Generalised Linear Models 2: Modelling Binary and Proportional Data

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March 15, 2021

Biol 520C: Statistical modelling for biological data

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- If you haven't already picked a dataset for Paper 2 I'd recommend giving it some thought.

Generalised Linear Models

Review





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- 3. Formally specify the 'link' between the mean of Y_i and the deterministic part based on your distributional assumption.



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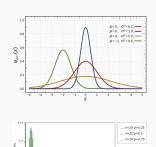
We then saw how to fit GLMs to count data in R using the glm() function.

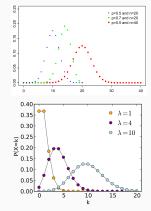
GLM review cont.

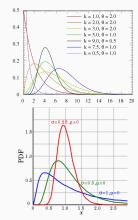




Because R functions streamline the process of fitting GLMs, the key step that's left in your hands is knowing when you will need to switch from a Gaussian model to a GLM, and identifying the correct distribution











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Logistic Regression





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- 3. The independent variables are measured without error.
- 4. The observations are independent.
- 5. There is no collinearity in the independent variables.





If you are going to be working with 0-1 data a lot in your career, you might consider reading a book focused entirely on this subject. A good place to start would be:

Agresti, A. (2018). An introduction to categorical data analysis. John Wiley & Sons.

Logistic Regression on Presence-Absence Data

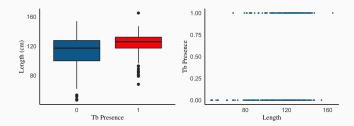
Tuberculosis in Wild Boar



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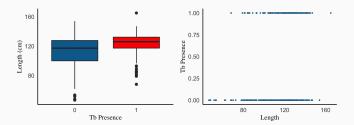
We're going to work with a dataset collected by Vicente et al. (2006). They analyzed the distribution on tuberculosis-like lesions in wild boar (*Sus scrofa*) for potential importance of persistence of tuberculosis in south central Spain.



Tuberculosis in Wild Boar



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With these data we're interested in knowing what whether body size related to Tb prevalence.





We already know a Gaussian model isn't a great choice, but let's see what that would look like.



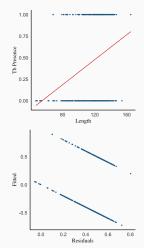
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library(nlme)
Fit_Linear <- gls(Tb ~ Length,
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summary (Fit_Linear)
Generalized least squares fit by maximum likelihood
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       AIC
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Coefficients:
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(Intercept) -0.3912856 0.13589236 -2.879379 0.0042
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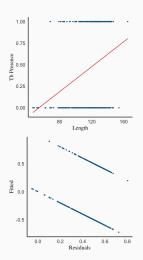


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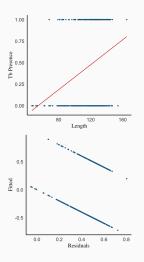
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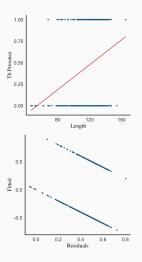






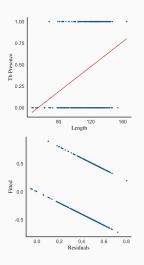
We found a significant relationship between body length and the presence/absence of Tb



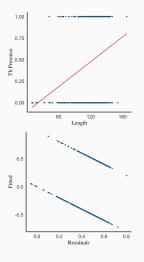


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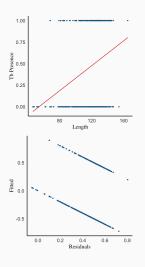
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Our fitted model is:

$$\mathrm{Tb}_i = -0.39 + 0.007 \times \mathrm{Length}_i + \varepsilon_i$$

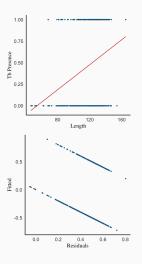


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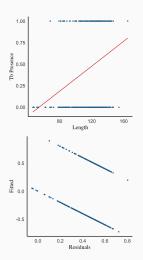
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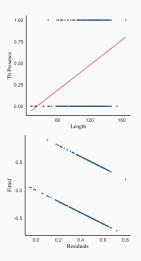
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Can a boar have 0.45 Tb?



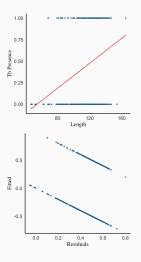






Animals are either infected or not and there is no in between, so we need to redefine what our fitted values mean.

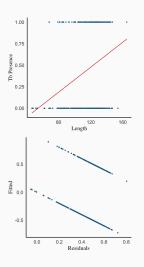




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Instead we re-define our response as π_i , where π_i represents the probability of have Tb, so a 120cm long boar will have a \sim 0.45 chance of having Tb.



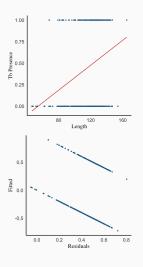


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But our model also says that a 50cm boar has a -0.04 chance of having Tb...

We need a deterministic function that maps the values between 0 and 1, and a dist. that makes more sense.

Step 1: Distributional assumption



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Can you think of a good candidate for 0,1 data?

Binomial distribution

The binomial distribution describes the probability of obtaining k yes/no successes in a sample of size n, or in other words, the distribution of the number of successful trials among a defined number of trials.

Parameters: n and p

Type: Discrete

Biological scenarios: Mark recapture data, live vs dead survival data, killed by a predator or not, yes/no behavioural outcomes, anything with a discrete yes/no outcome.

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PMF: $\binom{n}{k} p^k (1-p)^{n-k}$

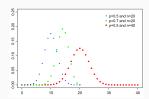
where

$$\binom{n}{k} = \frac{n!}{k!(n-k)!}$$

Range: discrete $(0 \le x \le n)$

Mean: np

Variance: np(1-p)



Step 2: Specify the deterministic model



The second step is to specify the deterministic model (same as always)

$$\pi_i = \beta_0 + \beta_1 \times \text{Length}$$

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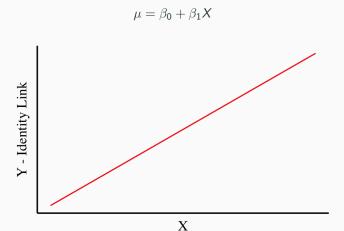
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The last step is to specify a link function that maps the values between 0 and 1.

The logit link

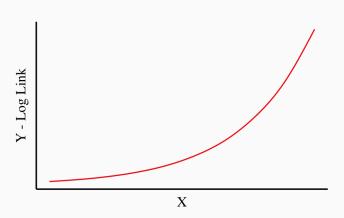


Standard linear regr. with an 'identity link' maps values between $-\infty, \infty$.



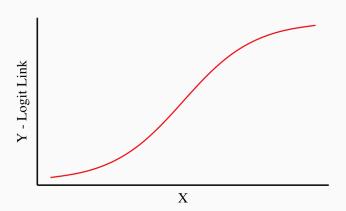
Last lecture we saw how a 'log link' maps values between $0, \infty$.

$$\mu = e^{\beta_0 + \beta_1 X}$$



The 'logit link' is a link function that maps values between 0,1. (How?)

$$\mu = \frac{e^{\beta_0 + \beta_1 X}}{1 + e^{\beta_0 + \beta_1 X}}$$



Logistic regression on 0-1 data



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Now we have all the pieces we need for fitting our GLM:

$$Y_i \sim Binomial(1, \pi_i)$$
 $E(Y_i) = \pi_i$ and $(Y_i) = \pi_i \times (1 - \pi_i)$ $\pi_i = \frac{e^{\beta_0 + \beta_1 \times \text{Length}}}{1 + e^{\beta_0 + \beta_1 \times \text{Length}}}$

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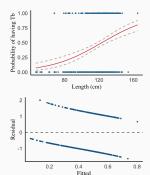
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Coefficients:

Estimate Std. Error z value Pr(>|z|) (Intercept) -4.137107 0.695381 -5.949 2.69e-09 Length 0.033531 0.005767 5.814 6.09e-09

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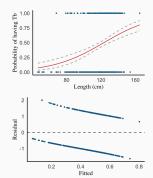
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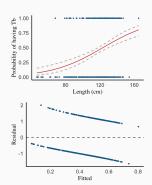
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Are the residuals normally distributed?



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Are the residuals normally distributed? Should they be?



$$Y_i \sim Binomial(1,\pi_i) \qquad E(Y_i) = \pi_i \quad \text{and} \quad (Y_i) = \pi_i \times (1-\pi_i)$$

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ATC: 645 23



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Null deviance: 681.25 on 493 degrees of freedom Residual deviance: 641.23 on 492 degrees of freedom AIC: 665.23



$$Y_i \sim Binomial(1, \pi_i)$$
 $E(Y_i) = \pi_i$ and $(Y_i) = \pi_i \times (1 - \pi_i)$ $\pi_i = \frac{e^{\beta_0 + \beta_1 \times \text{Length}}}{1 + e^{\beta_0 + \beta_1 \times \text{Length}}}$

summary (Fit_Logistic)

Coefficients:

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$$R^2 = \frac{\text{Null dev.-Resid. dev.}}{\text{Null dev.}}$$

$$R^2 = \frac{681.25 - 641.23}{681.25}$$



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We can try and calculate a Pseudo-R² just like we did when we modelled count data:

$$R^2 = \frac{\text{Null dev.-Resid. dev.}}{\text{Null dev.}}$$

$$R^2 = \frac{681.25 - 641.23}{681.25} \sim 6\%$$

In logistic regression Pseudo-R² are almost always going to be low. Not very informative!





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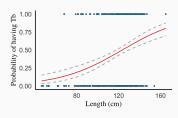
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library(DAAG)

CVbinary(Fit_Logistic)

Fold: 3 8 4 5 9 10 2 6 7 1

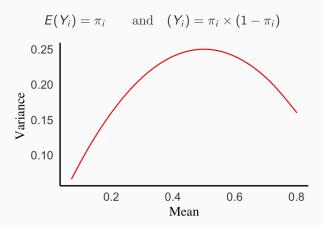
Internal estimate of accuracy = 0.615

Cross-validation estimate of accuracy = 0.615

Mean and Variance



For our model the mean and variance are given by:



Variance is largest for intermediate values of π_i .

Logistic Regression on

Proportion Data





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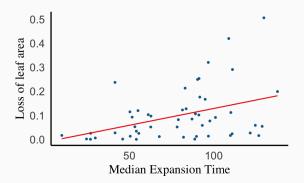
Most of the time, ecologists model proportion data using an $\arcsin(\sqrt{p})$ transformation, but this is not an ideal solution:

Warton, D. I., & Hui, F. K. (2011). The arcsine is asinine: the analysis of proportions in ecology. Ecology, 92(1), 3-10.

Leaf data



This example forms part of a paper asking whether plant species with small leaves have shorter expansion times than large leaved counterparts (Moles & Westoby 2000). The data we're going to work with are the percentage loss of leaf area in relation to median expansion time.



Note A linear regression shows a significant relationship Biol 520C: Statistical modelling for biological data





Leaf Loss_i ~ Binomial(1,
$$\pi_i$$
) $E(Y_i) = \pi_i$ and $(Y_i) = \pi_i \times (1 - \pi_i)$ $\pi_i = \frac{e^{\beta_0 + \beta_1 \times \text{Expansion Time}}}{1 + e^{\beta_0 + \beta_1 \times \text{Expansion Time}}}$



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Fit_Logistic <- glm(loss ~ expansion, family=binomial(link="logit"), data = data)

summary(Fit_Logistic)

Coefficients:

Estimate Std. Error z value Pr(>|z|)
(Intercept) -3.64244    1.58926   -2.292    0.0219
expansion    0.01687    0.01667    1.012    0.3116
---

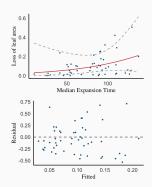
(Dispersion parameter for binomial family taken to be 1)

Null deviance: 6.1273    on 50    degrees of freedom Residual deviance: 5.0196    on 49    degrees of freedom AIC: 17.703
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For example, for our boar data we would say that we assumed our data were binomially distributed and that we modelled them using a logit link:

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Present all of your model outputs in a table, put diagnostic plots in supp. material, if you performed model selection make it clear how you got from A to B.

Where to from here?





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We covered log links and logit links, but there are a number of different link functions that you can use when fitting GLMs. The General idea stays the same, use the one that maps the response variable onto the right scale (e.g., don't use a log-link for a binomial GLM).

Where to now? cont.





We also saw how if our models are of the form:

$$Y_i = \beta_0 + \beta_1 X_i + \varepsilon_i, \qquad \varepsilon_i \sim \mathcal{N}(0, \sigma_i^2)$$

modifying the off-diagonals of the correlation matrix can correct for various forms of autocorrelation.

$$V = \sigma^{2} \begin{bmatrix} 1 & 0 & \cdots & 0 \\ 0 & 1 & \cdots & 0 \\ \vdots & \vdots & \ddots & \vdots \\ 0 & 0 & \cdots & 1 \end{bmatrix}$$

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... but because we're working with different distributions now those approaches don't translate cleanly.

Where to now? cont.



If you find yourself with zero-inflated data you might need to use mixture models that are comprised of combinations of different distributions. See Zuur et al. (2009) Ch. 11.