Lecture 11: Introduction to Generalized Linear Models

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Outline

- 1. Introduction (motivation and history)
- 2. Review ordinary linear regression
- 3. Components of a GLM
- 4. Natural Exponential family

		Response Variable	
Explanatory Variables	Binary	Nominal	Continuous
Binary	2×2 table	Contingency tables	t-tests
	logistic regression	log-linear models	
Nominal	Logistic regression	Contingency tables	ANOVA
	Log-linear models	log-linear models	
Continuous	Dose-response models	It depends	Multiple regression
	logistic regression		
Some Continuous	Logistic regression	It depends	ANCOVA
and some categorical			Multiple regression

Note, in general, most common analyses can be approached from a "modelling" approach. Some such as the log-linear and logistic are topics for this class. Why do we want to "model" data?

- The structural form of the model describes the patterns of interactions or associations in data.
- Inference for the model parameters provides a way to evaluate which explanatory variable(s) are related to the response variable(s) while statistically controlling for the other variables.
- Estimated model parameters provide measures of the strength and importance of effects.
- A model's predicted values "smooth" the data That is, they provide good estimates of the mean of the response variable.
- Modeling enables use to examine general extensions to the methods we have studied thus far.

Suppose you have a continuous response variable (Y) and continuous and/or discrete explanatory variables (X's).

You want to model the responses as a function of the explanatory variables (covariates). For example,

$$Y_i = \beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + e_i$$

where

- 1. Y_i is the response for the i^{th} subject
- 2. $\vec{\beta} = (\beta_0, \beta_1, \beta_2)'$ is a (column) vector of constants (or parameters) that describe the shape of the regression "line" (line, curve, etc)
- 3. $\vec{X}_i = (1, x_{1i}, x_{2i})$ is the (row) vector of explanatory variables for the i^{th} subject.
- 4. e_i is the random error assumed to be distributed as $N(0, \sigma^2)$

In general, you can view the previous regression model as,

$$Y = E(Y) + \epsilon$$

Where

$$E(Y) = \beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i}$$

or in more general terms
$$= X_{n \times p} \beta_{p \times 1}$$

Thus, E(Y) is the $n \times 1$ vector of expectations.

Note,

$$X = \begin{bmatrix} X_1 \\ X_2 \\ \dots \\ X_n \end{bmatrix} = \begin{bmatrix} x_{11}, x_{12}, \dots, x_{1p} \\ x_{21}, x_{22}, \dots, x_{2p} \\ \dots \\ x_{n1}, x_{n2}, \dots, x_{np} \end{bmatrix}$$

is called the design matrix.

The analysis of continuous data has relied heavily on the linear model presented. These reflect just a few applications of the linear model.

- 1. Simple linear regression
- 2. Multiple regression
- 3. ANOVA
- 4. ANCOVA

The least squares estimator for β is

$$\tilde{\beta} = (X'X)^{-1}X'Y$$

The predicted value of Y (denoted as \hat{Y}) is

 $\hat{Y} = X\tilde{\beta}$

Diagnostic of the regression fit can be accomplished with the Hat Matrix

 $H = X(X'X)^{-1}X'$

As we develop our 'generalized' approach, you will notice many similarities.

- 1. For OLS, we are dependent on the distribution of Y being normal.
- 2. For categorical data (by definition), the normality assumption is rarely feasible.
- 3. We may also be interested in other relations of the $X\beta$ with Y. Other mapping functions that ensure the range of Y remains valid is one of the key justifications.

In terms of a GLM, we have three components related to these limitations.

Three Components of a GLM

There are 3 components of a generalized linear model (or GLM)

- 1. Random Component (the outcome)
- 2. Systematic Component (the design matrix multiplied by the parameter vector)
- 3. Link function (the function, $g(\cdot)$ that "links" the systematic component to the random component)

Nelder & Wedderburn (1972) are generally given credit for unifying a broad class (of existing) models into the GLM definitions.

They showed that provided the random component was part of the 'Exponential Class', the MLEs for all of the models could be obtained using the same algorithm.

Random Component

- The random component of a GLM consists of a response variable Y with the independent observations (y_1, y_2, \ldots, y_n) .
- For a GLM, Y needs to have a distribution in the natural exponential family.
- Recall from theory, an exponential class distribution is of the form

$$f(y_i; \theta_i) = a(\theta_i)b(y_i)exp[y_iQ(\theta_i)]$$

This, in terms of common language, is

- $a(\theta_i)$ is a function only dependent on the unknown parameter
- $b(y_i)$ is a function of the observed sample
- $Q(\theta_i)$ is a function only dependent on the unknown parameter

Easy Example Exponential Class Variable

Suppose,

 $Y \sim \mathsf{Poisson}(\lambda)$

Then,

$$f(y,\lambda) = \frac{e^{-\lambda}\lambda^y}{y!}$$

$$= e^{-\lambda} (\frac{1}{y!}) e^{y \log \lambda}$$

Here $\theta = \lambda$, $a(\theta) = a(\lambda) = e^{-\lambda}$, b(y) = 1/y! and $Q(\pi) = \log \lambda$ Thus,

A Poisson random variable is of the exponential class variable.

Slightly more complicated example

Suppose,

$$Y_i \sim Bern(\pi$$

where $P(Y_i = 1) = \pi$ and $P(Y_i = 0) = 1 - \pi$

Then,

$$f(y_i;\pi) = \pi^{y_i} (1-\pi)^{1-y_i}$$

$$= \frac{\pi^{y_i}}{(1-\pi)^{y_i}(1-\pi)^{-1}}$$

$$= (1-\pi)(\frac{\pi}{1-\pi})^{y_i}$$

$$= (1-\pi)(1)e^{(y_i \log \frac{\pi}{1-\pi})}$$

Here $\theta = \pi$, $a(\theta) = a(\pi) = (1 - \pi)$, b(y) = 1 and $Q(\pi) = \log(\pi/(1 - \pi))$

Thus, a Bernoulli random variable is a member of the exponential class.

Exponential Class Variables in General

In general,

- The majority of distributions we will be interested in are exponential class variables
- This includes the more common examples of
 - 1. Normal
 - 2. Poisson
 - 3. Binomial
 - 4. Multinomial
- It also includes the less common examples of
 - 1. Gamma
 - 2. Negative Binomial

Examples

- 1. Dichotomous (binary) with a fixed number of trials
 - MI / No MI
 - Success/Failure
 - Correct/Incorrect
 - Agree/Disagree

These responses have a **Bernoulli** distribution.

- 2. Counts (including cells in a contingency table)
 - Number of babies born at MUSC daily
 - Number of car wrecks per year in Charleston County

These responses have a **Poisson** distribution.

Although, many distributions are members of the Exponential Class,

For the most part, we will focus on the

- 1. Binomial
- 2. Poisson

distributions.

However, the approach we will discuss works equally well for all exponential class distributions.

Systematic Component

Denote a new vector $(\eta_1, \eta_2, \ldots, \eta_n)$ such that

$$\eta_i = \sum_j \beta_j x_{ij}, \quad i = 1, \dots, n$$
$$= X_i \beta$$

- Recall, previously, we let η_i be the E(Y).
- However, this results in a linear regression model.
- If we want to minimize this dependency, let

 $\eta = f(E(Y))$

where $f(\cdot)$ is a function.

Denote,

$$E(Y) = \mu_i$$

Let, $g(\cdot)$ be a monotonic and differentiable function such that

 $\eta_i = g(\mu_i)$

Thus,

$$g(\mu_i) = \sum_j \beta_j x_{ij}, \quad i = 1, \dots, N$$

In words, we are now modeling a function of the mean (or Expectation) as a combination of linear predictors.

Link Function

- The function $g(\cdot)$ is called the "link function" since it links the $E(Y_i)$ to the set of explanatory variables and their estimates.
- For example, if we let g(x) = x (the identify link) and Y is distributed as a Normal R.V., then, we are back to the linear model (either simple linear or multiple linear regression)
- For GLM, you generally have the flexibility to choose what ever link you desire.
- However, there is a **Special** link that we need to consider

 $f(y_i; \theta_i) = a(\theta_i)b(y_i)exp[y_iQ(\theta_i)]$

If we revisit the density function for an exponential, we see a function $Q(\theta_i)$ that looks interesting.

 $Q(\theta)$ is defined as the natural parameter of the distribution.

If we let $g(\cdot)$ be defined such that it transforms the mean to the natural parameter, we have the **Canonical link**

Suppose

$$Y_i \sim Bern(\pi)$$

where $P(Y_i = 1) = \pi$ and $P(Y_i = 0) = 1 - \pi$

Then we previously showed that

$$f(y_i;\pi) = (1-\pi)(1)e^{(y_i \log \frac{\pi}{1-\pi})}$$

with $Q(\pi) = \log(\pi/(1 - \pi))$

So, if we would let

$$g(\pi) = \log(\pi/(1-\pi)) = \sum_{j} \beta_j x_j$$

We would have the canonical link of a Bernoulli/Binomial distribution.

Recall, the function

$$g(\pi) = \log(\pi/(1-\pi))$$

was previously introduced as the 'log odds' and was called the logit.

Lets recap what we have just accomplished.

If we let the random component be **Bernoulli/Binomial** and consider the linking function as **logit**, we can model the log odds ratio as a linear function of covariates using

$$g(\pi) = \sum_{j} \beta_j x_j$$

Since $g(\pi) = \log(\pi/1 - \pi)$, we can write the success probability as

$$\frac{\pi}{1-\pi} = e^{X\beta}$$
$$\pi = e^{X\beta} - \pi e^{X\beta}$$
$$\pi(1+e^{X\beta}) = e^{X\beta}$$

$$\pi \qquad \qquad = \quad \frac{e^{X\beta}}{1+e^{X\beta}}$$

For the logistic model, we have

$$g(\pi) = \log(\pi/(1-\pi)) = \sum_{j} \beta_j x_j$$

To answer the question, consider a model in which you have one predictor (i.e., treatment) and you observe the response MI/No MI.

Let

$$x_{1i} = \begin{cases} 1 & \text{if subject i received the active treatment} \\ 0 & \text{else} \end{cases}$$

and

 $X_i = [1, x_{i1}]$

Thus, if subject i is on active drug,

$$X_i = [1, 1]$$

and if on placebo

Then, the odds for a person on placebo would be

$$\frac{\pi}{1-\pi} = e^{\beta_0 + \beta_1 \cdot 0} = e^{\beta_0}$$

and for a subject on active drug, the log odds would be

$$\frac{\pi}{1-\pi} = e^{\beta_0 + \beta_1 \cdot 1} = e^{\beta_0 + \beta_1}$$

Thus, the odds ratio of a success for comparing active treatment to placebo could be written as

$$OR = \frac{e^{\beta_0 + \beta_1}}{e^{\beta_0}} = e^{\beta_1}$$

or that $log(OR) = \beta_1$.

If you recall, we introduced this notation when we introduced RD, RR and OR.

An implied *advantage* of the GLM formulation is that you can specify other links to derive additional parameter interpretations.

For example, suppose you used the "log" link ($log(\pi) = X\beta$) instead of the "logit" link.

Now, the log link is not the canonical link, but that is OKAY. Then,

$$log(\pi) = \beta_0 + \beta_1 x_{1i}$$

or

$$\pi = e^{\beta_0 + \beta_1 x_{1i}}$$

Therefore, the RR could be viewed as

$$RR = \frac{\pi | x = 1}{\pi | x = 0} = \frac{e^{\beta_0 + \beta_1}}{e^{\beta_0}} = e^{\beta_1}$$

 β_1 can now be interpreted as log Relative Risk.

Recall our famous MI example.

	Myocardial Infarction			
	Fatal Attack or No			
	Nonfatal attack	Attack		
Placebo	189	10845		
Aspirin	104	10933		

Previously, we estimated the OR to be

$$OR = (189 \cdot 10933) / (104 \cdot 10845) = 1.832$$

which indicates that subjects taking placebo had 1.8 times the odds of having an MI when compared to subjects taking aspirin.

For this analysis, we want to use the aspirin group as the reference group and estimate.

Therefore in terms of our regression dummy codes, we want

$$x_{1i} = \begin{cases} 1 & \text{if subject i received PLACEBO} \\ 0 & \text{if subject i received ASPIRIN} \end{cases}$$

with the response coding of

$$Y_i = \begin{cases} 1 & \text{if subject i has either a Fatal MI or a Non Fatal MI} \\ 0 & \text{if subject i does not have an MI} \end{cases}$$

Inputting Data

And we could input this data into SAS as
data one;
 input y x1 count;
 cards;
1 1 189
1 0 104
0 1 10845
0 0 10933
;
run;

And use PROC GENMOD (generalized linear models) to fit the data

```
proc genmod descending;
freq count;
model y = x1 /dist = bin link=logit;
estimate 'X1' x1 1 /exp;
run;
```

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Notes:

- 1. We have specified our design matrix to include just X1. GENMOD automatically includes an intercept unless you tell it not to.
- 2. We used "dist=bin" to specify the distribution of Y as binomial
- 3. We used "link = logit" to fit the canonical link (logistic link)
- 4. The estimate statement invokes a contrast and exponentiates the parameter estimates

Don't worry, we'll become very familiar with GENMOD over the next few weeks.

Response Profile

Ordered Total

Value y Frequency

1 1 293 2 0 21778

PROC GENMOD is modeling the probability that y='1'.

Note: The most important line is the one that indicates what level of the response is considered a success. In this case, we used "DESCENDING" to specify y=1 as the success. (by default, SAS takes the first sorted response category as the success)

Analysis Of Parameter Estimates

			Standard	Wald 95% (
Parameter	DF	Estimate	Error	Limi	ts
Intercept	1	-4.6552	0.0985	-4.8483	-4.4620
xl	1	0.6054	0.1228	0.3647	0.8462

Contrast Estimate Results

		Standard			
Label	Estimate	Error	Alpha	Confidenc	e Limits
X1	0.6054	0.1228	0.05	0.3647	0.8462
Exp(X1)	1.8321	0.2251	0.05	1.4400	2.3308
Therefore our estimate of $OR = 1.832$ with a 95% CI of (1.44, 2.33).					

Using PROC LOGISTIC

proc logistic descending; freq count; model y = x1; *estimate X1 x1 1 /exp; run;

THE OUTPUT

Analysis of Maximum Likelihood Estimates

			Standard	Wald	
Parameter	DF	Estimate	Error	Chi-Square	Pr > ChiSq
Intercept	1	-4.6551	0.0985	2232.4885	<.0001
xl	1	0.6054	0.1228	24.2911	<.0001

Odds Ratio Estimates

	Point	95% Wald	
Effect	Estimate	Confidence	Limits
xl	1.832	1.440	2.331
Difference?	GENMOD is mo	ment-based, but L	OGISTIC uses ML!

Deviance

For a particular GLM for observations $y = (y_1, \ldots, y_n)$, let

 $l(\mu, y)$

denote the log likelihood function expressed in terms of the means $\mu = (\mu_1, \dots, \mu_n)$

Let

 $l(\hat{\mu},y)$

denote the maximum of the log likelihood for the model.

If we have n observations and fit a model with n parameters, we have a saturated model.

We then have a 'perfect fit' of the data (i.e., no degrees of freedom).

Denote the likelihood under the saturated model as

l(y,y)

Then, the DEVIANCE of a model is

$$D = -2[l(\hat{\mu}, y) - l(y, y)]$$

Note:

- 1. D is distributed as χ^2 with df = N p
- 2. *p* is the number of parameters estimated under the alternative (or fitted model).
- 3. Recall, N in the saturated model is the number of parameters included (one for each observation).
- 4. Therefore, using the rules for calculating the df of a contingency table we developed earlier, df equals the difference in parameters estimated under the null (saturated model) and the alternative (at least one β not equal to zero)
- 5. For contingency tables $D \equiv G^2$
- 6. As we proceed, the Deviance will be used to provide a measure of model fit.

If you recall, we used a **Poisson log linear** model to calculate the Pearson residuals for a contingency table.

A saturated model for a contingency table is one that contains an interaction term. For

```
example;
proc genmod;
  model count = x1 y x1*y /dist=poi link = log;
run;
produces a saturated model.
```

Proof

The design matrix for this model would be

 $X = [1, x_{1i}, y_i, x_{1i}y_i]$

Since $x_{1i} = y_i = (0, 1)$ cell (1,1) has X = (1, 1, 1, 1)cell (1,2) has X = (1, 1, 0, 0)cell (2,1) has x = (1, 0, 1, 0)cell (2,2) has x = (1, 0, 0, 0)

That is, counts for all cells are determined by a combination of $X\beta$.

Criteria For Assessing Goodness Of Fit

Criterion	DF	Value	Value/DF	
Deviance	0	0.0000	•	
Scaled Deviance	0	0.0000	•	
Pearson Chi-Square	0	0.0000	•	
Scaled Pearson X2	0	0.0000	•	
Log Likelihood		181840.4662		
Note: $df = 0$ since we are fitting the saturated model ($df = N - N$)				

l(y, y) = 181840.4662

Analysis	Of	Parameter	Estimates
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			Standard	Wald 95% (Confidence
Parameter	DF	Estimate	Error	Limi	lts
Intercept	1	9.2995	0.0096	9.2808	9.3183
xl	1	-0.0081	0.0136	-0.0346	0.0185
У	1	-4.6552	0.0985	-4.8483	-4.4620
xl*y	1	0.6054	0.1228	0.3647	0.8462
— , (

Therefore, cell (1,1)'s count would be

$$log(\text{count cell 1,1}) = 9.2995 - 0.0081 - 4.6552 + 0.6054$$

= 5.2416
or
count cell 1,1 = $e^{5.2416}$
= 189

The Alternative Model

```
proc genmod;
model count = x1 y /dist=poi link = log;
run;
```

Criteria For Assessing Goodness Of Fit

Criterion	DF	Value	Value/DF
Deviance	1	25.3720	25.3720
Scaled Deviance	1	25.3720	25.3720
Pearson Chi-Square	1	25.0139	25.0139
Scaled Pearson X2	1	25.0139	25.0139
Log Likelihood		181827.7802	
$l(\hat{\mu}, y) = 181827.7802$			

Analysis Of Parameter Estimates

Daramatar	DF	Estimate	Standard Error	Wald 95% (Limi	
Parameter	Dr	ESCIMALE	ELLOL		
Intercept	1	9.2956	0.0096	9.2769	9.3144
xl	1	-0.0003	0.0135	-0.0267	0.0261
У	1	-4.3085	0.0588	-4.4238	-4.1932
Thus the even		Leavestfereall (A			

Thus, the expected cell count for cell (1,1) would be

log(expected count cell 1,1)	=	9.2956 - 0.0003 - 4.3085
	=	4.9868
		or
count cell 1,1	=	$e^{4.9868}$
	=	146.467

This is the same values as (with some rounding error) 293 * 11034/22071 = 146.48

Therefore,

$$D = -2(181827.7802 - 181840.4662)$$

= 25.37

on df = 4 - 3 = 1

Summary of Canonical Links

Distribution	Natural Parameter	Canonical Link
Poisson	$\log(\lambda)$	log
Normal	μ	identity
Binomial	$\log(\pi/(1-\pi))$	logit

As stated before, just because the natural parameter suggests a certain link, there is no requirement to model only using the canonical link.

Recap

Some key summary points:

- 1. We generalized linear models by allowing for the specification of the distribution of Y and the relationship of the expectation to the design matrix
- 2. We do not need normality for the regression model
- 3. GLMs provide a unified theory of modeling that encompasses most of the important models for continuous and discrete variables
- 4. As we will see next, model parameters can be estimated by ML
- 5. By restricting the distributions to only exponential class distributions, we can use the same algorithm for ML estimation