + \int S(\theta)^2 f(y;\theta,\phi) \, \mathrm{d}y = 0$$

$$\mathrm{E}[-I(\theta;y)] + \mathrm{E}[S(\theta;y)^2] = 0$$

Now for the exponential family, we apply above results and obtain:

$$\begin{split} & \mathbf{E}\big[S(\theta)\big] = 0, \\ & \mathbf{E}\bigg[\frac{Y - b'(\theta)}{a(\phi)}\bigg] = 0, \\ & \mathbf{E}[Y] = b'(\theta), \end{split}$$

$$\begin{split} \mathbf{E}\big[S(\theta)^2\big] &= \mathbf{E}\big[I(\theta)\big],\\ \mathbf{E}\bigg[\bigg(\frac{Y-b'(\theta)}{a(\phi)}\bigg)^2\bigg] &= \mathbf{E}\bigg[\frac{b''(\theta)}{a(\phi)}\bigg],\\ \frac{1}{a(\phi)^2}\,\mathbf{E}\Big[\big(Y-\mathbf{E}[Y]\big)^2\bigg] &= \frac{b''(\theta)}{a(\phi)},\\ \mathbf{Var}(Y) &= b''(\theta)a(\phi). \end{split}$$

Mean and Variance for the Exponential Family

• Mean: $E[Y] = b'(\theta) = \mu$.

• Variance: $Var(Y) = b''(\theta)a(\phi)$.

Note that:

• $b'(\theta) = \mu$ tells the relationship between *canonical* parameter θ and μ .

• $b''(\theta)$ is a function of θ and hence can be also expressed as a function of μ .

• Thus, we write $b''(\theta) = V(\mu)$ and call $V(\mu)$ the variance function.

• Subsequently, we have:

$$Var(Y) = b''(\theta)a(\phi) = V(\mu)a(\phi),$$

which is the mean-variance relationship for the exponential family.

LINK FUNCTIONS

Definition (Link Function)

The link function relates the linear predictor $\eta = x^{\top} \beta$ to the expected value μ of the random variable Y, i.e.,

$$g(\mu) = \eta = \boldsymbol{x}^{\top} \boldsymbol{\beta},$$

where $g(\cdot)$ is the link function.

Definition (Canonical Link Function)

When Y is a member of the exponential family we define the canonical link function to be:

$$g(\mu) = \theta = \eta = \boldsymbol{x}^{\top} \boldsymbol{\beta}$$

(i.e., the choice of $g(\cdot)$ that sets canonical parameter = linear predictor).

EXAMPLES

Recall that $POI(\lambda)$ is a member of exponential family,

$$f(y; \lambda) = \frac{\lambda^y e^{-\lambda}}{y!} = \exp\left\{\frac{y \log(\lambda) - \lambda}{1} - \log(y!)\right\}$$

where $\theta = \log(\lambda)$, $\phi = 1$, $b(\theta) = \lambda = e^{\theta}$, and $a(\phi) = 1$. Now to find the mean, variance function, and canonical link function:

- Mean: $E[Y] = b'(\theta) = e^{\theta} = \mu \implies \theta = \log(\mu)$.
- Variance Function: $V(\mu) = b''(\theta) = e^{\theta} \implies V(\mu) = \mu$.
- Variance: $Var(Y) = V(\mu)a(\phi) = \mu$ (mean-variance relationship).
- Canonical link: set $\theta = \eta$ using $\theta = \log(\mu) = \eta = \boldsymbol{x}^{\top}\boldsymbol{\beta}$, i.e., $g(\mu) = \log(\mu)$ where $\log(\cdot)$ is the canonical link.

Moving forward, we consider a log-linear model: $\log(\mu_i) = \boldsymbol{x}_i^{\top} \boldsymbol{\beta}$.

REMARKS ON LINK FUNCTION

- We can choose any function $q(\cdot)$ as the link function in theory.
- The canonical link is a special link function, we often choose to use canonical link for its good statistical properties.
- Context and goodness of fit should motivate the choice of link function in practice.

GENERALIZED LINEAR MODELS

Definition (Generalized Linear Model (GLM))

A Generalized Linear Model (GLM) is composed of three components:

- Random Component: The responses Y_1, \ldots, Y_n are independent random variables and each Y_i is assumed to come from a parametric distribution that is a member of the exponential family.
- Systematic Component (or linear predictor):

$$\eta_i = \boldsymbol{x}_i^{\top} \boldsymbol{\beta},$$

a linear combination of explanatory variables x_i and regression parameters β .

• Link function:

$$g(\mu_i) = \eta_i = \boldsymbol{x}_i^{\top} \boldsymbol{\beta},$$

a function that relates the mean of response to the linear predictor.

TOPIC SUMMARY

- 1. Definition of the Exponential Family.
 - Exponential form of the probability density (or mass) function.
 - Derivation of Score and Information.
 - Properties of exponential family, mean-variance relationship.
 - · Definition of canonical link.

2. Definition of a Generalized Linear Model.

Next Topic: 2b Estimation for Generalized Linear Models.

Topic 2b: Maximum Likelihood Estimation for Generalized Linear Models

GENERALIZED LINEAR MODELS

Suppose for each subject $i=1,\ldots,n$ in a random sample:

- Y_i is the response variable.
- x_{i1}, \ldots, x_{ip} are explanatory variables associated with Y_i .

We consider a Generalized Linear Model (GLM) for the data, by definition the GLM is composed following three components:

1 Random Component:

 $Y_i \sim \text{exponential family,} \qquad Y_1, \dots, Y_n \text{ are independent.}$

2 Systematic Component (or linear predictor):

$$\eta_i = \boldsymbol{x}_i^{\top} \boldsymbol{\beta} = \beta_0 + \beta_1 x_{i1} + \dots + \beta_p x_{ip}.$$

- $x_i = (1, x_{i1}, \dots, x_{ip})^{\top}$ is a covariate vector.
- $\boldsymbol{\beta} = (\beta_0, \beta_1, \dots, \beta_p)^{\top}$ is a vector of regression coefficients.
- 3 Link function: a function $g(\cdot)$ links $E[Y_i] = \mu_i$ to a linear prediction η_i :

$$g(\mu_i) = \eta_i = \boldsymbol{x}_i^{\top} \boldsymbol{\beta}.$$

EXAMPLE: A POISSON REGRESSION MODEL

Suppose $Y_i \overset{\text{ind}}{\sim} \text{POI}(\lambda_i)$ with mean $\mathrm{E}[Y_i] = \lambda_i$, $i = 1, \dots, n$:

$$f(y_i) = \frac{e^{-\lambda_i} \lambda_i^{y_i}}{y_i!} = \exp\{y_i \log(\lambda_i) - \lambda_i - \log(y_i!)\}.$$

Poisson distribution is a member of exponential family with:

- Canonical parameter: $\theta_i = \log(\lambda_i)$.
- Canonical link: $\theta_i = \eta_i \Longrightarrow \log(\lambda_i) = \boldsymbol{x}_i^{\top} \boldsymbol{\beta}$ (log link).

A Poisson regression model with the canonical link takes the form:

$$\log(\lambda_i) = \boldsymbol{x}_i^{\top} \boldsymbol{\beta} = \beta_0 + \beta_1 x_{i1} + \dots + \beta_n x_{in}$$
 (log-linear model).

EXAMPLE: A NORMAL REGRESSION MODEL

Assume $Y_i \stackrel{\text{ind}}{\sim} \mathcal{N}(\mu_i, \sigma^2)$ and σ^2 is known, $i = 1, \dots, n$:

$$f(y_i) = (2\pi\sigma^2)^{-1/2} \exp\left\{-\frac{(y_i - \mu_i)^2}{2\sigma^2}\right\}$$
$$= \exp\left\{\frac{y_i \mu_i - \mu_i^2/2}{\sigma^2} - \frac{1}{2}\left(\frac{y_i^2}{\sigma^2} + \log(2\pi\sigma^2)\right)\right\}.$$

A Normal distribution (σ^2 known) is a member of exponential family with:

- Canonical parameter: $\theta_i = \mu_i$.
- Canonical link: $\theta_i = \eta_i \Longrightarrow \mu_i = \boldsymbol{x}_i^{\top} \boldsymbol{\beta}$ (identity link).

A Normal regression model with the canonical link takes the form:

$$\mu_i = \boldsymbol{x}_i^{\top} \boldsymbol{\beta} = \beta_0 + \beta_1 x_{i1} + \dots + \beta_p x_{ip}$$
 (linear model).

Linear regression model (STAT 331) is a Normal GLM using the canonical link!

LIKELIHOOD FOR GENERALIZED LINEAR MODELS

We wish to use likelihood methods for the estimation of the regression parameter β from the GLM: $g(\mu_i) = \boldsymbol{x}_i^{\top} \boldsymbol{\beta}$. Consider the log-likelihood for a *single* observation from the exponential family:

$$\ell(\theta, \phi; y) = \frac{y\theta - b(\theta)}{a(\phi)} + c(y; \phi).$$

- ℓ is a function of θ (assume that ϕ is known).
- θ is related to μ through the result:

$$\mu = b'(\theta)$$
.

• η can be expressed in terms of μ through the link function:

$$q(\mu) = \eta$$
.

• β can be expressed in terms of η through the linear predictor:

$$\eta = \boldsymbol{x}^{\top} \boldsymbol{\beta}.$$

SCORE VECTOR

To find the maximum likelihood estimator for β , we must solve $S(\beta) = \frac{\partial \ell}{\partial \beta} = 0$. Consider taking derivative with respect to β_i using the chain rule:

$$\frac{\partial \ell}{\partial \beta_j} = \frac{\partial \ell}{\partial \theta} \frac{\partial \theta}{\partial \mu} \frac{\partial \mu}{\partial \eta} \frac{\partial \eta}{\partial \beta_j},$$

where

$$\begin{split} \frac{\partial \ell}{\partial \theta} &= \frac{y - b'(\theta)}{a(\phi)}, \\ \frac{\partial \theta}{\partial \mu} &= \left(\frac{\partial \mu}{\partial \theta}\right)^{-1} = \frac{1}{b''(\theta)} \\ \frac{\partial \mu}{\partial \eta} &= \frac{\partial \mu}{\partial \eta}, \\ \frac{\partial \eta}{\partial \beta_j} &= x_j \end{split} \qquad \text{since } \mu = b'(\theta), \end{split}$$

Hence, we have:

$$\begin{split} \frac{\partial \ell}{\partial \beta_{j}} &= \frac{y - b'(\theta)}{a(\phi)} \frac{1}{b''(\theta)} \frac{\partial \mu}{\partial \eta} x_{j} \\ &= \frac{y - \mu}{\text{Var}(Y)} \frac{\partial \mu}{\partial \eta} x_{j} \qquad \text{since } \mu = b'(\theta), \text{Var}(Y) = a(\phi)b''(\theta) \\ &= \frac{y - \mu}{\text{Var}(Y)} \left(\frac{\partial \mu}{\partial \eta}\right)^{2} \frac{\partial \eta}{\partial \mu} x_{j} \qquad \text{since } \frac{\partial \mu}{\partial \eta} \frac{\partial \eta}{\partial \mu} = 1 \\ &= (y - \mu) \left(\text{Var}(Y) \left(\frac{\partial \mu}{\partial \eta}\right)^{2}\right)^{-1} \frac{\partial \eta}{\partial \mu} x_{j} \\ &= (y - \mu)W \frac{\partial \eta}{\partial \mu} x_{j}, \end{split}$$

where $W^{-1} = \mathrm{Var}(Y) \left(\frac{\partial \eta}{\partial \mu}\right)^2$. Note that generally $\frac{\partial \eta}{\partial \mu}$ is easier to calculate than $\frac{\partial \mu}{\partial \eta}$ since we define the link as $\eta = g(\mu)$.

For a random sample Y_1, \dots, Y_n from exponential family and each Y_i has a probability density function

$$f(y_i; \theta, \phi) = \exp\left\{\frac{y_i \theta_i - b(\theta_i)}{a(\phi)} + c(y_i, \phi)\right\}.$$

We write likelihood and log-likelihood functions as:

$$L = \prod_{i=1}^{n} f(y_i; \theta_i, \phi) = \prod_{i=1}^{n} \exp\left\{\frac{y_i \theta_i - b(\theta_i)}{a(\phi)} + c(y_i, \phi)\right\},$$
$$\ell = \sum_{i=1}^{n} \ell_i = \sum_{i=1}^{n} \frac{y_i \theta_i - b(\theta_i)}{a(\phi)} + c(y_i, \phi).$$

The element of the score vector is:

$$\left[\mathbf{S}(\boldsymbol{\beta}) \right]_{j} = \frac{\partial \ell}{\partial \beta_{j}} = \sum_{i=1}^{n} \frac{\partial \ell_{i}}{\partial \beta_{j}} = \sum_{i=1}^{n} (y_{i} - \mu_{i}) W_{i} \frac{\partial \eta_{i}}{\partial \mu_{i}} x_{ij}$$

where $W^{-1} = \operatorname{Var}(Y_i)(\frac{\partial \eta_i}{\partial \mu_i})^2$, $g(\mu_i) = \eta_i = \boldsymbol{x}_i^{\top}\boldsymbol{\beta}$. In vector and matrix form we can write:

$$S(\beta) = XWA(y - \mu),$$

where

- $\boldsymbol{y} = (y_1, \dots, y_n)^{\top}$ and $\boldsymbol{\mu} = (\mu_1, \dots, \mu_n)^{\top}$ are $n \times 1$ vectors,
- $X = (x_1, \dots, x_n)$ is a $(p+1) \times n$ matrix,

•
$$\mathcal{W}=\mathrm{diag}(W_1,\ldots,W_n)=egin{bmatrix} W_1 & 0 & \cdots & 0 \\ \vdots & \ddots & \ddots & \vdots \\ 0 & \cdots & 0 & W_n \end{bmatrix}$$
, and

•
$$\mathcal{A} = \operatorname{diag}\left(\frac{\partial \eta_1}{\partial \mu_1}, \dots, \frac{\partial \eta_n}{\partial \mu_n}\right)$$
.

Example: Poisson Regression Model (Problem 1.4)

For a random sample from Poisson distribution, $Y_i \sim POI(\lambda_i)$, i = 1, ..., n

$$\ell_i = \log(f(y_i; \lambda_i)) = (y_i \log(\lambda_i) - \lambda_i - \log(y_i!)).$$

Poisson regression with a log-link:

$$\log(\lambda_i) = \eta_i = \boldsymbol{x}_i^{\top} \boldsymbol{\beta}.$$

To write down the score vector for the regression coefficients β , we may calculate the derivative using standard methods, i.e.,

$$[S(\beta)]_{j} = \sum_{i} \frac{\partial \ell_{i}}{\partial \beta_{j}}$$

$$= \sum_{i} \frac{\partial}{\partial \beta_{j}} (y_{i} \log(\lambda_{i}) - \lambda_{i} - \log(y_{i}!))$$

$$= \sum_{i} (y_{i} x_{ij} - e^{x_{i}^{\top} \beta} x_{ij}).$$

Or we can use the general results derived for the GLMs on the previous slides.

Solving $S(\beta) = 0$ for MLE

1 Newton Raphson update equation is:

$$\hat{\beta}^{(r+1)} = \hat{\beta}^{(r)} + I^{-1}(\hat{\beta}^{(r)})S(\hat{\beta}^{(r)}).$$

where I is the observed information matrix.

- This requires us to find and repeatedly evaluate the information *I* (possibly computationally intensive).
- Fisher suggested using the expected information matrix $\mathcal I$ rather than the observed information matrix.
- 2 Fisher Scoring update equation is:

$$\hat{\boldsymbol{\beta}}^{(r+1)} = \hat{\boldsymbol{\beta}}^{(r)} + \boldsymbol{\mathcal{I}}^{-1}(\hat{\boldsymbol{\beta}}^{(r)})\boldsymbol{S}(\hat{\boldsymbol{\beta}}^{(r)}).$$

Information Matrix

Consider the (j, k) element of the Information matrix:

$$\begin{split} I_{jk} &= -\frac{\partial^2 \ell}{\partial \beta_j} \frac{\partial \ell}{\partial \beta_k} \\ &= -\frac{\partial}{\partial \beta_k} \frac{\partial \ell}{\partial \beta_j} \\ &= \sum_i -\frac{\partial}{\partial \beta_k} \left[(y_i - \mu_i) W_i \left(\frac{\partial \eta_i}{\partial \mu_i} \right) x_{ij} \right] \\ &= \sum_i -(y_i - \mu_i) \left\{ \frac{\partial}{\partial \beta_k} \left[W_i \left(\frac{\partial \eta_i}{\partial \mu_i} \right) x_{ij} \right] \right\} - W_i \left(\frac{\partial \eta_i}{\partial \mu_i} \right) x_{ij} \left(\frac{\partial}{\partial \beta_k} (y_i - \mu_i) \right) \\ &= \sum_i -(y_i - \mu_i) \left\{ \frac{\partial}{\partial \beta_k} \left[W_i \left(\frac{\partial \eta_i}{\partial \mu_i} \right) x_{ij} \right] \right\} + W_i \left(\frac{\partial \eta_i}{\partial \mu_i} \right) x_{ij} \frac{\partial \mu_i}{\partial \eta_i} \frac{\partial \eta_i}{\partial \beta_k} \\ &= \sum_i -(y_i - \mu_i) \frac{\partial}{\partial \beta_k} \left[W_i \left(\frac{\partial \eta_i}{\partial \mu_i} \right) x_{ij} \right] + x_{ij} W_i x_{ik}. \end{split}$$

FISHER INFORMATION

To get an element of the Expected/Fisher Information matrix:

$$\begin{split} \mathcal{I}_{jk} &= \sum_{i} \mathbf{E} \left[-\frac{\partial^{2} \ell}{\partial \beta_{j} \, \partial \beta_{k}} \right] \\ &= \sum_{i} \mathbf{E} \left[-(y_{i} - \mu_{i}) \frac{\partial}{\partial \beta_{k}} \left[W_{i} \left(\frac{\partial \eta_{i}}{\partial \mu_{i}} \right) x_{ij} \right] + x_{ij} W_{i} x_{ik} \right] \\ &= \sum_{i} -\mathbf{E} \left[(y_{i} - \mu_{i}) \right] \frac{\partial}{\partial \beta_{k}} \left[W_{i} \left(\frac{\partial \eta_{i}}{\partial \mu_{i}} \right) x_{ij} \right] + x_{ij} W_{i} x_{ik} \\ &= \sum_{i} x_{ij} W_{i} x_{ik}. \end{split}$$

Therefore, we can write the (j, k) element of the Fisher information as:

$$\mathcal{I}_{jk} = \sum_{i=1}^{n} x_{ij} W_i x_{ik} = [\boldsymbol{X} \boldsymbol{\mathcal{W}} \boldsymbol{X}^{\top}]_{jk}$$

where again, $\mathbf{W} = \operatorname{diag}(W_1, \dots, W_n)$ and $W_i^{-1} = \operatorname{Var}(Y_i) \left(\frac{\partial \eta_i}{\partial \mu_i}\right)^2$.

WHEN IS FISHER SCORING EQUIVALENT TO NEWTON RAPHSON?

Recall information matrix:

$$I_{jk} = \sum_{i} -(y_i - \mu_i) \frac{\partial}{\partial \beta_k} \left[W_i \left(\frac{\partial \eta_i}{\partial \mu_i} \right) x_{ij} \right] + x_{ij} W_i x_{ik}.$$

Now examine:

$$W_{i} \left(\frac{\partial \eta_{i}}{\partial \mu_{i}} \right) x_{ij} = \left(\operatorname{Var}(Y_{i}) \left(\frac{\partial \eta_{i}}{\partial \mu_{i}} \right)^{2} \right)^{-1} \left(\frac{\partial \eta_{i}}{\partial \mu_{i}} \right) x_{ij}$$

$$= \left(a(\phi)b''(\theta_{i}) \frac{\partial \eta_{i}}{\partial \mu_{i}} \right)^{-1} x_{ij} \qquad \text{since } \operatorname{Var}(Y_{i}) = a_{i}(\phi)b''(\theta_{i})$$

$$= \left(a(\phi) \frac{\partial \mu_{i}}{\partial \theta_{i}} \frac{\partial \eta_{i}}{\partial \mu_{i}} \right)^{-1} x_{ij} \qquad \text{since } b'(\theta_{i}) = \mu_{i}, b''(\theta_{i}) = \frac{\partial \mu_{i}}{\partial \theta_{i}}$$

$$= \left(a(\phi) \right)^{-1} x_{ij} \qquad \text{under the canonical link } \theta_{i} = \eta_{i}.$$

So under the canonical link,

$$\frac{\partial}{\partial \beta_k} \left[W_i \left(\frac{\partial \eta_i}{\partial \mu_i} \right) x_{ij} \right] = \frac{\partial}{\partial \beta_k} \left[\left(a(\phi) \right)^{-1} x_{ij} \right] = 0,$$

therefore information matrix is same as the Fisher information:

$$I_{jk} = \sum_{i} x_{ij} W_i x_{ij} = \mathcal{I}_{jk}$$

and Fisher Scoring is equivalent to Newton Raphson.

ITERATIVELY REWEIGHTED LEAST SQUARES

The Fisher Scoring is also called iteratively reweighted least squares (IRWLS). The reason is that the update equation can be rewritten as:

$$\hat{\boldsymbol{\beta}}^{(r+1)} = \left(\boldsymbol{X}\boldsymbol{\mathcal{W}}\big(\hat{\boldsymbol{\beta}}^{(r)}\big)\boldsymbol{X}^{\top}\right)^{-1}\!\boldsymbol{X}\boldsymbol{\mathcal{W}}\big(\hat{\boldsymbol{\beta}}^{(r)}\big)\boldsymbol{Z}\big(\hat{\boldsymbol{\beta}}^{(r)}\big)$$

where Z is a transformation of the response vector Y such that:

$$oldsymbol{Z} = oldsymbol{\eta} + (oldsymbol{Y} - oldsymbol{\mu}) * rac{\partial oldsymbol{\eta}}{\partial oldsymbol{\mu}}$$

- See manipulation in Section 1.2.3 of course notes.
- Same form as the weighted LS estimate of β with dependent variable Z and weight matrix \mathcal{W} .
- Z and \mathcal{W} are updated at each iteration.

TOPIC SUMMARY

2b Maximum Likelihood Estimation of Generalized Linear Models:

 When Y_i come from a distribution in the exponential family, we can use the theory of Generalized Linear Models to fit the regression equations of the form:

$$g(\mu_i) = \boldsymbol{x}_i^{\top} \boldsymbol{\beta}.$$

- The link function $g(\cdot)$ may be the canonical link, but its choice should come from model interpretation and fit.
- Can use Fisher Scoring (also known as IRWLS) to estimate the regression parameters β from any GLM based on general forms for $I(\beta)$ and $S(\beta)$.
- PRACTICE: Chapter 1 review problems.

Topic 3a: Binary Data and Odds Ratios

BINARY DATA SET-UP

Consider the simplest case with two binary variables:

- COVID-19: infected or not infected (response).
- Vaccination: yes or no (explanatory variable).

Use a 2×2 table to summarize the data:

	COVID-19			
Vaccination	infected	not infected		
yes	y_1	$m_1 - y_1$	m_1	
no	y_2	m_2-y_2	m_2	
Total	y_{ullet}	$m_{\bullet} - y_{\bullet}$	m_{ullet}	

Treat m_1 and m_2 as fixed, assume Y_1 and Y_2 are independent binomial r.v.'s

$$Y_k \sim \text{BIN}(m_k, \pi_k), \qquad k = 1, 2,$$

where $\pi_k = P(\text{infection} \mid \text{group } k)$.

How do we measure the associate between COVID-19 infection and vaccination?

MEASURES OF ASSOCIATION

Definition (Odds Ratio)

The Odds Ratio (OR) is the ratio of the odds of an event occurring in one group to the odds of the event in another group (e.g., not vaccinated):

Odds Ratio =
$$\frac{\pi_1/(1-\pi_1)}{\pi_2/(1-\pi_2)}$$
.

Interpretation of OR:

Relative Risk (RR)

The Relative Risk (RR) is the ratio of the probability of an event occurring in one group versus another group:

Relative Risk =
$$\frac{\pi_1}{\pi_2}$$

In the case of a rare disease (i.e., when π_1 and π_2 are very small),

OR =
$$\frac{\pi_1/(1-\pi_1)}{\pi_2/(1-\pi_2)} = \frac{\pi_1}{\pi_2} \underbrace{\left(\frac{1-\pi_2}{1-\pi_1}\right)}_{\approx 1} \approx \frac{\pi_1}{\pi_2} = RR,$$

then

$$OR \approx RR$$
.

MAXIMUM LIKELIHOOD ESTIMATION OF ODDS RATIO

Goal: Estimate odds ratio $\psi = \frac{\pi_1/(1-\pi_1)}{\pi_2/(1-\pi_2)}$ using likelihood method. Based on "grouped" binomial data, $Y_k \sim \text{BIN}(m_k, \pi_k), \ k = 1, 2,$

$$\begin{split} L(\pi_1,\pi_2) &= \binom{m_1}{y_1} \pi_1^{y_1} (1-\pi_1)^{m_1-y_1} \binom{m_2}{y_2} \pi_2^{y_2} (1-\pi_2)^{m_2-y_2} \\ &\propto \left(\frac{\pi_1/(1-\pi_1)}{\pi_2/(1-\pi_2)}\right)^{y_1} \left(\frac{\pi_2}{1-\pi_2}\right)^{y_2+y_1} (1-\pi_1)^{m_1} (1-\pi_2)^{m_2}. \end{split}$$

Note that $\pi_1, \pi_2 \in [0, 1]$ and odds ratio $\psi \in (0, \infty)$ are restricted, we consider re-parameterize:

$$\theta_1 = \log\left(\frac{\pi_1/(1-\pi_1)}{\pi_2/(1-\pi_2)}\right) = \log(\psi), \qquad \theta_2 = \log\left(\frac{\pi_2}{1-\pi_2}\right),$$

and now $\theta_1, \theta_2 \in (-\infty, \infty)$.

Our re-parameterization implies:

$$\pi_1 = \frac{e^{\theta_1 + \theta_2}}{1 + e^{\theta_1 + \theta_2}}, \qquad \pi_2 = \frac{e^{\theta_2}}{1 + e^{\theta_2}}.$$

Now the likelihood becomes:

$$L(\theta_1, \theta_2) = (e^{\theta_1})^{y_1} (e^{\theta_2})^{y_1 + y_2} (1 + e^{\theta_1 + \theta_2})^{m_1} (1 + e^{\theta_2})^{-m_2},$$

$$\ell(\theta_1, \theta_2) = y_1 \theta_1 + (y_1 + y_2) \theta_2 - m_1 \log(1 + e^{\theta_1 + \theta_2}) - m_2 \log(1 + e^{\theta_2}).$$

The score vector is:

$$S(\theta_1, \theta_2) = \begin{pmatrix} \frac{\partial \ell}{\partial \theta_1} \\ \frac{\partial \ell}{\partial \theta_2} \end{pmatrix} = \begin{pmatrix} y_1 - m_1 \left(\frac{e^{\theta_1 + \theta_2}}{1 + e^{\theta_1 + \theta_2}} \right) \\ y_1 + y_2 - m_1 \left(\frac{e^{\theta_1 + \theta_2}}{1 + e^{\theta_1 + \theta_2}} \right) - m_2 \left(\frac{e^{\theta_2}}{1 + e^{\theta_2}} \right) \end{pmatrix}.$$

Solving $S(\theta_1, \theta_2) = \mathbf{0}$ gives us the MLEs:

$$\hat{\theta}_1 = \log\left(\frac{y_1/(m_1 - y_1)}{y_2/(m_2 - y_2)}\right), \qquad \hat{\theta}_2 = \log\left(\frac{y_2}{m_2 - y_2}\right).$$

So by the invariance property of MLEs, we have:

$$\hat{\pi}_1 = \frac{y_1}{m_1}, \qquad \hat{\pi}_2 = \frac{y_2}{m_2}, \qquad \hat{\psi} = \frac{\hat{\pi}_1/(1-\hat{\pi}_1)}{\hat{\pi}_2/(1-\hat{\pi}_2)} = \frac{y_1/(m_1-y_1)}{y_2/(m_2-y_2)}.$$

INFERENCE FOR ODDS RATIO

In order to do inference we will need the Information Matrix:

$$m{I}(heta_1, heta_2) = egin{bmatrix} I_{11} & I_{12} \ I_{21} & I_{22} \end{bmatrix} \qquad ext{where } I_{jk} = -rac{\partial^2}{\partial heta_j\,\partial heta_k}\ell(heta_1, heta_2).$$

Here, we have:

$$I_{11} = m_1 \left(\frac{e^{\theta_1 + \theta_2}}{(1 + e^{\theta_1 + \theta_2})^2} \right),$$

$$I_{12} = I_{21} = m_1 \left(\frac{e^{\theta_1 + \theta_2}}{(1 + e^{\theta_1 + \theta_2})^2} \right),$$

$$I_{22} = m_1 \left(\frac{e^{\theta_1 + \theta_2}}{(1 + e^{\theta_1 + \theta_2})^2} \right) + m_2 \left(\frac{e^{\theta_2}}{(1 + e^{\theta_2})^2} \right).$$

We are interested in doing inference on $\theta_1 = \log(\psi)$ (while θ_2 is nuisance). Recall the asymptotic distribution result of a Wald statistic:

Wald Statistic

For a vector $\boldsymbol{\theta} = (\theta_1, \theta_2)^{\top}$ where $\theta_1 = \log(\psi)$ is a scalar parameter of interest:

$$(\hat{\theta}_1 - \theta_1)^2 (I^{11}(\hat{\theta}_1, \hat{\theta}_2))^{-1} \sim \chi^2_{(1)},$$

where I^{11} is the (1,1) element of I^{-1} evaluated at MLE $\hat{\theta}_1$ and $\hat{\theta}_2$.

• Calculation of I^{11} by using a general result:

$$I = \begin{pmatrix} I_{11} & I_{12} \\ I_{21} & I_{22} \end{pmatrix}, \qquad I^{-1} = \begin{pmatrix} I^{11} & I^{12} \\ I^{21} & I^{22} \end{pmatrix}, \qquad I^{11} = \begin{pmatrix} I_{11} - I_{12}I_{21}^{-1}I_{21} \end{pmatrix}^{-1}.$$

• We can use the Wald result to find a confidence interval for $\theta_1 = \log(\psi)$.

CONFIDENCE INTERVAL FOR ODDS RATIO

Here, we obtain:

$$I^{11}(\hat{\theta}_1, \hat{\theta}_2) = \frac{1}{y_1} + \frac{1}{m_1 - y_1} + \frac{1}{y_2} + \frac{1}{m_2 - y_2}.$$

Thus, a Wald-based 95 % confidence interval for $\theta_1 = \log(\psi)$ is:

$$\hat{\theta}_1 \pm 1.96 \sqrt{\frac{1}{y_1} + \frac{1}{m_1 - y_1} + \frac{1}{y_2} + \frac{1}{m_2 - y_2}} = (\hat{\theta}_{1\mathrm{L}}, \hat{\theta}_{1\mathrm{U}}).$$

A $95\,\%$ confidence interval for the Odds Ratio ψ is:

$$\left(\exp\{\hat{\theta}_{1L}\},\exp\{\hat{\theta}_{1U}\}\right).$$

EXAMPLE: PRENATAL CARE FROM TWO CLINICS

Consider the data below for the relationship between:

- Response: Fetal Mortality.
- Explanatory variable: Level of Care.

	Fetal		
Level of Care	Died	Survived	Total
Intensive Regular	20 46	316 373	336 419
	66	689	755

• Using the above data, we obtain MLE of odds ratio ψ :

$$\hat{\psi} = \frac{y_1/(m_1 - y_1)}{y_2/(m_2 - y_2)} = \frac{20/316}{46/373} = 0.51.$$

 $\hat{\psi} = 0.51 < 1$, the risk of mortality is lower with intensive care.

• A 95 % CI for $\theta_1 = \log(\psi)$:

$$\log(0.51) \pm 1.96\sqrt{\frac{1}{20} + \frac{1}{316} + \frac{1}{46} + \frac{1}{373}} = (-1.219, -0.127).$$

• A 95 % CI for odds ratio ψ :

$$(\exp\{-1.219\}, \exp\{-0.127\}) = (0.30, 0.89).$$

Note that the CI does not cover the value $\psi=1$ (no association), so we reject the null hypothesis of no association between fetal mortality and level of care. In other words, there is evidence of association.

EXAMPLE: PRENATAL CARE FROM TWO CLINICS

There is an additional explanatory variable: Clinic (A vs B).

Prenatal Care Data Stratified by Clinic

1 1 6	Clinic A			D: 1	Clinic B	l m . 1
Level of Care	Died	Survived	Total	Died	Survived	Total
Intensive	16	293	309	4	23	27
Regular	12	176	188	34	197	231
	28	469	497	38	220	258

- $\hat{\psi}_A=0.80~(0.37,1.73)$ and $\hat{\psi}_B=1.01~(0.33,3.10)$. These cover value 1, different from the results from the pooled analysis on the previous slide.
- These results do NOT agree with the results from the pooled analysis on the previous slide.

Association Between Clinic and Level of Care

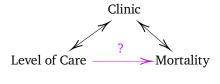
	A	В	
Intensive Regular	309 118	27 231	336 419
	497	258	755

• $\hat{\psi} = 14.06 \ (9.12, 21.76).$

Association Between Clinic and Mortality

	A	В	
Died	28	38	66
Survived	469	220	689
	497	258	755

- $\hat{\psi} = 0.35 \ (0.21, 0.58).$
- The initial strong association between Level of Care and Infant Morality disappeared when we stratified by clinic.



- Instead of having to examine multiple 2×2 tables we'd like to estimate the OR and compute associations using a multiple regression model.
- One way to do this is by fitting a Binomial GLM to the data.

WEEK 4 0927 to 1st October

Topic 3b: Binomial Regression Models for Binary Data

RECALL TOPIC 3A: BINARY DATA AND ODDS RATIOS

Last week, we introduce a simple method for association between two binary variables, 2×2 contingency table analysis: Measure of Association: $OR = \psi = \frac{\pi_1/(1-\pi_1)}{\pi_2/(1-\pi_2)}$,

		ortality	
Level of Care	Died	Survived	
Intensive Regular	y_1 y_2	$m_1 - y_1$ $m_2 - y_2$	$Y_1 \sim \text{BIN}(m_1, \pi_1)$ $Y_2 \sim \text{BIN}(m_2, \pi_2)$

- OR = 1 (equal risk).
- 0 < OR < 1 (lower risk in group 1).
- OR > 1 (higher risk in group 1).

Maximum likelihood estimator for OR is:

$$\hat{\psi} = \frac{y_1/(m_1 - y_1)}{y_2/(m_2 - y_2)},$$

and a Wald-based 95% CI is:

$$\exp\left\{\log(\hat{\psi}_1) \pm 1.96 \underbrace{\sqrt{\frac{1}{y_1} + \frac{1}{m_1 - y_1} + \frac{1}{y_2} + \frac{1}{m_2 - y_2}}}_{\text{se}(\log(\hat{\psi}))}\right\}$$

Prenatal Care Data Example: However, Mortality and Care are also related to another variable, Clinic:

Table 1: $1 \notin (0.30, 0.89) \implies$ evidence of association between Mortality and Care.

OR (Mortality and Clinic)	Est.	$95\%~\mathrm{CI}$
Intensive vs Regular	0.35	(0.12, 0.58)

Table 2: Association between Mortality and Clinic.

OR (Care and Clinic)	Est.	95% CI
Intensive vs Regular	14.06	(9.12, 21.76)

Table 3: Association between Care and Clinic.

- Therefore, we wish to consider how a variable, e.g., Mortality (Y), is related to multiple explanatory variables together, e.g., Care (x_1) and Clinic (x_2) .
- This can be done using multiple regression methodology for binary data \implies Topic 3b: Binomial Regression Models for Binary Data.

MULTIPLE REGRESSION FOR BINARY DATA

- Often we need to consider the relationship between a binary outcome and multiple explanatory variables, using multiple regression methodology.
- This is because we may want to:
 - control for cofounding variables and hence want to examine the effect of several variables simultaneously;
 - examine the effect of categorical variables (> 2 levels) or continuous covariates;
 - develop sophisticated models that describe complex relationship.
- Suppose *subject level data* is binary with a value of 1 indicating that an event of interest occurs and a value of 0 indicating that event doesn't occur.
- Subjects can be classified according to the values of explanatory variables into *n* groups (i.e., common covariates values within each group), so we have *grouped data* such that:
 - m_i denotes number of subjects in group i;
 - Y_i denotes number of subjects experienced the event in group i;
 - x_{i1}, \ldots, x_{ip} denote the covariates values associated with group i where $i = 1, \ldots, n$.

SET-UP OF A BINOMIAL REGRESSION MODEL

Response Variable: $Y_i \sim \text{BIN}(m_i, \pi_i)$, i = 1, ..., n, and Binomial distribution is a member of Exponential family!

$$f(y_i) = {m_i \choose y_i} \pi_i^{y_i} (1 - \pi_i)^{m_i - y_i}$$

= $\exp \left\{ y_i \log \left(\frac{\pi_i}{1 - \pi_i} \right) + m_i \log (1 - \pi_i) + \log \left(\binom{m_i}{y_i} \right) \right\},$

where

$$\theta_i = \log\left(\frac{\pi_i}{1 - \pi_i}\right),$$

$$a(\phi) = \phi = 1,$$

$$b(\theta_i) = -m_i \log(1 - \pi_i) = m_i \log(1 + e^{\theta_i}).$$

$$c(y_i; \phi) = \log\left(\binom{m_i}{y_i}\right).$$

(2) Linear Predictor:

$$\eta_i = \boldsymbol{x}_i^{\top} \boldsymbol{\beta} = \beta_0 + \beta_1 x_{i1} + \dots + \beta_p x_{ip}.$$

3 Link Function: Recall that for Binomial distribution, we have $E[Y_i] = \mu_i = m_i \pi_i$, therefore we typically re-write the link function in terms of π_i ,

$$g(\pi_i) = \boldsymbol{x}_i^{ op} \boldsymbol{eta}.$$

As $\pi_i \in (0,1)$, any function $g:(0,1) \to (-\infty,\infty)$ may work, and here are some link functions we can consider:

log-log	$g(\pi) = \log(-\log(\pi))$
complementary log-log	$g(\pi) = \log(-\log(1-\pi))$
$Probit^a$	$g(\pi) = \Phi^{-1}(\pi)$
Logit (canonical)	$g(\pi) = \log(\pi/(1-\pi))$

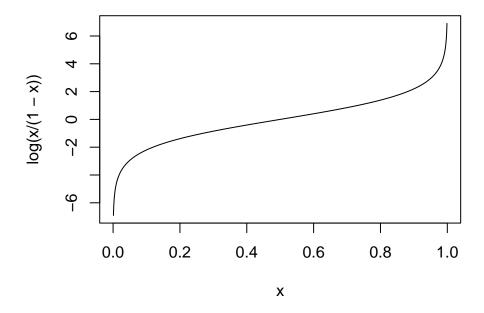
^aFor the Probit link, $\Phi(\cdot)$ is the *CDF* of $\mathcal{N}(0,1)$.

CANONICAL LINK AND LOGISTIC REGRESSION

Recall for Binomial distribution $\theta_i = \log\left(\frac{\pi_i}{1-\pi_i}\right)$, and by setting $\theta_i = \eta_i$, we have:

$$\log\left(\frac{\pi_i}{1-\pi_i}\right) = \eta_i.$$

The Logit link, $g(\pi_i) = \log(\pi_i/(1-\pi_i))$, is the canonical link for the Binomial!



This leads to a Logistic Regression Model:

$$\log \left(\frac{\pi_i}{1-\pi_i}\right) = \boldsymbol{x}_i^{\top} \boldsymbol{\beta} = \beta_0 + \beta_1 x_{i1} + \dots + \beta_p x_{ip}.$$

PREDICTION FROM LOGISTIC REGRESSION

Aside: The inverse of the logit function is called the expit function:

$$\operatorname{logit}(\pi_i) = \operatorname{log}\left(\frac{\pi_i}{1 - \pi_i}\right) = \boldsymbol{x}_i^{\top} \boldsymbol{\beta} \iff \pi_i = \frac{\exp\{\boldsymbol{x}_i^{\top} \boldsymbol{\beta}\}}{1 + \exp\{\boldsymbol{x}_i^{\top} \boldsymbol{\beta}\}} = \operatorname{expit}(\boldsymbol{x}_i^{\top} \boldsymbol{\beta}).$$

Suppose we have found MLE $\hat{\beta}$ using Fisher scoring, then the fitted value for the probability of response π_i given explanatory variables x_i is:

$$\hat{\pi}_i = \frac{\exp\{\boldsymbol{x}_i^{\top} \hat{\boldsymbol{\beta}}\}}{1 + \exp\{\boldsymbol{x}_i^{\top} \hat{\boldsymbol{\beta}}\}}.$$

The predicted number of responses are: $\hat{Y}_i = m_i \hat{\pi}_i$.

Interpretation of eta in Logistic Regression

• Consider a simple logistic model with a single binary explanatory variable:

$$\log\left(\frac{\pi_i}{1-\pi_i}\right) = \beta_0 + \beta_1 x_{i1},$$

where $x_{i1} = 0$ (group 0) and $x_{i1} = 1$ (group 1).

- Let's compare the model when $x_{i1} = 1$ vs $x_{i1} = 0$.
- We subtract line 2 from line 1 to isolate β_1 and find its interpretation.
- $\beta_1 = \log$ odds ratio of response for subjects with $x_{i1} = 1$ vs $x_{i1} = 0$.
- Please see Section 2.4.2 for general interpretations of β 's in multiple logistic regression models.

Group
$$x_i^{\top}$$
 $\eta_i = \log(\pi_i/(1-\pi_i))$

$$\begin{array}{cccc} 1 & (1,1)^{\top} & \beta_0 + \beta_1 & = \log(\pi_1/(1-\pi_1)) \\ 0 & (1,0)^{\top} & \beta_0 & = \log(\pi_0/(1-\pi_0)) \\ \end{array}$$

$$\beta_1 = \log\left(\frac{\pi_1/(1-\pi_1)}{\pi_0/(1-\pi_0)}\right) = \log(OR)$$

LOGISTIC REGRESSION FOR PRENATAL CARE EXAMPLE

• Response: Fetal Mortality, that is,

$$Y_i \sim BIN(m_i, \pi_i), i = 1, 2,$$

• Explanatory Variables:

$$x_{i1} = \begin{cases} 1 & \text{Intensive Care} \\ 0 & \text{Regular Care} \end{cases}$$

$$x_{i2} = \begin{cases} 1 & \text{Clinic A} \\ 0 & \text{Clinic B} \end{cases}$$

$$x_{i3} = x_{i1}x_{i2} = \begin{cases} 1 & \text{Intensive care and Clinic A} \\ 0 & \text{Otherwise} \end{cases}$$

- We will use the context of this example to illustrate how to:
 - fit (simple and multiple) logistic regression models using R, and
 - interpret regression parameters.

MODEL 1: LEVEL OF CARE ONLY MODEL

$$\log\left(\frac{\pi_i}{1-\pi_i}\right) = \beta_0 + \beta_1 x_{i1}.$$

Level of Care	Clinic	\boldsymbol{x}_i^\top	$\log(\pi_i/(1-\pi_i))$
Intensive	_	$(1,1)^{\top}$	$\beta_0 + \beta_1$
Regular	_	$(1,0)^{\top}$	eta_0

- $\beta_0 = \log \text{ odds}$ of mortality for babies born to mothers treated with regular care.
- $\beta_1 = \log \text{ odds ratio}$ of mortality for babies born to mothers treated with intensive vs regular care.

MODEL 2: MAIN EFFECTS MODEL

$$\log\left(\frac{\pi_i}{1-\pi_i}\right) = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2}.$$

- $\beta_0 = \log \text{ odds}$ of mortality with regular care at Clinic B.
- $\beta_1 = \log$ odds ratio of mortality for babies born to mothers treated with intensity vs regular care at the *same clinic*.
- $\beta_2 = \log$ odds ratio of mortality for babies born to mothers treated at Clinic A vs Clinic B at the *same* level of care.

Level of Care	Clinic	\boldsymbol{x}_i^\top	$\log(\pi_i/(1-\pi_i))$
Intensive	Α	$(1,1,1)^{\top}$	$\beta_0 + \beta_1 + \beta_2$
Regular	Α	$(1,0,1)^{\top}$	$\beta_0 + \beta_2$
Intensive	В	$(1,1,0)^{\top}$	$\beta_0 + \beta_1$
Regular	В	$(1,0,0)^{\top}$	eta_0

MODEL 3: INTERACTION MODEL

$$\log\left(\frac{\pi_i}{1-\pi_i}\right) = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2} + \beta_3 x_{i3}.$$

Level of Care	Clinic	\boldsymbol{x}_i^\top	$\log(\pi_i/(1-\pi_i))$
Intensive	Α	$(1,1,1)^{\top}$	$\beta_0 + \beta_1 + \beta_2 + \beta_3$
Regular	Α	$(1,0,1)^{\top}$	$\beta_0 + \beta_2$
Intensive	В	$(1,1,0)^{\top}$	$\beta_0 + \beta_1$
Regular	В	$(1,0,0)^{\top}$	eta_0

- $\beta_0 = \log$ odds ratio of mortality for babies born to mothers treated with intensity vs regular care at *Clinic B*.
- $\beta_1 + \beta_3 = \log$ odds ratio of mortality for babies born to mothers treated with intensity vs regular care at *Clinic A*.
- $\beta_2 = \log$ odds ratio of mortality for babies born to mothers treated at Clinic A vs Clinic B with *regular* care.
- $\beta_2 + \beta_3 = \log$ odds ratio of mortality for babies born to mothers treated at Clinic A vs Clinic B with *intensive* care.
- β_3 represents the difference in log odds ratios.
- If $\beta_3 = 0$ then association between mortality and level of care does not dependent on clinic.
- Equivalently, if $\beta_3 = 0$ then the association between mortality and clinic does not depend on level of care.

Data file prenatal.dat

```
clinic loc y m
1 0 0 34 231
2 0 1 4 27
3 1 0 12 188
4 1 1 16 309
```

- The first line contains the variable names/labels.
- We are using indicator variables for the explanatory variables:

$$x_{i1} = loc$$
 (1 for Intensive, 0 for Regular)
 $x_{i2} = clinic$ (1 for Clinic A, 0 for Clinic B)

• The variable y records the number of deaths (events).

FIT GLMs USING R

The glm() function in R is used to fit the generalized linear models:

```
fit = glm(formula, family = (link = ), data = ).
```

• formula: a linear formula describing the model, e.g.,

```
resp ~loc + clinic.
```

• family: a description of the exponential family distribution and link function to be used in the model, e.g.,

```
family = binomial, gaussian, poisson, Gamma, etc..
link = logit, log, loglog, cloglog, identity, probit, etc..
```

• The default is the canonical link.

R CODE AND OUTPUT FOR ANALYSIS OF PRENATAL CARE DATA

```
# read file prenatal.data
prenatal.dat = read.table("prenatal.dat", header = T)
# construct the binomial response for the logistic
# regression analysis
prenatal.dat$resp = cbind(prenatal.dat$y, prenatal.dat$m - prenatal.dat$y)
prenatal.dat
 clinic loc y m resp.1 resp.2
     0 0 34 231
                      34
                            197
                            23
                      4
      0 1 4 27
      1 0 12 188
                      12
                            176
      1 1 16 309
                            293
                  16
```

The logistic regression models are fit using the glm() commands like:

```
# fit the logistic model using the glm function
model1 = glm(resp ~ loc, family = binomial(link = logit), data = prenatal.dat)
summary(model1)
```

FIT OF MODEL 1: LEVEL OF CARE MODEL

$$\log\left(\frac{\pi_i}{1-\pi_i}\right) = \beta_0 + \beta_1 x_{i1}.$$

Components of the summary() output for glm objects:

- Estimate: the maximum likelihood estimates of the regression coefficients $\hat{\beta}_0$ and $\hat{\beta}_1$.
- Std. Error: estimated standard errors, the square root of the diagonal of the inverse of the Information matrix:

$$\operatorname{se}(\hat{\beta}_j) = \sqrt{\left[\boldsymbol{I}^{-1}(\hat{\boldsymbol{\beta}})\right]_{jj}} = \sqrt{I^{jj}(\hat{\boldsymbol{\beta}})}.$$

• z value: Wald-type test statistics for testing the hypotheses:

$$H_0$$
: $\beta_j = 0$ vs H_A : $\beta_j \neq 0$.

• Pr(>|z|): p-value for above Wald test.

For this model:

• β_1 is the log odds ratio of mortality for infants born to mothers treated with intensive versus regular care.

Hypothesis test for β_i

• We may wish to test:

$$H_0$$
: $\beta_j = \beta^*$ versus H_A : $\beta_j \neq \beta^*$.

• The general Wald result for a single parameter β_j is:

$$(\hat{\beta}_i - \beta^*)^2 (I^{jj}(\hat{\boldsymbol{\beta}}))^{-1} \sim \chi_1^2,$$

equivalently
$$\frac{\hat{\beta}_j - \beta^\star}{\operatorname{se}(\hat{\beta}_j)} \sim \mathcal{N}(0, 1)$$
 where $\operatorname{se}(\hat{\beta}_j) = \sqrt{I^{jj}(\hat{\beta})}$.

• We can find the *p*-value of this test using:

$$p = 2 \operatorname{P} \left(Z > \frac{|\hat{\beta}_j - \beta^*|}{\operatorname{se}(\hat{\beta}_j)} \right).$$

• The summary() output gives the test statistics and p-values for testing

$$H_0$$
: $\beta_i = 0$ vs H_A : $\beta_i \neq 0$.

Hypothesis test for β_1 from Model 1: Level of Care Model

• We wish to test:

$$H_0$$
: $\beta_1 = 0$ vs H_A : $\beta_1 \neq 0$

• Wald test:

$$z = \frac{\hat{\beta}_1 - 0}{\text{se}(\hat{\beta}_1)} = \frac{-0.6671}{0.2785} = -2.3949$$

p-value:

$$p = 2 P(Z > |-2.3949|) = 0.0166 < 0.05$$

- Therefore, we reject the null hypothesis that $\beta_1 = 0$.
- Estimate of OR for Mortality for Intensive vs Regular Care:

$$\hat{\psi} = \exp{\{\hat{\beta}_1\}} = \exp{\{-0.6670729\}} = 0.51.$$

• Confidence Interval for OR:

$$\exp\{\hat{\beta}_1 \pm 1.96 \operatorname{se}(\hat{\beta}_1)\} = \exp\{-0.6671 \pm 1.96(0.2785)\}$$
$$= (\exp\{-1.2130\}, \exp\{-0.1211\})$$
$$= (0.30, 0.89)$$

• The estimate and Wald 95% CI here match those found previously from the 2×2 table analysis. That is, the 2×2 table analysis is equivalent to a simple logistic regression with a single binary covariate.

FIT OF MODEL 2: MAIN EFFECTS MODEL

$$\log\left(\frac{\pi_i}{1-\pi_i}\right) = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2}.$$

What is the OR for mortality for Intensive vs Regular Care, now controlling for Clinic?

$$\widehat{OR} = \hat{\psi} = \exp\{-0.1503\} = 0.86.$$

• 95 % CI:

$$\exp\{-0.1503 \pm 1.96 \times 0.3302\} = (0.4505, 1.6436).$$

FIT OF MODEL 3: INTERACTION MODEL

$$\log\left(\frac{\pi_i}{1-\pi_i}\right) = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2} + \beta_3 x_{i3}.$$

```
model3 <- glm(resp ~ loc + clinic + loc * clinic, family = binomial(link = logit), data = prenatal.dat)

summary(model3)$coefficients

Estimate Std. Error z value Pr(>|z|)

(Intercept) -1.756843204 0.1857092 -9.46018403 3.074017e-21
loc 0.007643349 0.5726827 0.01334657 9.893513e-01
clinic -0.928734141 0.3514300 -2.64272868 8.224091e-03
loc:clinic -0.229649891 0.6949054 -0.33047646 7.410400e-01
```

Level of Care	Clinic	\boldsymbol{x}_i^\top	$\log(\pi_i/(1-\pi_i))$
Intensive	Α	$(1,1,1)^{\top}$	$\beta_0 + \beta_1 + \beta_2 + \beta_3$
Regular	Α	$(1,0,1)^{\top}$	$\beta_0 + \beta_2$
Intensive	В	$(1,1,0)^{\top}$	$\beta_0 + \beta_1$
Regular	В	$(1,0,0)^{\top}$	eta_0

• What is the OR for Mortality for Intensive vs Regular Care at Clinic A?

$$OR = \psi = \exp{\{\beta_1 + \beta_3\}} \implies \hat{\psi} = \exp{\{0.0076 - 0.2296\}} = 0.8.$$

- $se(\hat{\beta}_1 + \hat{\beta}_3)$ is required for calculation of 95 % CI.
 - Recall $\operatorname{Var}(\hat{\beta}) = I^{-1}(\hat{\beta})$, now for any linear function of β 's, e.g., $c\beta$ where c is a row vector of constants, then MLE of $c\beta$ is $c\hat{\beta}$, and $\operatorname{se}(\hat{c\beta}) = \sqrt{cI^{-1}(\hat{\beta})c^{\top}}$.
- Therefore, $\log(\psi) = \beta_1 + \beta_3 = c\beta$, $c = (0, 1, 0, 1)^{\mathsf{T}}$. In R, vcov(model3) gives $I^{-1}(\hat{\beta})$.
- What is OR for Mortality for Intensive vs Regular Care at Clinic B?

$$OR = \psi = \exp{\{\beta_1\}} \implies \hat{\psi} = \exp{\{0.0076\}} = 1.01.$$

Topic 3c: Likelihood Ratio Test for Logistic Regression Models

LOGISTIC REGRESSION MODELS

Recall major developments of Binomial logistic regression from last topic 3b: $Y_i \sim BIN(m_i, \pi_i)$, i = 1, ..., n independently, with covariate vector x_i and

$$\log\!\left(rac{\pi_i}{1-\pi_i}
ight) = oldsymbol{x}_i^ opoldsymbol{eta}.$$

- Estimation: $\hat{\beta}$ come from Fisher scoring using R function glm().
- Interpretation: $\exp\{\beta_i\}$ has OR interpretation.
- Hypothesis tests of H_0 : $\beta_j = 0$ using Wald statistic.
- Confidence Intervals: $\hat{\beta}_j \pm z_{1-\alpha/2} \operatorname{se}(\hat{\beta}_j)$.

LIKELIHOOD FOR LOGISTIC REGRESSION MODELS

• Log-likelihood for Binomial Distribution:

$$\ell = \log \left(\prod_{i=1}^{n} \pi_i^{y_i} (1 - \pi_i)^{m_i - y_i} \right)$$
$$= \sum_{i=1}^{n} y_i \log \left(\frac{\pi_i}{1 - \pi_i} \right) + m_i \log (1 - \pi_i).$$

• Using logit link we can re-parameterize the log-likelihood in terms of β :

$$\log \left(\frac{\pi_i}{1 - \pi_i}\right) = \boldsymbol{x}_i^{\top} \boldsymbol{\beta}, \qquad \pi_i = \frac{\exp\{\boldsymbol{x}_i^{\top} \boldsymbol{\beta}\}}{1 + \exp\{\boldsymbol{x}_i^{\top} \boldsymbol{\beta}\}}.$$

• Log likelihood for logistic regression:

$$\ell = \sum_{i=1}^{n} y_i \boldsymbol{x}_i^{\top} \boldsymbol{\beta} - m_i \log (1 + \exp{\{\boldsymbol{x}_i^{\top} \boldsymbol{\beta}\}}).$$

- Maximization of log-likelihood $\ell(\beta)$ gives MLE $\hat{\beta}$, and
 - estimated probability of response:

$$\hat{\pi}_i = e^{\boldsymbol{x}_i^{\top}\hat{\boldsymbol{\beta}}}/(1 + e^{\boldsymbol{x}_i^{\top}\hat{\boldsymbol{\beta}}}) = \operatorname{expit}(\boldsymbol{x}_i^{\top}\hat{\boldsymbol{\beta}}),$$

- estimated number of responses: $\hat{y}_i = m_i \hat{\pi}_i$.
- Questions:
 - How good is the model? How well do the estimated number of events \hat{y}_i approximate the observed data y_i ? (goodness of fit).
 - How much worse is the fit of a model when several of the covariates are excluded? (nested models):

$$H_0: \beta_k = \beta_{k+1} = 0 \text{ vs } H_a: \beta_k \neq 0 \text{ or } \beta_{k+1} \neq 0.$$

LIKELIHOOD RATIO TEST (GENERAL SETTING)

- Suppose $\ell(\theta)$ is the likelihood for a q-dimension parameter vector θ and let
 - $\tilde{\theta}$ be the q-dim MLE of θ (unconstrained/saturated, q = n),
 - $\hat{\theta}$ be the *p*-dim MLE of θ (constrained/unsaturated, p < q).
- Hypotheses:
 - H_0 : the constrained model is adequate (i.e., as good as the unconstrained).
 - H_A : constrained model is not adequate.
- Recall the Likelihood Ratio (LR) result:

Under
$$H_0$$
: $-2\log\left(\frac{L(\hat{\boldsymbol{\theta}})}{L(\tilde{\boldsymbol{\theta}})}\right) = -2\left[\ell(\hat{\boldsymbol{\theta}}) - \ell(\tilde{\boldsymbol{\theta}})\right] \sim \chi_{q-p}^2$.

• Reject H_0 at θ if

$$p\text{-value} = \mathrm{P}(\chi^2_{q-p} > -2\big[\ell(\hat{\boldsymbol{\theta}}) - \ell(\tilde{\boldsymbol{\theta}})\big]) < \alpha.$$

LIKELIHOOD RATIO TEST (LOGISTIC REGRESSION MODEL)

• Saturated (unconstrained) model MLEs:

$$\tilde{\pi}_i = \frac{y_i}{m_i}, \ i = 1, \dots, n.$$

- Binomial MLE without imposing any constraint.
- We will have $\tilde{y}_i = m_i \tilde{\pi}_i = y_i$, a perfect fit!
- Unsaturated (constrained) model MLEs:

$$\hat{\pi}_i = \operatorname{expit}(\boldsymbol{x}_i^{\top} \hat{\boldsymbol{\beta}}).$$

– Regression models are a way of imposing constraints on the estimation of π_i through p-dim regression coefficients β .

- We will have fitted number of responses $\hat{y}_i = m_i \hat{\pi}_i = m_i \operatorname{expit}(\boldsymbol{x}_i^{\top} \hat{\boldsymbol{\beta}})$.
- Hypotheses:
 - H_0 : the p-dim model, e.g., $\operatorname{logit}(\pi_i) = \boldsymbol{x}_i^{\top} \boldsymbol{\beta}$ is adequate.
 - H_A : the p-dim model, e.g., $\operatorname{logit}(\pi_i) = \boldsymbol{x}_i^{\top} \boldsymbol{\beta}$ is not adequate compared to the n-dim saturated model.
- Likelihood Ratio Statistic (also referred to as the Deviance):

$$D = -2 \left[\ell(\hat{\pi}) - \ell(\tilde{\pi}) \right]$$

$$= -2 \left(\sum_{i=1}^{n} \left(y_i \log(\hat{\pi}_i) + (m_i - y_i) \log(1 - \hat{\pi}_i) \right) - \sum_{i=1}^{n} \left(y_i \log(\tilde{\pi}_i) + (m_i - y_i) \log(1 - \tilde{\pi}_i) \right) \right)$$

$$= -2 \sum_{i=1}^{n} \left(y_i \log \left(\frac{y_i}{m_i \hat{\pi}_i} \right) + (m_i - y_i) \log \left(\frac{m_i - y_i}{m_i (1 - \hat{\pi}_i)} \right) \right).$$

• The LR/Deviance can also be written in a general form as:

$$D = 2\sum_{i=1}^{n} \sum_{j=1}^{2} \left(O_{ij} \log \left(\frac{O_{ij}}{E_{ij}} \right) \right).$$

- $O_{i1} = y_i$, $E_{i1} = m_i \hat{\pi}_i$ (observed and expected # of events).
- $O_{i2}=m_i-y_i,\,E_{i2}=m_i(1-\hat{\pi}_i)$ (observed and expected # of non-events).
- We expect $D \sim \chi^2_{n-p}$ under H_0 , and reject H_0 if $P(\chi^2_{n-p} > D) < \alpha$.
 - Unfortunately, this is not a great approximation.
 - Approximation is much better for testing nested unsaturated models though.

Example: Prenatal Care Data

• Model 2: Main Effects Model,

$$logit(\pi_i) = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2}.$$

- H_0 : Model 2 is adequate.
- H_A : Model 2 is not adequate compared to the saturated model.
- In R, the summary() output D is reported as the Residual Deviance.

- Deviance: D = 0.10693.

- p-value: $P(\chi_{n-p}^2 > D) = P(\chi_1^2 > D) = 0.7436689 \gg 0.05$.

• Do not reject the null hypothesis that Model 2 is adequate.

PEARSON STATISTIC

 The Pearson statistic is another statistic that can be used for assessing "overall" fit (or goodness of fit) of a Binomial model:

$$P = \sum_{i=1}^{n} \frac{(y_i - m_i \hat{\pi}_i)^2}{m_i \hat{\pi}_i (1 - \hat{\pi}_i)}.$$

- As with LR/Deviance statistic, $P \sim \chi^2_{n-p}$ under H_0 : the model is adequate.
- Note that *P* has the general form:

$$P = \sum_{i} \frac{(O_i - E_i)}{V_i}.$$

- The χ^2 approximation is a bit better than for deviance statistic D.
- Both are poor if the sample size (m_i) is small though.

TESTING NESTED NON-SATURATED MODELS

- The previous LR/Deviance test was for an unsaturated model vs a saturated model.
- Now consider two unsaturated models (p < q < n).

$$logit(\pi_i) = \beta_0 + \beta_1 x_{i1} + \dots + \beta_{p-1} x_{ip-1}$$
(1)

$$logit(\pi_i) = \beta_0 + \beta_1 x_{i1} + \dots + \beta_{p-1} x_{ip-1} + \dots + \beta_{q-1} x_{iq-1}$$
(2)

- Model (1) is *nested* within Model (2).
- H_0 : Model (1) fits the data as well as Model (2).

-
$$H_0$$
: $\beta_p = \cdots = \beta_{q-1} = 0$.

- H_A : Model (1) is inadequate compared to Model (2).
 - H_A : at least one of $\beta_p, \ldots, \beta_{q-1} \neq 0$.

Model	Dimension	MLEs
(1) Reduced model	p	$\hat{\pi}_i$
(2) Full model	q	π_i
Saturated model	n	$\tilde{\tau}_{\overline{t}}$

• LR/Deviance test of Model (1) vs Saturated Model:

$$D_0 = -2(\ell(\hat{\boldsymbol{\pi}}) - \ell(\tilde{\tilde{\boldsymbol{\pi}}})).$$

• LR/Deviance test of Model (2) vs Saturated Model:

$$D_{\mathsf{A}} = -2\big(\ell(\tilde{\boldsymbol{\pi}}) - \ell(\tilde{\tilde{\boldsymbol{\pi}}})\big).$$

• Now, we wish to conduct LR test of Model (1) vs Model (2):

$$\Delta D = D_0 - D_{\mathbf{A}} = -2(\ell(\hat{\boldsymbol{\pi}}) - \ell(\tilde{\boldsymbol{\pi}})).$$

• It can be shown that under H_0 : Model (1) is as adequate as Model (2),

$$\Delta D \sim \chi_{q-p}^2$$
.

- This approximation is much better than when testing an unsaturated model vs the saturated model.
- If $p = P(\chi_{q-p}^2 > \Delta D) < \alpha$ then reject H_0 .
 - Reduced model does not fit the data as well as Full model.
 - One or more of covariates x_{ip}, \ldots, x_{iq-1} is important (i.e., associated with the response).

EXAMPLE: PRENATAL CARE DATA

• Summary of Deviance ("residual deviance") from R output:

Model	Covariates	Deviance	Parameters	n-p
1	loc	10.814378	2	2
2	loc + clinic	0.106928	3	1
3	<pre>loc + clinic + loc*clinic</pre>	0	4	0
4	clinic	0.314841	2	2

- Compare nested models:
 - Model 2: $logit(\pi_i) = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2}$.
 - Model 4: $logit(\pi_i) = \beta_0 + \beta_2 x_{i2}$.
- Is level of care associated with fetal mortality after accounting for clinic?
 - H_0 : Model 4 is as adequate as Model 2 (e.g., $\beta_1 = 0$).
 - H_A : Model 4 is inadequate compared to Model 2 (e.g., $\beta_1 \neq 0$).
- LR test for comparing Model 4 vs Model 2, or equivalently testing hypotheses:

$$H_0$$
: $\beta_1 = 0$ vs H_A : $\beta_1 \neq 0$.

- We do not reject H_0 of no association between level and care and fetal mortality after controlling for Clinic.

```
model2 = glm(resp ~ loc + clinic, family = binomial, data = prenatal.dat)
model4 = glm(resp ~ clinic, family = binomial, data = prenatal.dat)
D = model4$deviance - model2$deviance
1 - pchisq(D, 2 - 1)
[1] 0.6484081
```

- This implies that level of care is no longer important when clinic is included in the model.
- It also implies that Model 4 is as adequate compared to Model 2.
- Finally, when testing a single parameter, e.g., H_0 : $\beta_1 = 0$, LR/Deviance test result is consistent with the Wald test result provided in the R output:

```
model2 = glm(resp ~ loc + clinic, family = binomial, data = prenatal.dat)
summary(model2)
Call:
glm(formula = resp ~ loc + clinic, family = binomial, data = prenatal.dat)
Deviance Residuals:
-0.08521 0.25805 0.13909 -0.11719
Coefficients:
         Estimate Std. Error z value Pr(>|z|)
(Intercept) -1.7410 0.1785 -9.755 < 2e-16 ***
          -0.1503 0.3302 -0.455 0.64894
loc
          clinic
Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
(Dispersion parameter for binomial family taken to be 1)
   Null deviance: 16.91763 on 3 degrees of freedom
Residual deviance: 0.10693 on 1 degrees of freedom
AIC: 23.262
Number of Fisher Scoring iterations: 3
```

SUMMARY OF LR/DEVIANCE TEST FOR LOGISTIC REGRESSION

• For Binomial GLM with logit link the LR/Deviance test statistic is:

$$D = \sum_{i=1}^{n} 2\left(y_i \log\left(\frac{y_i}{m_i \hat{\pi}_i}\right) + (m_i - y_i) \log\left(\frac{m_i - y_i}{m_i (1 - \hat{\pi}_i)}\right)\right).$$

- This is reported as the "Residual Deviance" in R glm summary output.
- Deviance statistic *D* can be used to:
 - Test adequacy/goodness of fit of a non-saturated logistic model:

$$D \stackrel{H_0}{\sim} \chi^2_{n-p}.$$

- Compare the fit of two nested-non saturated logistic models:

$$\Delta D = D_0 - D_{\rm A} \stackrel{H_0}{\sim} \chi^2_{q-p}.$$

WEEK 5 3rd to 8th October

Topic 3d: Residuals for Binomial Data and Neuroblastoma Example

RECALL: RESIDUALS IN LINEAR REGRESSION MODELS