basic generalized linear models

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library(ggplot2)
theme_set(theme_bw())
library(ggExtra)
library(cowplot)
library(dotwhisker)

Linear models

- foundation for (G)LM(M)s, other complex models
- flexible, robust, computationally efficient, standard
- includes (multiple) regression, ANOVA, ANCOVA, ...
- natural ways to express dependence, interactions

Linear models: assumptions

- response variables:
 - Gaussian (normally distributed)
 - independent
 - conditionally homoscedastic (equal variance)
 - univariate
- predictor variables
 - numeric or categorical (nominal)

Linear models: math

$$z = a + bx + cy + \epsilon$$

or (more predictor variables)

$$y = \beta_0 + \beta_1 x_1 + \beta_2 x_2 + \ldots + \epsilon$$

or (more flexible distribution syntax)

$$y \sim \text{Normal}(\beta_0 + \beta_1 x_1 + \beta_2 x_2 + \dots, \sigma^2)$$

or (more complex sets of predictors)

$$\mu = \mathbf{Xfi}$$
 $y_i \sim \text{Normal}(\mu_i, \sigma^2)$

what does "linear" mean?

- *y* is a linear function of the *parameters* $(\partial^2 y/\partial^2 \beta_i = 0)$
- e.g. polynomials: $y = a + bx + cx^2 + dx^3$
- or sinusoids: $y = a \sin(x) + b \cos(x)$
- but **not**: power-law (ax^b) , exponential $(a \exp(-bx))$

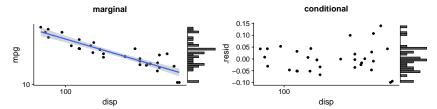
marginal vs. conditional distributions

- common mistake: worry about the overall distribution of the response,
 - rather than the *conditional* distribution (i.e., residuals)
- if only categorical predictors, can mean-correct each group, then look at residuals
- otherwise have to fit the model first!

example

MPG vs displacement for cars

 $cars_lm <- lm(log10(mpg) \sim log10(disp), mtcars)$



(We'll come back to how to judge this later)

categorical predictors

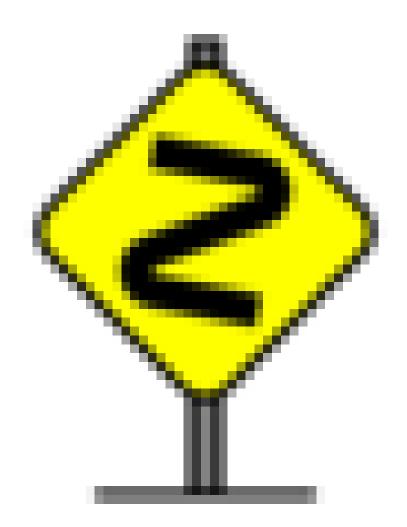
- how do categorical predictors fit into this scheme?
- *dummy variables*: convert to 0/1 values
- R does this automatically with formula syntax
- e.g. for two levels:

```
dd <- data.frame(flavour = rep(c("chocolate",</pre>
    "vanilla"), c(2, 3)))
print(dd)
       flavour
## 1 chocolate
## 2 chocolate
       vanilla
## 4
       vanilla
## 5
       vanilla
```

model.matrix(~flavour, dd)

```
##
     (Intercept) flavourvanilla
## 1
## 2
               1
## 3
               1
                              1
## 4
               1
## 5
                              1
## attr(,"assign")
## [1] 0 1
## attr(,"contrasts")
## attr(,"contrasts")$flavour
## [1] "contr.treatment"
```

• first alphabetical level (chocolate) used as default (use relevel() or factor(...,levels=...) to change default)



ordered factors are handled differently

R formulas

- Wilkinson and Rogers (1973)
- response ~ predictor1 + predictor2 + ...
- numeric variables used "as is"
- categorical variables (factors) converted to dummy variables
- intercept added automatically (1+ ...)
- interaction: : multiplies relevant columns
- a*b: main effect plus interactions
- model.matrix(formula, data)

Formulas, continued

- y~f: 1-way ANOVA
- y~f+g: 2-way ANOVA (additive)
- y~f*g: 2-way ANOVA (with interaction)
- y~x: univariate regression
- y~f+x: ANCOVA (parallel slopes)
- y~f*x: ANCOVA (with interaction, non-parallel slopes)
- y~x1+x2: multivariate regression (additive)
- y~x1*x2: multiv. regression with interaction

If confused, (1) try to write out the equation; (2) model.matrix()

Contrasts

- Machinery for translating categorical variables to dummy (0/1) variables
- treatment contrasts (default):
 - β_1 = intercept = expected value of first level (by default, "aardvark")
 - β_i = difference between level i + 1 and baseline
- sum-to-zero contrasts:
 - β_1 = intercept = unweighted mean of all levels
 - β_i = difference between level *i* and mean; last level not included (!)

too many ways to change contrasts (globally via options(); as attribute of factor; contrasts argument in lm())

Example 1 (treatment contrasts)

Data on ant colonies from Gotelli and Ellison (2004):

```
ants <- data.frame(place = rep(c("field", "forest"),</pre>
    c(6, 4)), colonies = c(12, 9, 12, 10, 9, 6,
    4, 6, 7, 10))
aggregate(colonies ~ place, data = ants, FUN = mean)
      place colonies
## 1 field 9.666667
## 2 forest 6.750000
pr <- function(m) printCoefmat(coef(summary(m)),</pre>
    digits = 3, signif.stars = FALSE)
pr(lm1 <- lm(colonies ~ place, data = ants))</pre>
##
               Estimate Std. Error t value
                  9.667
                             0.958 10.09
## (Intercept)
## placeforest
                 -2.917
                              1.515 -1.92
##
               Pr(>|t|)
## (Intercept)
                  8e-06
## placeforest
                   0.09
Ants: sum-to-zero contrasts
pr(lm2 <- update(lm1, contrasts = list(place = contr.sum)))</pre>
               Estimate Std. Error t value
## (Intercept)
                  8.208
                              0.758
                                      10.83
## place1
                  1.458
                              0.758
                                       1.92
##
               Pr(>|t|)
## (Intercept) 4.7e-06
## place1
                   0.09
data(lizards, package = "brglm")
```

Interactions: example

- Bear road-crossing
- Predictor variables: sex (categorical: M/F), road type (categorical: major/minor), road length (continuous)
- Two-way interactions
 - sex × road length: "are females more sensitive to amount of road than males?"
 - sex × road type: "do females prefer major over minor roads more than males?"
 - road type × road length: "does amount of road affect crossings differently for different road types?"
- Three-way interaction: does the difference of the effect of road length between road types differ between sexes?

Centering (Schielzeth 2010)

- in interaction models, interpretation of main effects **depends on** the center-point of the predictors
- *centering* makes main effects much more interpretable
 - numeric predictors (subtracting the mean by default; other choices could be sensible)
 - categorical predictors: sum-to-zero (weighted or unweighted)
- e.g. if Gregorian year is a predictor, the intercept is at year o (!)
- also improves model stability, decorrelates coefficients

Scaling (Schielzeth 2010)

- scaling parameters improves interpretability
- standard deviation scaling: parameter magnitudes = importance

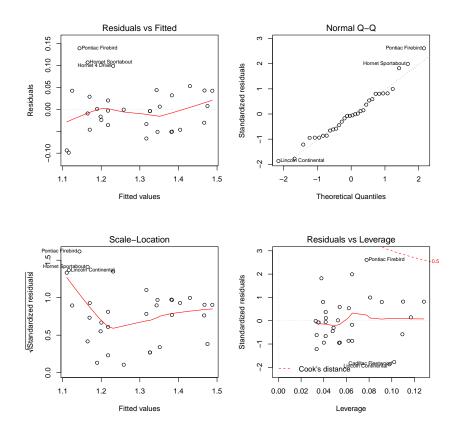
```
mtcars_big <- lm(mpg ~ ., data = mtcars)</pre>
mtcars_big_sc <- lm(mpg ~ ., data = as.data.frame(scale(mtcars)))</pre>
dwfun <- function(., title) {</pre>
    dwplot(., order_vars = names(sort(coef(.)))) +
        geom_vline(xintercept = 0, linetype = 2) +
        ggtitle(title)
}
plot_grid(dwfun(mtcars_big, "unscaled"), dwfun(mtcars_big_sc,
    "scaled"))
                 unscaled
                                                     scaled
      -i5
          -10
```

LM diagnostics

- fitted vs. residual: pattern in mean? (linearity)
- scale-location: pattern in variance? (homoscedasticity)
- Q-Q plot: Normality of residuals
- leverage/Cook's distance: influential points?
- independence is often hard to test
- Normality is the **least important** of these assumptions

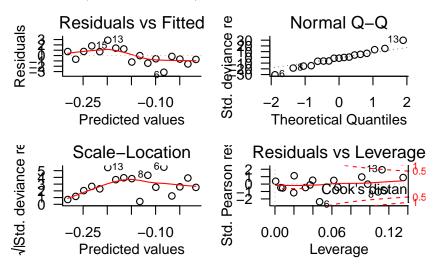
LM diagnostics

```
par(mfrow = c(2, 2))
plot(cars_lm)
```



- problems are not independent
- deal with problems in order (location > scale > outliers > distribu-
- smooth lines help interpretation
- highlighted points are 3 most extreme (id.n argument)

a bad model (Tiwari et al. 2006)



```
original data/fit ...
```

```
tmod <- glm(cbind(notdu, du) ~ density - 1, subset = density >
    0, family = quasibinomial(link = "log"), data = dat)
predframe <- data.frame(density = seq(0, 0.18,
    by = 0.001)
pp <- predict(tmod, newdata = predframe, se.fit = TRUE)</pre>
predframe$duprop <- 1 - exp(pp$fit)</pre>
predframe$low <- 1 - exp(pp$fit + 1.96 * pp$se.fit)</pre>
predframe$high <- 1 - exp(pp$fit - 1.96 * pp$se.fit)</pre>
ggplot(dat, aes(x = density, y = duprop)) + geom_point(aes(size = tot),
    alpha = 0.7, colour = "blue") + geom_line(data = predframe) +
    geom_ribbon(data = predframe, aes(ymin = low,
        ymax = high), colour = NA, alpha = 0.2)
      0.3
                                                        tot
      0.2
                                                            100
                                                            200
                                                             300
      0.1
      0.0
                     0.05
                               0.10
           0.00
                                         0.15
                          density
```

Diagnostics

- statisticians: "don't use p-values to evaluate LM assumptions"
- everyone else: "so what should I do?"
- statisticians: "look at pictures"
- everyone else: "how do I decide whether to worry?"
- statisticians: "..."

testing hypotheses and interpreting results

- parameter-by-parameter: summary() (*t* test)
- multi-parameter comparisons: anova(), car::Anova() (*F* test)
- order matters
- interactions/main effects matter

From LM to GLM

Why GLMs?

- assumptions of LMs do break down sometimes
- count data: discrete, non-negative
- proportion data: discrete counts, $0 \le x \le N$

https://twitter.com/thedavidpowell/status/984432764215754753

GLMs in action

- vast majority of GLMs
 - logistic regression (binary/Bernoulli data)
 - Poisson regression (count data)
- lots of GLM theory carries over from LMs
 - formulas
 - parameter interpretation (partly)
 - diagnostics (partly)

Family

- family: what kind of data do I have?
 - from first principles: family specifies the relationship between the mean and variance
 - binomial: proportions, out of a total number of counts; includes binary (Bernoulli) ("logistic regression")
 - Poisson (independent counts, no set maximum, or far from the maximum)
 - other (Normal ("gaussian"), Gamma)
- default family for glm is Gaussian

link functions

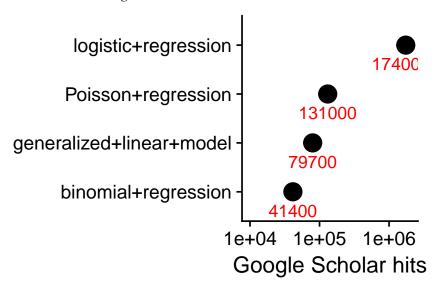
- transform *prediction*, not response
- e.g. rather than $\log(\mu) = \beta_0 + \beta_1 x$, use $\mu = \exp(\beta_0 + \beta_1 x)$
- in this case log is the link function, exp is the inverse link func-
- extreme observations don't cause problems (usually)

family definitions

- link function plus variance function
- typical defaults

- Poisson: log (exponential)
- binomial: logit/log-odds (logistic)

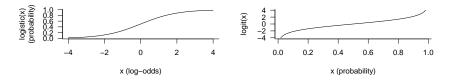
Most GLMs are logistic



log link

- proportional scaling of effects
- small values of coefficients (< 0.1) \approx proportionality
- otherwise change per unit is $exp(\beta)$
- large parameter values (> 10) mean some kind of trouble

logit link/logistic function



- qlogis() function (plogis() is logistic/inverse-link)
- log-odds (log(p/(1-p)))
- most natural scale for probability calculations
- interpretation depends on base probability
 - small probability: like log (proportional)
 - large probability: like log(1-p)
 - intermediate (0.3 < p < 0.7): effect $\approx \beta/4$

diagnostics

• a little harder than linear models: plot is still somewhat useful

- binary data especially hard (e.g. arm::binnedplot)
- goodness of fit tests, R^2 etc. hard (can always compute cor(observed, predict(model, type="response")))
- residuals are *Pearson residuals* by default ((obs exp)/V(exp)); predicted values are on the effect scale (e.g. log/logit) by default (use type="response" to get data-scale predictions)
- also see DHARMa package

back-transformation

- confidence intervals are symmetric on link scale
- can back-transform estimates and CIs for log
- logit is hard (must pick a reference level)
- don't back-transform standard errors!

estimation

- iteratively re-weighted least-squares
- usually Just Works

inference

like LMs, but:

- one-parameter tests are usually *Z* rather than *t*
- CIs based on standard errors are approximate (Wald)
- confint.glm() computes likelihood profile CIs

Common(est?) glm() problems

- neglecting overdispersion
- binomial/Poisson models with non-integer data
- equating negative binomial with binomial rather than Poisson
- failing to specify family (\rightarrow linear model); using glm() for linear models (unnecessary)
- predictions on effect scale
- using (k, N) rather than (k, N k) in binomial models
- worrying about overdispersion unnecessarily (binary/Gamma)
- back-transforming SEs rather than CIs
- ignoring random effects
- Poisson for *underdispersed* responses

Example

```
AIDS (Australia: Dobson & Barnett)
aids <- read.csv("../data/aids.csv")</pre>
aids <- transform(aids, date = year + (quarter -</pre>
    1)/4)
print(gg0 <- ggplot(aids, aes(date, cases)) +</pre>
    geom_point())
    150 -
    100
    50
                      1985
                                                1987
                                                             1988
        1984
                                       date
```

Easy GLMs with ggplot

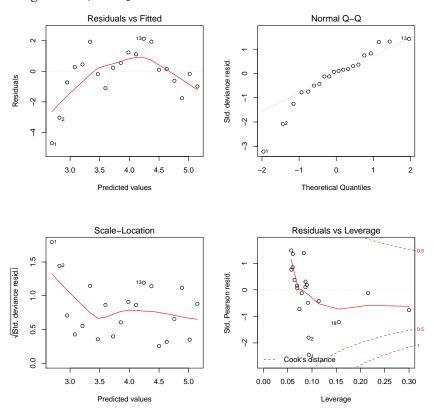
```
print(gg1 <- gg0 + geom_smooth(method = "glm",</pre>
    colour = "red", method.args = list(family = "quasipoisson")))
    200 -
    150
  cases
         1984
                      1985
                                   1986
                                                1987
                                                              1988
                                        date
```

Equivalent code

```
g1 <- glm(cases ~ date, aids, family = quasipoisson(link = "log"))</pre>
summary(g1)
##
## Call:
```

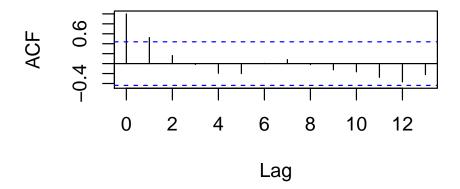
```
## glm(formula = cases ~ date, family = quasipoisson(link = "log"),
##
      data = aids)
##
## Deviance Residuals:
      Min
           1Q Median
                                 30
                                         Max
## -4.7046 -0.7978 0.1218 0.6849 2.1217
##
## Coefficients:
                Estimate Std. Error t value
## (Intercept) -1.023e+03 6.806e+01 -15.03
## date
             5.168e-01 3.425e-02 15.09
              Pr(>|t|)
## (Intercept) 1.25e-11 ***
## date
              1.16e-11 ***
## ---
## Signif. codes:
    0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
## (Dispersion parameter for quasipoisson family taken to be 2.354647)
##
      Null deviance: 677.26 on 19 degrees of freedom
## Residual deviance: 53.02 on 18 degrees of freedom
## AIC: NA
## Number of Fisher Scoring iterations: 4
```

Diagnostics (plot(g1))



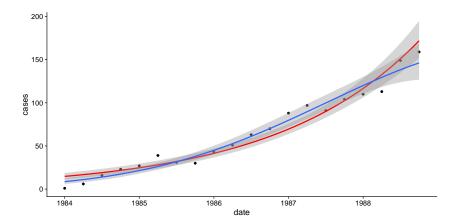
acf(residuals(g1)) ## check autocorrelation

Series residuals(g1)



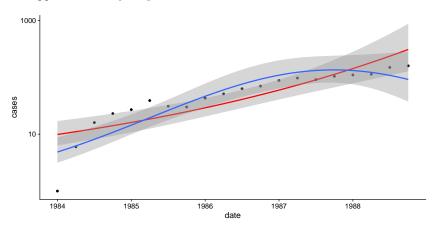
ggplot: check out quadratic model

```
print(gg2 <- gg1 + geom_smooth(method = "glm",</pre>
    formula = y \sim poly(x, 2), method.args = list(family = "quasipoisson")))
```



on log scale

print(gg2 + scale_y_log10())

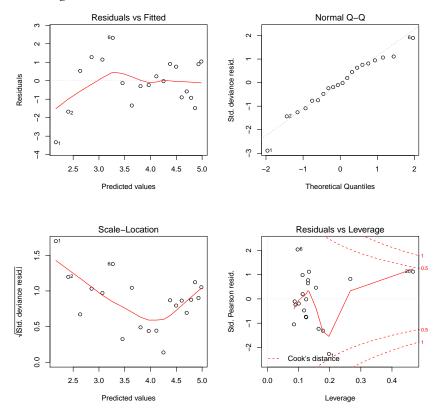


improved model

```
g2 \leftarrow update(g1, . \sim poly(date, 2))
summary(g2)
##
## glm(formula = cases \sim poly(date, 2), family = quasipoisson(link = "log"),
##
       data = aids)
##
## Deviance Residuals:
##
       Min
                 1Q
                      Median
                                    30
                                             Max
## -3.3290 -0.9071 -0.0761
                                0.8985
                                          2.3209
##
## Coefficients:
                   Estimate Std. Error t value
##
                    3.86859
                               0.05004 77.311
## (Intercept)
```

```
## poly(date, 2)1 3.82934 0.25162 15.219
## poly(date, 2)2 -0.68335 0.19716 -3.466
##
                 Pr(>|t|)
## (Intercept) < 2e-16 ***
## poly(date, 2)1 2.46e-11 ***
## poly(date, 2)2 0.00295 **
## ---
## Signif. codes:
   0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for quasipoisson family taken to be 1.657309)
##
##
      Null deviance: 677.264 on 19 degrees of freedom
## Residual deviance: 31.992 on 17 degrees of freedom
## AIC: NA
##
## Number of Fisher Scoring iterations: 4
anova(g1, g2, test = "F") ## for quasi-models specifically
## Analysis of Deviance Table
##
## Model 1: cases ~ date
## Model 2: cases ~ poly(date, 2)
    Resid. Df Resid. Dev Df Deviance
## 1
           18
                 53.020
## 2
           17
                  31.992 1 21.028 12.688
##
      Pr(>F)
## 1
## 2 0.002399 **
## ---
## Signif. codes:
## 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

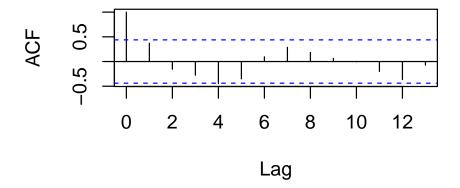
new diagnostics



autocorrelation function

acf(residuals(g2)) ## check autocorrelation

Series residuals(g2)



References

Gotelli, Nicholas J., and Aaron M. Ellison. 2004. A Primer of Ecological Statistics. Sunderland, MA: Sinauer.

Schielzeth, Holger. 2010. "Simple Means to Improve the Interpretability of Regression Coefficients." Methods in Ecology and Evolution 1: 103-13. doi:10.1111/j.2041-210X.2010.00012.x.

Tiwari, Manjula, Karen A. Bjorndal, Alan B. Bolten, and Benjamin M. Bolker. 2006. "Evaluation of Density-Dependent Processes and Green Turtle Chelonia Mydas Hatchling Production at Tortuguero, Costa Rica." Marine Ecology Progress Series 326: 283-93.

Wilkinson, G. N., and C. E. Rogers. 1973. "Symbolic Description of Factorial Models for Analysis of Variance." Applied Statistics 22 (3): 392-99. doi:10.2307/2346786.