## BIOS621 Session 4 - loglinear regression part 1

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#### Welcome and outline - session 4

- brief review of GLMs
- Motivating example for log-linear models
  - ► Poisson regression
- ▶ Checking model assumptions and fit: Residual Analysis
- Note on collinearity

Reading: Vittinghoff textbook chapter 8.1-8.3

## Learning Objectives

- Define log-linear models in GLM framework
- Identify situations that motivate use of log-linear models
- Assess model fit of log-linear models
- Define multi-collinearity

## Components of GLM

- ▶ Random component specifies the conditional distribution for the response variable - it doesn't have to be normal but can be any distribution that belongs to the "exponential" family of distributions
- Systematic component specifies linear function of predictors (linear predictor)
- ▶ Link [denoted by g(.)] specifies the relationship between the expected value of the random component and the systematic component, can be linear or nonlinear

## Linear Regression as GLM

The model:

$$y_i = E[y|x] + \epsilon_i = \beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + \dots + \beta_p x_{pi} + \epsilon_i$$

- ▶ **Random component** of  $y_i$  is normally distributed:  $\epsilon_i \stackrel{iid}{\sim} N(0, \sigma_{\epsilon}^2)$
- Systematic component (linear predictor):  $\beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + ... + \beta_n x_{ni}$
- ▶ **Link function** here is the *identity link*: g(E(y|x)) = E(y|x). We are modeling the mean directly, no transformation.

# Logistic Regression as GLM

► The model:

$$Logit(P(x)) = log\left(\frac{P(x)}{1 - P(x)}\right) = \beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + \dots + \beta_p x_{pi}$$

- ▶ Random component: *y<sub>i</sub>* follows a Binomial distribution (outcome is a binary variable)
- ► Systematic component: linear predictor

$$\beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + ... + \beta_p x_{pi}$$

▶ Link function: logit (log of the odds that the event occurs)

$$g(P(x)) = logit(P(x)) = log\left(\frac{P(x)}{1 - P(x)}\right)$$

$$P(x) = g^{-1} (\beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + \dots + \beta_p x_{pi})$$

### Additive vs. Multiplicative models

- Linear regression is an additive model
  - e.g. for two binary variables  $\beta_1 = 1.5$ ,  $\beta_2 = 1.5$ .
  - ▶ If  $x_1 = 1$  and  $x_2 = 1$ , this adds 3.0 to E(y|x)
- ► Logistic regression is a *multiplicative* model
  - If  $x_1 = 1$  and  $x_2 = 1$ , this adds 3.0 to  $log(\frac{P}{1-P})$
  - Odds-ratio  $\frac{P}{1-P}$  increases 20-fold: exp(1.5+1.5) or exp(1.5)\*exp(1.5)

## Motivating example for log-linear models

- ▶ Effectiveness of a new case-management program for depression
  - ► can the new treatment reduce the number of needed visits to the emergency room, compared to standard care?
- outcome: # of emergency room visits for each patient in the year following initial treatment
- predictors: race (white or nonwhite), treatment (treated or control), amount of alcohol consumption (numerical measure), drug use (numerical measure)

# Motivating example (cont'd)

- Statistical issues:
  - ▶ about 1/3 of observations are exactly 0 (did not return to the emergency room within the year)
  - highly nonnormal and cannot be transformed to be approximately normal
  - even  $log(y_i + 1)$  transformation will have a "lump" at zero
    - ightharpoonup over 1/2 the transformed data would have values of 0 or log(2)
  - a linear regression model would give negative predictions for some covariate combinations
  - some subjects die or cannot be followed up on for a whole year

# Motivating example (cont'd)

- ► A *multiplicative* model will allow us to make inference on *ratios* of mean emergency room usage
- ▶ Modeling log of the mean emergency usage ensures positive means, and does not suffer from log(0) problem
- ▶ Random component of GLM, or residuals (was  $\epsilon_i \stackrel{iid}{\sim} N(0, \sigma_\epsilon^2)$  for linear regression) may still not be normal, but we can choose from other distributions

# Motivating example: proposed model without time

$$log(E[Y_i]) = \beta_0 + \beta_1 RACE_i + \beta_2 TRT_i + \beta_3 ALCH_i + \beta_4 DRUG_i$$
 Or equivalently:

$$E[Y_i] = \exp(\beta_0 + \beta_1 RACE_i + \beta_2 TRT_i + \beta_3 ALCH_i + \beta_4 DRUG_i)$$

where  $E[Y_i]$  is the expected number of emergency room visits for patient i.

▶ Important note: Modeling  $log(E[Y_i])$  is not equivalent to modeling  $E(log(Y_i))$ 

# Motivating example: accounting for time of follow-up

Instead, model mean count per unit time:

$$log(E[Y_i]/t_i) = \beta_0 + \beta_1 RACE_i + \beta_2 TRT_i + \beta_3 ALCH_i + \beta_4 DRUG_i$$

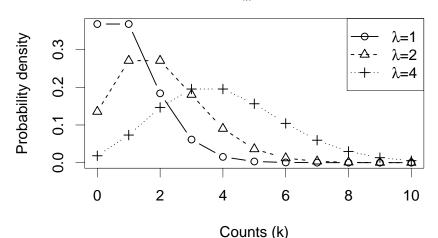
Or equivalently:

$$log(E[Y_i]) = \beta_0 + \beta_1 RACE_i + \beta_2 TRT_i + \beta_3 ALCH_i + \beta_4 DRUG_i + log(t_i)$$

 $\triangleright$   $log(t_i)$  is not a covariate, it is called an *offset* 

## Motivating example: Choice of Distribution

- ▶ Count data are often modeled as Poisson distributed:
  - $\blacktriangleright$  mean  $\lambda$  is greater than 0
  - ightharpoonup variance is also  $\lambda$
  - Probability density  $P(k,\lambda) = \frac{\lambda^k}{k!} e^{-\lambda}$



## Motivating example: the Poisson GLM

- Model the number of counts per unit time as Poisson-distributed
  - **>** so the expected number of counts per time is  $\lambda_i$

$$E[Y_i]/t_i = \lambda_i$$

$$log(E[Y_i]/t_i) = log(\lambda_i)$$

$$log(E[Y_i]) = log(\lambda_i) + log(t_i)$$

Recalling the log-linear model systematic component:

$$log(E[Y_i]) = \beta_0 + \beta_1 RACE_i + \beta_2 TRT_i + \beta_3 ALCH_i + \beta_4 DRUG_i + log(t_i)$$

# Motivating example: the Poisson GLM

Then the systematic part of the GLM is:

$$log(\lambda_i) = \beta_0 + \beta_1 RACE_i + \beta_2 TRT_i + \beta_3 ALCH_i + \beta_4 DRUG_i$$

Or alternatively:

$$\lambda_{i} = exp\left(\beta_{0} + \beta_{1}RACE_{i} + \beta_{2}TRT_{i} + \beta_{3}ALCH_{i} + \beta_{4}DRUG_{i}\right)$$

# Motivating example: interpretation of coefficients

- Suppose that  $\hat{\beta}_1 = -0.5$  in the fitted model, where  $RACE_i = 0$  for white and  $RACE_i = 1$  for non-white.
- ► The mean rate of emergency room visits per unit time for white relative to non-white, all else held equal, is estimated to be:

$$\frac{\exp(\beta_0 + 0 + \beta_2 \text{TRT}_i + \beta_3 \text{ALCH}_i + \beta_4 \text{DRUG}_i)}{\exp(\beta_0 - 0.5 + \beta_2 \text{TRT}_i + \beta_3 \text{ALCH}_i + \beta_4 \text{DRUG}_i)}$$

$$= \frac{e^{\beta_0} e^0 e^{\beta_2 \text{TRT}_i} e^{\beta_3 \text{ALCH}_i} e^{\beta_4 \text{DRUG}_i}}{e^{\beta_0} e^{-0.5} e^{\beta_2 \text{TRT}_i} e^{\beta_3 \text{ALCH}_i} e^{\beta_4 \text{DRUG}_i}}$$

$$= \frac{e^0}{e^{-0.5}}$$

$$= e^{0.5} \approx 1.65$$

# Motivating example: interpretation of coefficients

- ▶ If  $\hat{\beta}_1 = -0.5$  with whites as the reference group:
  - ▶ after adjustment for treatment group, alcohol and drug usage, whites tend to use the emergency room at a rate 1.65 times higher than non-whites.
  - equivalently, the average rate of usage for whites is 65% higher than that for non-whites
- ► Multiplicative rules apply for other coefficients as well, because they are exponentiated to estimate the mean rate.

#### Example by simulation

```
simdat <- data.frame(race=sample(c("white", "non-white"), size=10000, replace=TRUE))
simdat$race <- factor(simdat$race, levels=c("white", "non-white"))
simdat$y <- rpois(10000, lambda=ifelse(simdat$race=="white", exp(3.5), exp(3)))
fit <- glm(y - race, data=simdat, family=poisson("log"))
summary(fit)</pre>
```

```
##
## Call:
## glm(formula = v ~ race, family = poisson("log"), data = simdat)
##
## Deviance Residuals:
      Min
               10 Median
                                 30
                                         Max
## -3.5323 -0.7127 -0.0246 0.6616 3.5473
##
## Coefficients:
##
                Estimate Std. Error z value Pr(>|z|)
## (Intercept) 3.500139 0.002446 1431.0 <2e-16 ***
## racenon-white -0 498900 0 004003 -124 6 <2e-16 ***
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for poisson family taken to be 1)
##
##
      Null deviance: 26157 on 9999 degrees of freedom
## Residual deviance: 10111 on 9998 degrees of freedom
## ATC: 60885
##
## Number of Fisher Scoring iterations: 4
```

#### Inference on deviance residuals

- ▶ The difference in total deviance between two nested models is  $\chi^2$  distributed under  $H_0$  that the more complex model is no better at explaining the response.
  - ► The difference in deviance residuals is (26157 10111) = 16046, with a difference of 1 degrees of freedom.

The critical threshold for rejection at p=0.05 is:

```
qchisq(0.95, df=1)
```

## [1] 3.841459

So we reject  $H_0$ 

#### Inference on deviance residuals 2

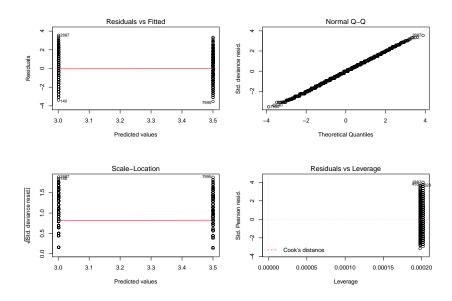
- lacktriangle Total residual deviance is  $\chi^2$  distributed if the model is correctly specified
  - What is the critical value for rejecting  $H_0$  at p < 0.05 with a  $\chi^2$  distribution of 9998 degrees of freedom?

```
qchisq(0.95, df=9998)
```

```
## [1] 10231.73
```

Here total residual deviance is 10111, so we do *not* exceed the threshold and do not reject  $H_0$  that the model is correctly specified.

### Example by simulation: Deviance Residuals Plots



## Example: Risky Drug Use Behavior

- Load the "needle\_sharing" dataset is available csv format
- Outcome is # times the drug user shared a syringe in the past month (shared\_syr)
- Predictors: sex, ethn, homeless

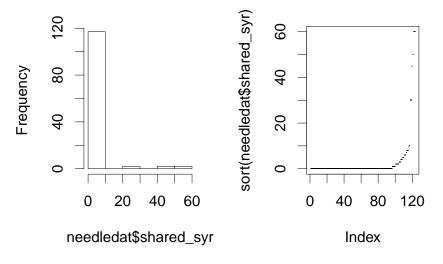
```
needledat = read.csv("needle_sharing.csv")
summary(needledat$shared_syr)

## Min. 1st Qu. Median Mean 3rd Qu. Max. NA's
## 0.000 0.000 0.000 2.976 0.000 60.000 5

var(needledat$shared_syr, na.rm=TRUE)
```

```
## [1] 106.5978
```

# Example: Risky Drug Use Behavior

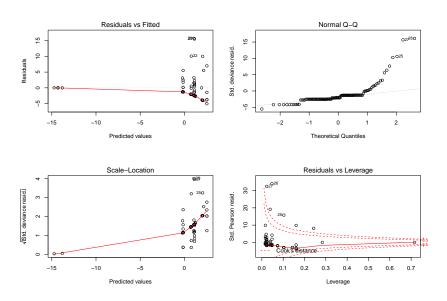


▶ There are a *lot* of zeros - Poisson model is not a good fit

# Risky Drug Use Behavior: fitting a Poisson model

```
##
## Call:
## glm(formula = shared_syr ~ sex + ethn + homeless, family = poisson(link = "log"),
      data = needledat)
##
##
## Deviance Residuals:
     Min
             10 Median
                           30
                                  Max
## -5.057 -2.506 -2.030 -1.279 15.721
##
## Coefficients:
##
                    Estimate Std. Error z value Pr(>|z|)
## (Intercept)
                   ## sexM
                   -0.92480 0.12133 -7.622 2.50e-14 ***
## sexTrans
                   -15.08655 773.78384 -0.019 0.9844
## ethnFilipino -14.52887 510.68253 -0.028 0.9773
## ethnHispanic
                   1.46454 0.16004 9.151 < 2e-16 ***
## ethnIndian
                   -14.10111 773.78385 -0.018 0.9855
## ethnIndian & White -15.02591 773.78384 -0.019 0.9845
## ethnWhite
                    0.06064 0.13348 0.454 0.6496
## ethnWhite & Hispa 0.86195 0.39872 2.162 0.0306 *
## homelessyes
                1.28543 0.12664 10.150 < 2e-16 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for poisson family taken to be 1)
##
      Null deviance: 1621.9 on 120 degrees of freedom
## Residual deviance: 1364.8 on 111 degrees of freedom
    (7 observations deleted due to missingness)
## ATC: 1483.8
##
## Number of Fisher Scoring iterations: 12
```

# Risky Drug Use Behavior: residuals plots



# Multicollinearity

- Multicollinearity exists when two or more of the independent variables in regression are moderately or highly correlated.
- Multicollinearity implies near-linear relationship among the predictors
- ► The presence of near-linear dependence dramatically impacts the ability to estimate regression coefficients
- High multicollinearity results in larger standard errors for regression coefficients
  - estimates of such regression coefficients will tend to be less stable over repeated sampling

### Concluding notes

- ► Inference from log-linear models is sensitive to the choice of link function (assumption on distribution of residuals)
- We will cover other options next week for when the Poisson model doesn't fit:
  - ▶ Variance proportional to mean, instead of equal
  - Negative Binomial
  - Zero Inflation