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Basque Center for Applied Mathematics

http://idaejin.github.io/bcam-courses/



GitHub: idaeiin

Introduction

- ► Response variable is a **count**
- ▶ e.g.: number of cases of a disease, Number of infected plants, firms going bankrupt etc ...
- When exploratory variables are categorical (i.e. you have a contingency table with counts in the cells), the models are usually called 'log-linear models'
- ► If they are numerical/continuous, convention is to call them **poisson** regression
- ► The Poisson distribution describes the count of the number of random events within a fixed interval of time or space with a known average rate.



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Introduction

- ▶ Poisson regression may also be appropriate for rate data, where the rate is a count of events occurring to a particular unit of observation, divided by some measure of that unit's **exposure**.
- ▶ e.g.: The number of deaths may be affected by the underlying population at risk, in demography death rates in geographic areas are modelled as the count of deaths divided by person-years, etc ...
- ▶ In Poisson regression this is handled as an offset.
- ▶ Then, we include the exposure (*exp*) in the denominator and form a rate such as:

$$Y/exp = rate \Rightarrow Y = rate \times exp$$



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D.-J. Lee (BCAM)

Introduction

► Linear regression methods (constant variance, normal errors) are not appropriate for count data for 5 main reasons:



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Introduction

- ► Linear regression methods (constant variance, normal errors) are not appropriate for count data for 5 main reasons:
  - 1. the linear model might lead to the prediction of negative counts
  - 2. the variance of the response variable is likely to increase with the mean
  - 3. the errors will not be normally distributed
  - 4. zeros are difficult to handle in transformations
  - 5. some distributions (e.g. log-normal or gamma) do not allow zeros



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#### Poisson distribution

- The Poisson distribution is widely used for the description of count data that refer to cases where we know how many times something happened, but we have no way of knowing how many times it did not happen.
- Recall that with the binomial distribution we know how many times something did not happen as well as how often it did happen.



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#### Poisson distribution

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- ► Recall that with the binomial distribution we know how many times something did not happen as well as how often it did happen.

#### Definition

lacktriangledown a discrete random variable Y is ditributed as Poisson with parameter  $\mu>0$ , if, for k=0,1,2,..., the probability mass function of Y is given by

$$\Pr(Y = k) = \frac{\mu^k e^{-\mu}}{k!}$$

 $ightharpoonup \mathbb{E}(Y) = \mathbb{V}ar(Y) = \mu$ 

D.-J. Lee (BCAM) Intro to GLM's with R GitHub: idaejin

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Components of a GLM

- ► The components of a GLM for a count response are:
  - 1. Random Component: Poisson distribution and  $\mathbb{E}(Y) = \mu$
  - 2. Systematic component: predictors/explanatory variables X
  - 3. Link function:
    - ▶ Identity link: which would give us  $\mu = \beta_0 + \beta_1 x_1 + ... + \beta_k x_k$  (but then we can get  $\mu < 0$ )
    - ▶ Log link: is the most common and canonical link

$$\log(\mu) = \beta_0 + \beta_1 x_1 + \dots + \beta_k x_k$$

• Since the  $\log(\mu)$  is a linear function of the X's and  $\mu$  is a multiplicative function of X:

$$\mu = e^{\beta_0 + \beta_1 x_1 + \dots + \beta_k x_k}$$
$$= e^{\beta_0} e^{\beta_1 x_1} \dots e^{\beta_p x_k} = \exp(\mathbf{X}\boldsymbol{\beta})$$



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D.-J. Lee (BCAM)

Components of a GLM

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log link and interpretation

$$\log(\mu) = \beta_0 + \beta_1 x_1 + \dots + \beta_k x_k$$

▶ What does it mean for  $\mu$ ? How do you interpret  $\beta$ ?



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log link and interpretation

$$\log(\mu) = \beta_0 + \beta_1 x_1 + \dots + \beta_k x_k$$

- ▶ What does it mean for  $\mu$ ? How do you interpret  $\beta$ ?
- ▶ The  $\log$  is a transformation of the mean such that  $\log(\mu)$  has a linear relationship with the predictors,  $e_i^\beta$  represents a multiplier effect of the  $i^{\rm th}$  predictor on the mean, such that a 1-unit increase on  $X_i$  has a multiplicative effect of  $e^{\beta_i}$  on  $\mu$ .



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- ▶ The  $\log$  is a transformation of the mean such that  $\log(\mu)$  has a linear relationship with the predictors,  $e_i^\beta$  represents a multiplier effect of the  $i^{\text{th}}$  predictor on the mean, such that a 1—unit increase on  $X_i$  has a multiplicative effect of  $e^{\beta_i}$  on  $\mu$ .
- ▶ Suppose of a single predictor x and 2 values of x (i.e.  $x_1$  and  $x_2$ ), with a difference of 1, i.e.  $x_1=10$  and  $x_1=11$ , where  $\mu_1=\mathbb{E}(Y|x=10)$  and  $\mu_2=\mathbb{E}(Y|x=11)$ 
  - ▶ If  $\beta = 0$ , then  $e^0 = 1$  and  $\mu_2$  is the same as  $\mu_1$ . That is  $\mu$  is not related to x.
  - ▶ If  $\beta > 0$ ,  $e^{\beta} > 1$  and  $\mu_2$  is  $e^{\beta}$  times larger than  $\mu_1$ .
  - ▶ If  $\beta < 0$ ,  $e^{\beta} < 1$  and  $\mu_2$  is  $e^{\beta}$  times smaller than  $\mu_1$ .



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Estimation, Deviances, Hypothesis testing, Goodness-of-Fit and residuals

- ► Once we are in a GLM setting with a log link, the parameters are estimated by Maximum Likelihood and Iterative Re-weighted Least Squares.
- ► The **Deviance** (discrepancy measure between observed and fitted values) for Poisson responses takes the form:

$$D = 2\sum \{y_i \log \left(\frac{y_i}{\hat{\mu}_i}\right) - (y_i - \hat{\mu}_i)\}$$

for large samples  $D \sim \chi^2_{n-p}$ .

- ▶ Goodness-of-Fit measure is the Pearson's chi-squared statistic  $\chi_p^2 = \sum \frac{(y_i \mu_i)^2}{\hat{\mu}_i}$
- ► Likelihood ratio tests can easily be constructed in terms of deviances, just as we did in logistic regression models.
- ▶ We will use standardized residuals  $r_i = \frac{y_i \hat{\mu}_i}{\sqrt{\hat{\mu}_i}} \sim \mathcal{N}(0,1)$

(bcam)

Case study: Agricultural experiment of species richness

▶ A longterm agricultural experiment had 90 grassland plots, each  $25m \times 25m$ , differing in biomass, soil pH and species richness (the count of species in the whole plot).



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- ▶ It is well known that species richness declines with increasing biomass, but the question addressed here is whether the slope of that relationship differs with soil pH. The plots were classified according to a 3-level factor as high, medium or low pH with 30 plots in each level.



D.-J. Lee (BCAM)

Case study: Agricultural experiment of species richness

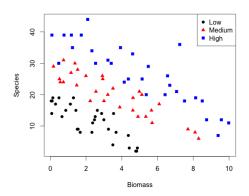
- ▶ A longterm agricultural experiment had 90 grassland plots, each  $25m \times 25m$ , differing in biomass, soil pH and species richness (the count of species in the whole plot).
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#### See Poisreg.R

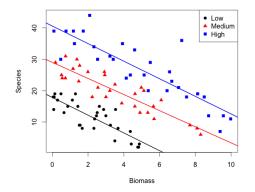
- > species<-read.table("data/species.txt",header=TRUE)
- > names(species)



We can plot the data for each level of soil pH and fit a lm



We can plot the data for each level of soil pH and fit a lm



There is a clear difference in mean species richness with declining soil pH, but there is little evidence of any substantial difference in the slope of the relationship between species richness and biomass on soils of differing pH.

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Case study: Agricultural experiment of species richness

▶ We start fitting each predictor separately, are they significant?



Case study: Agricultural experiment of species richness

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```
> m0=glm(Species~Biomass,family=poisson,data=species)
```

- > anova(m0,test="Chisq")
- > m1=glm(Species~pH,family=poisson,data=species)
- > anova(m1,test="Chisq")



Case study: Agricultural experiment of species richness

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```

► Include the interaction and test its significance



Case study: Agricultural experiment of species richness

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- ▶ Include the interaction and test its significance The effect of biomass on species richness depends on soil pH, and the strength of the relationship varies with soil pH



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Case study: Agricultural experiment of species richness

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> m1=glm(Species~pH,family=poisson,data=species)
> anova(m1,test="Chisq")
```

▶ Include the interaction and test its significance The effect of biomass on species richness depends on soil pH, and the strength of the relationship varies with soil pH

```
> m2 <- glm(Species~Biomass+pH,family=poisson,data=species)
> m3 <- glm(Species~Biomass*pH,family=poisson,data=species)
> anova(m2,m3,test="Chisq")
```

Hence assume that the slopes were the same is completely wrong: in fact, the slopes are very significantly different (p < 0.00035).

```
> summary(m3)
```

```
Coefficients:
```

Biomass:pHHigh 0.15503 0.04003 3.873 0.000108 \*\*\*
--Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for poisson family taken to be 1)

Null deviance: 452.346 on 89 degrees of freedom Residual deviance: 83.201 on 84 degrees of freedom

AIC: 514.39

Number of Fisher Scoring iterations: 4



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Case study: Agricultural experiment of species richness (cont.)

▶ The fitted model is

$$\begin{split} \log(\texttt{Species}) &= 2.95 - 0.262 \times \texttt{Biomass} + 0.484 \texttt{pH}_{medium} + 0.815 \times \texttt{pH}_{high} \\ &+ 0.123 \times \texttt{pH}_{medium} \times \texttt{Biomass} + 0.155 \times \texttt{pH}_{high} \times \texttt{Biomass} \end{split}$$



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Case study: Agricultural experiment of species richness (cont.)

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- Interpretation of the parameters:
  - A 1-unit change in the Biomass implies a number of species of exp(−0,262) = 0,769 times less in soil with low pH.



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  - A 1-unit change in the Biomass implies a number of species of exp(−0,262 + 0,123) = 0,870 times less in soil with medium pH.



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Case study: Agricultural experiment of species richness (cont.)

► The fitted model is

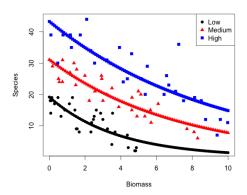
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- Interpretation of the parameters:
  - A 1-unit change in the Biomass implies a number of species of exp(−0,262) = 0,769 times less in soil with low pH.
  - A 1-unit change in the Biomass implies a number of species of  $\exp(-0.262 + 0.123) = 0.870$  times less in soil with **medium** pH.
  - A 1-unit change in the Biomass implies a number of species of exp(−0,262 + 0,155) = 0,898 times less in soil with high pH.



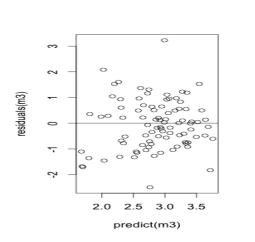
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We can plot the data for each level of soil pH



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And check the fitted VS residuals plot and the QQ-plot of the residuals



# Normal Q-Q Plot 3 2

Theoretical Quantiles

Overdispersion

- ▶ In Poisson regression, it is assumed that  $Var(Y) = \mu$
- ► Sometimes the data do not hold this assumption, this yields to wrong estimation of the standard errors in the parameters.
- ▶ Overdispersion is present if  $Var(Y) > \mu$
- A simple option to account for this phenomena is to include a **dispersion** parameter  $\phi$ , such that  $\mathbb{V}\mathrm{ar}(Y) = \phi \mu$ , then when  $\phi = 1$  we are in the Poisson case.
- $\blacktriangleright$  The dispersion parameter  $\phi$  can be estimated as

$$\hat{\phi} = \frac{\mathsf{Deviance}}{n-p}$$

▶ In our example, we had that Deviance = 83,2 and n-p=84, then  $\phi\approx 1$ .



Overdispersion (cont.)

▶ In R, we can specify the family=quasipoisson

```
> m4 <- glm(Species~Biomass*pH,family=quasipoisson,data=species)
```



Overdispersion (cont.)

▶ In R, we can specify the family=quasipoisson

```
> m4 <- glm(Species~Biomass*pH,family=quasipoisson,data=species)
```

► Or use the function dispersiontest in library(AER)

```
> library(AER)
> dispersiontest(m3)
```

Overdispersion test

data: m3

z = -0.38285, p-value = 0.6491

alternative hypothesis: true dispersion is greater than 1

sample estimates:

dispersion

0.9369549

► The test is

$$H_0: \phi^* = 1$$
  
 $H_1: \phi^* \neq 1$ 

$$H_1: \phi^* \neq$$



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Other alternatives for overdispersion

- One common cause of overdispersion is excess zeros, which in turn are generated by an additional data generating process. In this situation, zero-inflated Poisson model should be considered.
- ► Negative Binomial regression: it can be considered as a generalization of Poisson regression since it has the same mean structure as Poisson regression and it has an extra parameter to model the overdispersion. E.g.: glm.nb function in library (MASS)



for incidence rates

► GLM's for rates

$$g(\mu) = \beta_0 + \beta_1 x_1 + \beta_2 x_2 + \ldots + \beta_k x_k$$

- ▶ Random component: Response Y has a Poisson distribution, and t is index of the time or space; more specifically the expected value of rate Y/t, is  $\mathbb{E}(Y/t) = \mu$  that is  $\mathbb{E}(Y) = \mu t$ .
- ▶ Systematic component: Any set of  $X = (x_1, x_2, ..., x_k)$  can be explanatory variables. For now let's focus on a single variable x.
- ▶ Link function: Log of rate: log(Y/t)



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for incidence rates (cont.)

Poisson loglinear regression model for the expected rate of the occurrence of event is:

$$log(\mu/t) = \alpha + \beta x$$

This can be rearranged to:

$$\log(\mu) - \log(t) = \alpha + \beta x$$
  
$$\log(\mu) = \alpha + \beta x + \log(t)$$

- ▶ The term  $\log(t)$  is referred to as an **offset**. It is an adjustment term and a group of observations may have the same offset, or each individual may have a different value of t.
- $ightharpoonup \log(t)$  is an observation and it will change the value of estimated counts:

$$\mu = \exp(\alpha + \beta x + \log(t)) = t \times \exp(\alpha)\exp(\beta x)$$

This means that mean count is proportional to t.

Note that the interpretation of parameter estimates,  $\alpha$  and  $\beta$  will stay the same as for the model of counts; you just need to multiply the expected counts by t.

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for incidence rates (cont.)

- ▶ In poisson regression we also have relative risk (RR) interpretations, i.e. the estimated effect of an explanatory variable is multiplicative on the rate, and thus leads to a risk ratio or relative risk.
- ► This is similar to logistic regression, but then the effect of an explanatory variable is multiplicative on the odds and thus leads to an odds ratio.



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for incidence rates (cont.)

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- ► This is similar to logistic regression, but then the effect of an explanatory variable is multiplicative on the odds and thus leads to an odds ratio.
- ▶ Let  $\lambda_0$  be the incidence rate when  $x_0$  and  $\lambda_1$  when  $X = x_0 + 1$ , then

$$\log\left(\frac{\lambda_1}{\lambda_0}\right) = \log(\lambda_1) - \log(\lambda_0) = \beta$$

$$RR = \frac{\lambda_1}{\lambda_0} = \exp(\beta)$$

i.e.  $\exp(\beta)$  is the relative risk between the population with  $X=x_0$  and  $X=x_1$ , and  $\exp(\alpha)$  is the incidence rate when X=0



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Case study: Lung cancer incidence in Denmark 1968-1971

- ▶ This data set contains counts of incident lung cancer cases and population size in four neighbouring Danish cities by age group.
- ► The variables are:

```
city a factor with levels Fredericia, Horsens, Kolding, and Vejl age a factor with levels 40-54, 55-59, 60-64, 65-69, 70-74, and 75+. pop the number of inhabitants. cases the number of lung cancer cases.
```



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```

► How does the expected number of lung cancer counts vary by age?

#### See Poisreg.r

> lung <- read.table("data/lung.txt",header=TRUE)</pre>



Case study: Lung cancer incidence in Denmark 1968-1971

#### See Poisreg.r

> head(lung)

```
city age pop cases
1 Fredericia 40-54 3059 11
2 Horsens 40-54 2879 13
3 Kolding 40-54 3142 4
4 Vejle 40-54 2520 5
5 Fredericia 55-59 800 11
6 Horsens 55-59 1083 6
```

The incidence rate is denoted by  $\lambda = Y/E$ , then for people aged 40-54 years, in Fredericia, Y=11 and E=3059, and then

$$\lambda \ = \ \frac{11}{3059} = 0.003595946$$
 or  $3.595946$  for each 1000 inhab.



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Case study: Lung cancer incidence in Denmark 1968-1971

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▶ The incidence rate is denoted by  $\lambda = Y/E$ , then for people aged 40-54 years, in Fredericia, Y = 11 and E = 3059, and then

$$\lambda \quad = \quad \frac{11}{3059} = 0.003595946$$
 or  $3.595946$  for each 1000 inhab.

> boxplot(cases~age,data=lung,col="bisque")



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Case study: Lung cancer incidence in Denmark 1968-1971

▶ We start considering a model with age as covariate

$$\begin{split} \log(\lambda_i) &= \beta_0 + \beta_1 I(\mathsf{Age55-59}_i) + \beta_2 I(\mathsf{Age60-64}_i) + \\ &+ \beta_3 I(\mathsf{Age65-69}_i) + \beta_4 I(\mathsf{Age70-74}_i) + \beta_5 I(\mathsf{Age}{>}74_i) \end{split}$$

where  $I(\cdot)$  is a indicator (1 if TRUE, 0 otherwise) for each range of age, with Age40-45 is used as baseline.



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Case study: Lung cancer incidence in Denmark 1968–1971

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where  $I(\cdot)$  is a indicator (1 if TRUE, 0 otherwise) for each range of age, with Age40-45 is used as baseline.

> lungmod1 <- glm(cases ~ age, family=poisson, data=lung)</pre>



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```
> summary(lungmod1)
```

```
Call:
```

glm(formula = cases ~ age, family = poisson, data = lung)

#### Deviance Residuals:

Min 1Q Median 3Q Max -2.4661 -0.4488 -0.1532 0.7773 1.5242

#### Coefficients:

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

(Intercept) 2.11021 0.17408 12.122 <2e-16 \*\*\*
age55-59 -0.03077 0.24810 -0.124 0.901
age60-64 0.26469 0.23143 1.144 0.253
age65-69 0.31015 0.22918 1.353 0.176
age70-74 0.19237 0.23516 0.818 0.413
age75+ -0.06252 0.25012 -0.250 0.803

Signif. codes: (

Case study: Lung cancer incidence in Denmark 1968-1971

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$$\begin{split} \log(\lambda_i) &= 2{,}11021 - 0{,}03077 \times I(\mathsf{Age55-59}_i) + 0{,}26469 \times I(\mathsf{Age60-64}_i) + \\ &+ 0{,}31015 \times I(\mathsf{Age65-69}_i) + 0{,}19237(\mathsf{Age70-74}_i) - 0{,}06252 \times I(\mathsf{Age}{>}74_i) \end{split}$$

#### ► Interpretation:

- $\exp(2,11021) = 8,24$  is the expected count of cancer cases among individuals aged 40-54
- $\exp(2,11021-0,03077)=8,00$  is the expected count of cancer cases among individuals aged 55-59
- $\exp(-0.0377) = 0.97$  is the ratio of the expected counts comparing the 55-59 aged group to the baseline group of age 40-54.  $\exp(\hat{\beta}_1)$  is also the relative rate.



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Case study: Lung cancer incidence in Denmark 1968-1971

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- $\exp(2,11021) = 8,24$  is the expected count of cancer cases among individuals aged 40-54
- $\exp(2,11021-0,03077)=8,00$  is the expected count of cancer cases among individuals aged 55-59
- $\exp(-0.0377) = 0.97$  is the ratio of the expected counts comparing the 55-59 aged group to the baseline group of age 40-54.  $\exp(\hat{\beta}_1)$  is also the relative rate.

> lungmod1 <- glm(cases ~ age, family=poisson, data=lung)</pre>

D.-J. Lee (BCAM) Intro to GLM's with R GitHub: idaejin

Case study: Lung cancer incidence in Denmark 1968–1971

If we calculate the  ${\bf Cl}$ 's for all ages we find that all contain 0, is there any association between cancer and age?

```
> confint(lungmod1)
```

```
2.5 % 97.5 % (Intercept) 1.7484571 2.4330407 age55-59 -0.5200577 0.4572927 age60-64 -0.1863226 0.7247913 age65-69 -0.1357438 0.7664682 age70-74 -0.2671849 0.6587654 age75+ -0.5564824 0.4289194
```



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Case study: Lung cancer incidence in Denmark 1968-1971

#### We can perform a Likelihood Ratio Test:

```
> lungmod0 <- glm(cases ~ 1, family=poisson, data=lung)
> anova(lungmod0,lungmod1,test="Chisq")
```

#### Analysis of Deviance Table

```
Model 1: cases ~ 1
Model 2: cases ~ age
   Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1 23 27.704
2 18 22.756 5 4.9478 0.4223
```

and hence, we do not reject the hyphotesis of all  $\beta_i = 0$ , for i = 1, ..., 5.



Case study: Lung cancer incidence in Denmark 1968-1971

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- ► Can we improve the analysis?



Case study: Lung cancer incidence in Denmark 1968-1971

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- Each city and age group has a different population size.
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Case study: Lung cancer incidence in Denmark 1968–1971

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- ▶ It may be of more interest to know the rate per person, per 4 period of observation. We are interested in

$$r_i = rac{\lambda_i}{\mathsf{Pop}_i} = rac{\mathbb{E}(\mathsf{count}_i)}{\mathsf{Pop}_i}$$

and model it by a log-linear model



D.-J. Lee (BCAM)

Case study: Lung cancer incidence in Denmark 1968–1971

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► Then, our model is

$$Y_i \sim \mathcal{P} \mathsf{ois}(\lambda_i) = \mathcal{P} \mathsf{ois}(r_i \times \mathsf{Pop}_i)$$



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Case study: Lung cancer incidence in Denmark 1968-1971

On a log-scale, our model is:

$$\begin{split} \log\left(\frac{\lambda_i}{\mathsf{Pop}_i}\right) &= \beta_0 + \beta_1 I(\mathsf{Age55-59}_i) + \beta_2 I(\mathsf{Age60-64}_i) + \\ &+ \beta_3 I(\mathsf{Age65-69}_i) + \beta_4 I(\mathsf{Age70-74}_i) + \beta_5 I(\mathsf{Age}{>}74_i) \end{split}$$

Exponentiating we have:

$$\begin{split} \left(\frac{\lambda_i}{\mathsf{Pop}_i}\right) &= \exp\left(\beta_0 + \beta_1 I(\mathsf{Age55\text{-}59}_i) + \beta_2 I(\mathsf{Age60\text{-}64}_i) + \\ &+ \beta_3 I(\mathsf{Age65\text{-}69}_i) + \beta_4 I(\mathsf{Age70\text{-}74}_i) + \beta_5 I(\mathsf{Age}\text{>}74_i)) \end{split}$$

- ▶ All counts are restricted to the same period of 1968-1971, the values of  $\lambda_i$  are rates 'per 4-years'
- ► To obtain an easier interpretation of the rates we can divide by 4 to get rate per person-year and multiply by 10,000 to get a rate per 10,000 person-years, i.e. divide by 2500

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Case study: Lung cancer incidence in Denmark 1968-1971

This can be easily done by

```
> lungmod2 <- glm(cases ~ age + offset(log(pop/2500)),
+ family=poisson, data=lung)</pre>
```

- ▶ log(pop/2500) is the offset
- ▶ The offset accounts for the population size, which could vary by age, region, etc ...
- It gives a convenient way to model rates per person-years, instead of modeling the raw counts.



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Case study: Lung cancer incidence in Denmark 1968-1971

Null deviance: 129.908 on 23 degrees of freedom Residual deviance: 28.307 on 18 degrees of freedom

AIC: 136.69



Case study: Lung cancer incidence in Denmark 1968-1971

#### Confidence intervals for the parameters can also be obtained as:

```
> confint(lungmod2)
```

```
2.5 % 97.5 % (Intercept) 1.6000371 2.284621 age55-59 0.5930558 1.570406 age60-64 1.0506603 1.961774 age65-69 1.3043879 2.206600 age70-74 1.3876652 2.313616 age75+ 0.9143186 1.899720
```



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Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

- ► The inclusion of the offset, implies that the interpretation of the coefficients should be done in terms of  $log(\lambda_i) offset_i$
- ▶ In our example,  $\lambda_i$  is the expected number of cases observed in a particular age group, within a 4 year period of time.
- ► Hence in our case, with an offset of log(pop/2500), we should think of the outcome as log rate per 10,000 person years.
- ► Then:
  - $\triangleright$   $\beta_0$  is the log rate of cancer cases per 10,000 person years in the age group of 40-54 (baseline)



Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

- ▶ The inclusion of the offset, implies that the interpretation of the coefficients should be done in terms of  $\log(\lambda_i)$  – offset<sub>i</sub>
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  - $\triangleright$   $\beta_1$  is the log relative rate of cancer cases per 10,000 person years comparing the age group of 50-59 to the baseline age group 40-54



Intro to GLM's with R

Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

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- ► Then:
  - $\beta_0$  is the log rate of cancer cases per 10,000 person years in the age group of 40-54 (baseline)
  - $ightharpoonup eta_1$  is the log relative rate of cancer cases per 10,000 person years comparing the age group of 50-59 to the baseline age group 40-54
  - $ightharpoonup eta_2$  is the log relative rate of cancer cases per 10,000 person years comparing the age group of 60-64 to the baseline age group 40-54



Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

- ▶ Note that, including the offset, the regression coefficients are significant in the expected counts per year at age group compared to the baseline group.
- ► Now, with the offset, we are looking for differences in the expected counts per person-year, across age groups.



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Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

- ▶ Note that, including the offset, the regression coefficients are significant in the expected counts per year at age group compared to the baseline group.
- Now, with the offset, we are looking for differences in the expected counts per person-year, across age groups.
- ► Likelihood Ratio Test to test the effect of age

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Case study: Lung cancer incidence in Denmark 1968-1971 (cont.)

What are the difference between model with and without the offset?

Without offset	With offset
Do not reject H <sub>0</sub>	Reject H <sub>0</sub>
Expected cases across the age groups*	Expected cases per person year,
	across the age groups**

- There is no difference because the lower population Age75+ might be counterbalanced by high rate of cases with increasing age
- \*\* The offset let us compare the rate of cancer among those who are alive, i.e. taking into account the number of people within cohort of interest.



Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

► Given the model including the offset what can we say about the rate of lung cancer in Denmark?



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Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

- Given the model including the offset what can we say about the rate of lung cancer in Denmark?
- ▶ Predicted log expected count of cancer from 1968 1971 among 40 54 year old group of age:

$$\begin{split} \log(\lambda_i) &= \log\left(\frac{\text{Population Age 40-54}}{2500}\right) + \hat{\beta}_0 \\ &= \log\left(\frac{11600}{2500}\right) + 1,96 = 3,49 \end{split}$$

- ▶ where predicted rate of cancer per 10,000 person years among 40-54 year olds is  $\log(\lambda_i) = \hat{\beta}_0 = 1.96$
- ▶ Taking exponentials we have the predicted expected count of cancer cases 1968-1971 among age 40-54

$$\lambda_i = \exp(3.49) = 32.9$$

per 10,000 person years among 40-54, is  $\exp(\hat{\beta}_0)=7{,}11$ 



Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

per 10,000 person years among 40-54, is  $\exp(\hat{\beta}_0) = 7.11$ Then, based on this model, the prediction of cases of lung cancer per 10,0000 people aged 40-54 year old in Denmark, per year is 7,11.



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Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

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Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

per 10,000 person years among 40-54, is  $\exp(\hat{\beta}_0)=7.11$ Then, based on this model, the prediction of cases of lung cancer per 10,0000 people aged 40-54 year old in Denmark, per year is 7,11.And among 55-59 age group?

▶ Log expected count is

$$\begin{split} \log(\lambda_i) &= \log\left(\frac{\text{Population Age 55-59}}{2500}\right) + \hat{\beta}_0 + \hat{\beta}_1 \\ &= \log\left(\frac{3811}{2500}\right) + 1,96 + 1,08 = 3,46 \end{split}$$

and the predicted log rate of cancer per 10,000 person years among 55-59 years old  $\hat{\beta}_0+\hat{\beta}_1=1,96+1,08=3,04$ 



Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

▶ What is  $\exp(\hat{\beta}_1)$ ?



Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

- ▶ What is  $\exp(\hat{\beta}_1)$ ?
- $\exp(\hat{\beta}_1) = \exp(1,08) = 2,94$  is the relative rate of cancer cases per 10,000 person years comparing 55-59 and 40-54 groups, i.e.

$$\frac{\exp(\hat{\beta}_0 + \hat{\beta}_1)}{\exp(\hat{\beta}_0)} = \frac{20,9}{7,09} = 2,94$$



Case study: Lung cancer incidence in Denmark 1968–1971 (cont.)

- ▶ What is  $\exp(\hat{\beta}_1)$ ?
- $\exp(\hat{\beta}_1) = \exp(1,08) = 2,94$  is the relative rate of cancer cases per 10,000 person years comparing 55-59 and 40-54 groups, i.e.

$$\frac{\exp(\hat{\beta}_0 + \hat{\beta}_1)}{\exp(\hat{\beta}_0)} = \frac{20,9}{7,09} = 2,94$$

> exp(coef(lungmod2))

(Intercept) age55-59 age60-64 age65-69 age70-74 age75+ 7.112069 2.951584 4.489204 5.756252 6.342177 4.088919

There is a increasing trend in relative rates compared to the baseline age group 40-54 except for Age75+



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Some conclusions

- ▶ The offset allows for including the change the denominator or units of a rate
- ► When considering poisson regression on incidence rates, the model without an offset does not make much sense, and it fails to hold the Poisson assumptions
- ► We should try to use an offset if we suspect that the underlying population sizes differ for each of the observed counts

