Logistic and binomial regression

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//creativecommons.org/licenses/by-nc/3.0/). Please share & remix noncommercially,

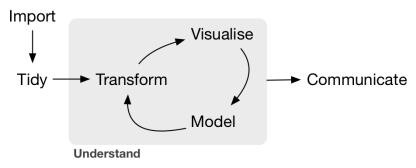
mentioning its origin.

modeling

data analysis road map

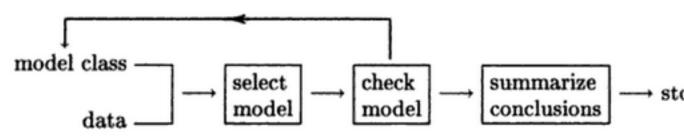
- 1. figure out the (subject-area) question
 - 2. design experiment/data collection (power analysis; simulation)
 - 3. collect data
 - 4. understand the data
 - 5. specify the model; write it down!
 - 6. inspect data (Q/A) (return to 5?
 - 7. fit model
 - 8. graphical diagnostics (return to 5?
 - 9. interpret parameters; inference; plot results

Alternatively



https://jules32.github.io/2016-07-12-0xford/dplyr_tidyr/

also:



(McCullagh and Nelder, 1989)

These are great but *doesn't address the snooping problem*

basics

In principle can use any smooth function from $(0,1) \to \mathbb{R}$ as the link function

- logistic regression: binary data with a logit link (inverse-link=logistic)
- binomial (or aggregated binomial regression: binomial data (maybe logit link, maybe other)
- probit regression: probit link

Binary data and aggregated (N > 1 data) are handled slightly differently.

```
## Error in ggplot(linkdata, aes(x, prob, colour = invlink)):
could not find function "ggplot"
```

```
library(ggplot2)
theme_set(theme_bw())
library(grid)
zmargin <- theme_update(panel.spacing=unit(0,"lines"))</pre>
library(dotwhisker)
library(descr) ## for R^2 measures
library(aods3) ## for overdispersion
```

Contraception data example

```
library(MEMSS)
data("Contraception",package="mlmRev")
head(Contraception)
##
    woman district use livch
                                age urban
## 1
       1
                1
                        3+ 18.4400
## 2
                         0 -5.5599
        2
                1
## 3
        3
                1 N 2 1.4400
                                       Υ
        4
## 4
                1
                    N
                        3+
                             8.4400
                                       Υ
## 5
                         0 -13.5590
                                       Υ
## 6
        6
                1 N
                         0 -11.5600
```

```
cc <- transform(Contraception,</pre>
                use_n=as.numeric(use)-1,
                age_cat=cut(age,breaks=age_breaks))
## Error in cut.default(age, breaks = age_breaks): object
'age_breaks' not found
```

```
gg0 <- ggplot(cc,aes(age,use_n,colour=urban))+
    stat_sum(alpha=0.5)+facet_wrap(~livch,labeller=label_both)
gg0+geom_smooth()
gg0+geom_smooth(method="gam",
                formula=y \sim s(x, k=20),
                method.args=list(family=binomial))
```

Alternative smoothing: binning

```
age_breaks <- seq(-15,20,by=5)
age_mids <- (age_breaks[-1]+age_breaks[-length(age_breaks)])/2</pre>
sumfun <- function(use) {</pre>
    prop <- mean(use)</pre>
    n <- length(use)</pre>
    se <- sqrt(prop*(1-prop)/n) ## approx binomial CIs</pre>
    c(prop=prop, n=n, se=se)
}
cc_agg <- aggregate(use_n~age_cat+urban+livch,</pre>
           data=cc,
           FUN=sumfun)
## ugh!
cc_agg[,c("prop","n","se")] <- cc_agg$use_n</pre>
cc_agg$age_mid <- age_mids[as.numeric(cc_agg$age_cat)]</pre>
gg0+geom_pointrange(data=cc_agg,
                      aes(x=age_mid,
                          y=prop,
                          ymin=prop-2*se,
                          ymax=prop+2*se,
                          size=n),
                      alpha=0.5)+
    scale_size(range=c(1,3))
```

Fit the model: quadratic with all interactions

```
glm(...)
```

```
Explore diagnostics (plot(); DHARMa::simulateResiduals();
arm::binnedplot; mgcv::qq.gam).
  Explore parameters.
```

Computing model predictions

Set up an example to use:

```
lizards <- read.csv("../data/lizards.csv")</pre>
lizards <- transform(lizards,</pre>
                      time=factor(time,levels=c("early","midday","late")))
g1 <- glm(gfrac~height+diameter+light+time,
          lizards,family=binomial,weight=N)
```

Predictions Predictions are fairly easy: set up the new model matrix and multiply by coefficients, then compute the inverse link. This is what predict does (use type="response" to get the backtransformed predictions).

```
newdata <- with(lizards,</pre>
                expand.grid(height=levels(height),
                             diameter=levels(diameter),
                             light=levels(light),
                             time=levels(time)))
## [-2] deletes the response variable; you could also use
## formula(delete.response(terms(g1)))
newX <- model.matrix(formula(g1)[-2], newdata)</pre>
pred0 <- newX %*% coef(g1) ## log-odds
pred <- plogis(pred0)</pre>
                        ## probability
head(c(pred))
## [1] 0.7497750 0.9026836 0.5829216 0.8122611 0.8748663 0.9558361
head(predict(g1,newdata,type="response"))
                     2
                               3
           1
## 0.7497750 0.9026836 0.5829216 0.8122611 0.8748663 0.9558361
```

If you use predict, keep in mind that predict produces predictions on the scale of the linear predictor (type="link") by default rather than on the scale of the original data (type="response").

Confidence intervals Confidence intervals: get new model matrix and compute XVX^T to get variances on the link-function scale. Then compute Normal CIs on the link scale, then back-transform. Or use se=TRUE in predict.

```
pvar <- newX %*% vcov(g1) %*% t(newX)</pre>
pse <- sqrt(diag(pvar))</pre>
```

Or equivalently for any model type where predict has an se.fit argument:

```
pse <- predict(g1,newdata=newdata,se.fit=TRUE)$se.fit</pre>
lwr0 <- pred0-1.96*pse ## or gnorm(0.025)</pre>
upr0 <- pred0+1.96*pse ## or qnorm(0.975)
lwr <- plogis(lwr0)</pre>
upr <- plogis(upr0)</pre>
```

Put the predictions and confidence intervals back into a data frame with the predictor variables:

```
predFrame <- data.frame(newdata,gfrac=pred,lwr,upr)</pre>
```

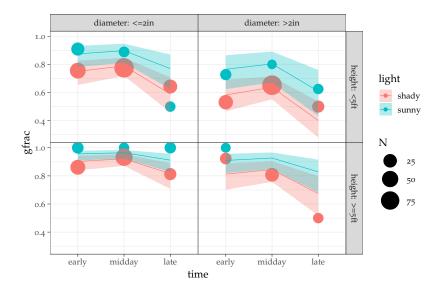
Note:

- back-transforming the standard errors via a logistic usually doesn't make sense: if you want to back-transform them (approximately), you have to multiply them by $(d\mu/d\eta)$, i.e. use dlogis.
- if you use response=TRUE and se.fit=TRUE, R computes the standard errors, scales them as above, and uses them to compute (approximate) symmetric confidence intervals. Unless your sample is very large and/or your predicted probabilities are near 0.5 (so the CIs don't get near o or 1), it's probably best to use the approach above

Getting CIs into ggplot Compute a new data frame, then use geom_ribbon (need to set alpha by hand, and use colour=NA to suppress lines at the edges of the ribbon), unlike when using geom_smooth).

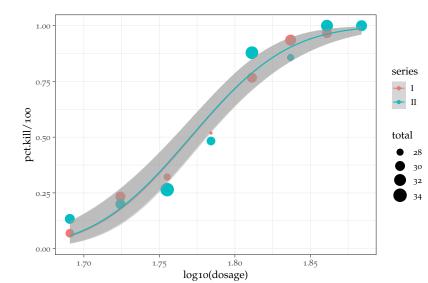
```
gplot0 <- ggplot(lizards,aes(time,gfrac,colour=light))+</pre>
    facet_grid(height~diameter,labeller=label_both)+geom_point(aes(size=N))+
    scale_size_continuous(range=c(3,9))
gplot0 + geom_line(data=predFrame,aes(group=light))+
    geom_ribbon(data=predFrame,
                aes(ymin=lwr,ymax=upr,group=light,fill=light),
                colour=NA,
                alpha=0.3)
```

¹ Dobson, A. J. and A. Barnett (2008, May). An Introduction to Generalized



CIs on nonlinear functions of parameters Tricky. An example presented in ¹, originally from ²:

```
Linear Models, Third Edition (3 ed.).
beetle <- read.csv("../data/beetle2.csv",comment="#")</pre>
                                                                             Chapman and Hall/CRC
## adjust percentages so number killed=integer
                                                                             <sup>2</sup> Bliss, C. I. (1935). The calculation of
beetle <- transform(beetle,</pre>
                                                                             the dosage-mortality curve. Annals of
                                                                             Applied Biology 22(1), 134âĂŞ167
    pct.kill=100/total*round(pct.kill*total/100))
(pplot <- ggplot(beetle,aes(log10(dosage),pct.kill/100,colour=series))+</pre>
    geom_point(aes(size=total))+
    geom_smooth(method=glm, aes(weight=total),
                  method.args=list(family=binomial(link="probit"))))
```



It looks like we can ignore the difference between the series ...

```
g2 <- glm(pct.kill/100~log10(dosage),
          data=beetle,
          family=binomial(link="probit"),
          weight=total)
```

Suppose we're interested in the LD50, i.e. the dose required to kill 50% of the population. Since

$$p = 0.5 = \text{logistic}(\beta_0 + \beta_1 x_{0.5})$$

 $0 = \beta_0 + \beta_1 x_{0.5}$
 $x_{0.5} = -\frac{\beta_0}{\beta_1}$

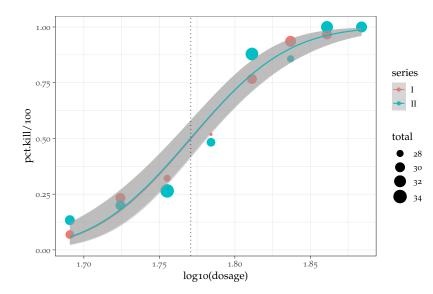
(the units work out correctly too)

So the estimate is

```
cc <- coef(g2)
ld50 <- -cc[1]/cc[2]</pre>
```

Double-check graphically:

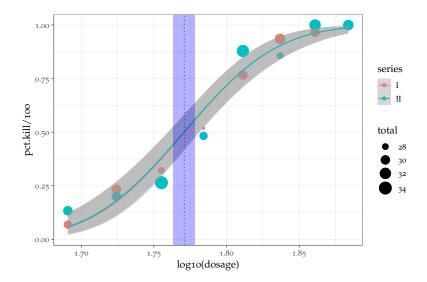




But what about the confidence intervals?

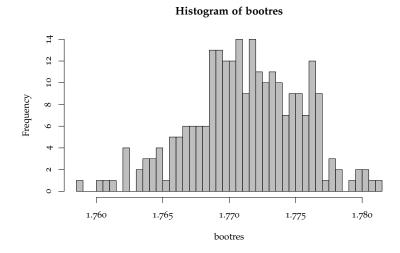
• **delta method**: If we want to compute the variance of f(x, y, z)and $g = (\frac{\partial f}{\partial x}, \frac{\partial f}{\partial y}, \frac{\partial f}{\partial z})$ then the variance is gVg^T (which reduces to $CV^2(f(x,y)) = CV^2(x) + CV^2(y)$ for the case of independent values when f(x,y) = x/y or xy):

```
grad <- c(-1/cc[2],cc[1]/cc[2]^2)
ld50_var <- t(grad) %*% vcov(g2) %*% grad
ld50_se <- c(sqrt(ld50_var)) ## c() converts from matrix to vector (= scalar)</pre>
deltaCI <- ld50+c(-1,1)*1.96*ld50_se</pre>
pplot+geom_vline(xintercept=ld50,linetype=3)+
    annotate("rect",
           xmin=deltaCI[1],
           xmax=deltaCI[2],
           ymin=-Inf,
           ymax=Inf,alpha=0.3,fill="blue",
             colour=NA)
```



• bootstrapping

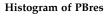
```
bootres <- numeric(250)</pre>
for (i in 1:250) {
    bootdat <- beetle[sample(nrow(beetle), replace=TRUE),]</pre>
    bootmodel <- update(g2,data=bootdat)</pre>
    bootcc <- coef(bootmodel)</pre>
    bootres[i] <- -bootcc[1]/bootcc[2]</pre>
hist(bootres,col="gray",breaks=50)
```

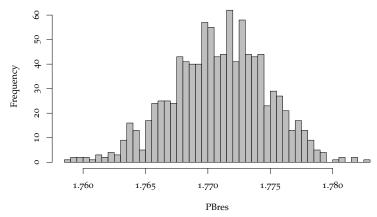


```
bootCI <- quantile(bootres,c(0.025,0.975))</pre>
```

• pseudo-Bayes: MVN sample from parameters

```
library(MASS)
PBsamp <- mvrnorm(1000, mu=coef(g2), Sigma=vcov(g2))
PBres <- -PBsamp[,1]/PBsamp[,2]
hist(PBres, col="gray", breaks=50)
```





```
PBCI <- quantile(PBres,c(0.025,0.975))</pre>
```

In this case the results are all extremely similar:

```
rbind(deltaCI,bootCI,PBCI)
              2.5%
##
                      97.5%
## deltaCI 1.763397 1.778304
## bootCI 1.762387 1.779041
## PBCI 1.763289 1.777993
```

Logistic example

```
## data from http://data.princeton.edu/wws509/datasets/cuse.dat
if (FALSE) {
    try(download.file("http://data.princeton.edu/wws509/datasets/cuse.dat",
                      dest="../data/cuse.dat"),
        silent=TRUE)
}
cuse <- read.table("../data/cuse.dat",header=TRUE)</pre>
```

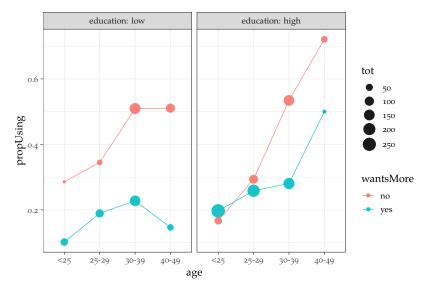
Add convenience variables (proportion and total in each group): change the education factor so that "low" rather than "high" is the baseline group:

```
cuse <- transform(cuse,</pre>
                   propUsing=using/(using+notUsing),
                   tot=using+notUsing,
                   education=relevel(education, "low"))
```

ggplot tricks:

- use label_both in the facet_grid specification to get the subplots labelled by their factor name as well as the level name
- use aes(x=as.numeric(age)) to convince ggplot to connect the factor levels on the x axis with lines; use size=0.5 to make the lines a little skinnier

```
(gg1 <- ggplot(cuse,aes(x=age,y=propUsing,size=tot,colour=wantsMore))+</pre>
  facet_grid(.~education,labeller=label_both)+
  geom_point(alpha=0.9)+
  geom_line(aes(x=as.numeric(age)), size=0.5)+zmargin)
```

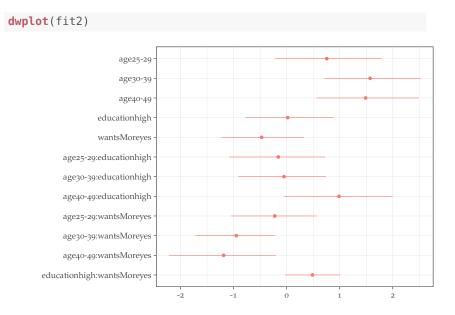


We could fit the three-way interaction, but it would be a bit silly because there would be as many parameters as observations (this is called a saturated model. It would probably be more sensible to worry only about two-way interactions:

```
fit2 <- glm(cbind(using,notUsing)~(age+education+wantsMore)^2,</pre>
            family=binomial,
            data=cuse)
library(aods3)
gof(fit2)
## D = 2.4415, df = 3, P(>D) = 0.4859584
## X2 = 2.5153, df = 3, P(>X2) = 0.4725266
```

There do indeed seem to be important two-way interactions:

```
drop1(fit2,test="Chisq")
## Single term deletions
##
## Model:
## cbind(using, notUsing) \sim (age + education + wantsMore)^2
                      Df Deviance
                                              LRT Pr(>Chi)
##
                                      AIC
## <none>
                            2.4415 99.949
                       3 10.8240 102.332 8.3826 0.03873 *
## age:education
## age:wantsMore
                        3 13.7639 105.272 11.3224 0.01010 *
                           5.7983 101.306 3.3568 0.06693 .
## education:wantsMore 1
## Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```



Interpreting parameters

Based on this model, I'm going to demonstrate how to answer a few specific questions:

- 1. what is the expected odds, and probability (according to the twoway interaction model), that a woman in the age=25-29/education=high/wantsMore=no category is using contraception?
- 2. what is the difference, in terms of log-odds, odds, and probabilities of contraceptive use, between women with low and high educations who are < 25 and don't want more children?
- 3. how would we find the average difference between low- and higheducation women in the same terms?
- 1. To get the log-odds we need to add (intercept+[age effect]+[education effect]+ [age:education]). We don't need to add [wantsMore] or the interactions involved with [wantsMore] because we are in the reference level (no) for that factor.

```
(rcoefs <- coef(fit2)[c("(Intercept)", "age25-29", "educationhigh",</pre>
                          "age25-29:educationhigh")])
##
               (Intercept)
                                           age25-29
                                                              educationhigh
##
               -1.47099549
                                         0.75577018
                                                                  0.01935151
## age25-29:educationhigh
               -0.15789791
##
```

```
(logodds <- sum(rcoefs))</pre>
## [1] -0.8537717
(odds <- exp(logodds)) ## odds</pre>
## [1] 0.4258059
(prob <- odds/(1+odds)) ## probability</pre>
## [1] 0.2986423
## or
(prob <- plogis(logodds))</pre>
## [1] 0.2986423
```

Or

```
newdata <- data.frame(age="25-29",education="high",wantsMore="no")</pre>
predict(fit2,newdata)
##
           1
## -0.8537717
predict(fit2,newdata,type="response")
##
           1
## 0.2986423
```

2. since "don't want more children" and "age < 25" are baseline levels and we are using treatment contrasts, we just need to look at the effects of education:

```
(logodds <- coef(fit2)["educationhigh"])</pre>
## educationhigh
      0.01935151
##
(odds <- exp(logodds))</pre>
## educationhigh
## 1.01954
```

The tricky part here is that we can't calculate the difference in probabilities from the difference coefficients alone; we need to go back and compute the individual probabilities.

```
(lowed_logodds <- coef(fit2)["(Intercept)"])</pre>
## (Intercept)
     -1.470995
(lowed_prob <- plogis(lowed_logodds))</pre>
## (Intercept)
     0.1867914
(highed_logodds <- sum(coef(fit2)[c("(Intercept)", "educationhigh")]))</pre>
## [1] -1.451644
(highed_prob <- plogis(highed_logodds))</pre>
## [1] 0.1897487
(probdiff <- highed_prob-lowed_prob)</pre>
## (Intercept)
## 0.002957333
```

3. Switch to sum contrasts:

```
options(contrasts=c("contr.sum","contr.poly"))
## re-fit the same model:
fit2S <- glm(cbind(using,notUsing)~(age+education+wantsMore)^2,</pre>
            family=binomial,
            data=cuse)
```

We double the difference between the grand mean and low-education women to get the overall difference between low- and higheducation women:

```
(logodds_eddiff <- abs(2*coef(fit2S)["education1"]))</pre>
## education1
## 0.4566298
## can calculate odds as above ...
```

To get the average log-odds for low-education women:

```
(logodds_loed <- sum(coef(fit2S)[c("(Intercept)", "education1")]))</pre>
## [1] -1.049534
```

To get the average log-odds for high-education women we have to subtract the coefficient:

```
(logodds_hied <- coef(fit2S)["(Intercept)"]-coef(fit2S)["education1"])</pre>
## (Intercept)
## -0.5929042
```

From here we can calculate the odds, probabilities, difference in probabilities as above ...

```
options(contrasts=c("contr.treatment","contr.poly")) ## restore defaults
```

Wald tests

Wald tests are based on the local curvature of the likelihood surface, and are the quickest but least reliable tests of significance. They are marginal ("type III") tests, so they test all effects in the presence of all other effects (including interactions).

```
summary(fit2)
##
## Call:
## glm(formula = cbind(using, notUsing) ~ (age + education + wantsMore)^2,
##
       family = binomial, data = cuse)
##
## Deviance Residuals:
##
          1
                    2
                              3
                                                   5
                                                             6
                                                                       7
                        0.21578
## -0.56081
              0.89772
                                 -0.46350
                                           -0.13981
                                                       0.18708
                                                                 0.07522
##
          8
                    9
                             10
                                       11
                                                  12
                                                            13
                                                                       14
## -0.10841
              0.50044
                       -0.39622 -0.42659
                                           0.41098
                                                      -0.26671
                                                                 0.12795
##
         15
                   16
   0.31058
            -0.21319
##
##
## Coefficients:
##
                              Estimate Std. Error z value Pr(>|z|)
                                           0.45337 -3.245 0.001176 **
## (Intercept)
                               -1.47100
```

```
## age25-29
                              0.75577
                                         0.50936 1.484 0.137869
## age30-39
                              1.57249
                                         0.45913 3.425 0.000615 ***
                                         0.48904 3.041 0.002358 **
## age40-49
                              1.48716
## educationhigh
                                         0.42172 0.046 0.963400
                              0.01935
## wantsMoreyes
                             -0.47338
                                         0.39674 -1.193 0.232802
## age25-29:educationhigh
                             -0.15790
                                         0.45838 -0.344 0.730496
## age30-39:educationhigh
                             -0.05178
                                         0.41840 -0.124 0.901501
## age40-49:educationhigh
                              0.98645
                                         0.52114
                                                  1.893 0.058378
## age25-29:wantsMoreyes
                             -0.22536
                                         0.41024 -0.549 0.582773
## age30-39:wantsMoreyes
                             -0.95014
                                         0.38199 -2.487 0.012870 *
## age40-49:wantsMoreyes
                             -1.19012
                                         0.51278 -2.321 0.020291 *
## educationhigh:wantsMoreyes 0.48617
                                         0.26522
                                                 1.833 0.066785 .
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
##
  (Dispersion parameter for binomial family taken to be 1)
##
      Null deviance: 165.7724 on 15 degrees of freedom
##
## Residual deviance: 2.4415 on 3 degrees of freedom
  AIC: 99.949
##
##
## Number of Fisher Scoring iterations: 4
```

This says that the log-odds of contraceptive use in the intercept (age < 25, low) group is significantly different from zero (which means significantly different from probability=0.5); as usual this is not particularly interesting.

The significant values for the main effect here are for the parameters in the presence of the other effects, which means they are tests of differences with respect to age in the baseline (low-education) group. The interpretation of these main effects depends on the contrasts, however.

The other disadvantage of summary, besides its using Wald tests, is that it gives separate tests of each parameter (contrast), rather than a test of the overall effect of the factor (this only matters if the factor has more than two levels). While we can construct Wald F statistics that test the combined effect of several parameters, we might as well use a more accurate likelihood ratio test, based on comparing the goodness of fit (deviance) of two nested models.

We have two choices: drop1 and anova.

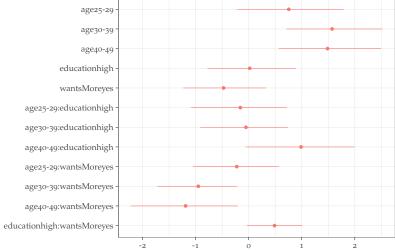
drop1 tries dropping each term out of the model, but it respects marginality, so it does not try to drop main effects if there is an interaction term in the model. If we use test="Chisq" it gives us a likelihood ratio test (otherwise it only provides the difference and

deviance and an AIC value, but not a significance test):

```
drop1(fit2,test="Chisq")
## Single term deletions
##
## Model:
## cbind(using, notUsing) ~ (age + education + wantsMore)^2
##
                     Df Deviance
                                     AIC
                                             LRT Pr(>Chi)
                          2.4415 99.949
## <none>
## age:education
                      3 10.8240 102.332 8.3826 0.03873 *
## age:wantsMore
                      3 13.7639 105.272 11.3224 0.01010 *
## education:wantsMore 1
                          5.7983 101.306 3.3568 0.06693 .
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```

The interaction term is weakly significant here. I am a little bit nervous about dropping it, because (among other things) the magnitudes of the interaction terms are not that much smaller than those of the main effects:





If we use anova it gives us a sequential analysis of deviance (analogous to an analysis of variance): we again need to specify test="Chisq":

```
anova(fit2,test="Chisq")
## Analysis of Deviance Table
##
## Model: binomial, link: logit
```

```
##
## Response: cbind(using, notUsing)
##
## Terms added sequentially (first to last)
##
##
                      Df Deviance Resid. Df Resid. Dev Pr(>Chi)
##
## NULL
                                        15
                                            165.772
                       3
                          79.192
                                        12 86.581 < 2.2e-16 ***
## age
## education
                       1
                          6.162
                                        11
                                               80.418 0.0130496 *
## wantsMore
                       1
                          50.501
                                        10
                                               29.917 1.191e-12 ***
                      3 6.766
## age:education
                                        7
                                               23.151 0.0797373 .
                       3
                         17.353
                                         4
                                               5.798 0.0005979 ***
## age:wantsMore
## education:wantsMore 1
                                         3
                                                2.441 0.0669289 .
                           3.357
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The significance of age quoted here is for age alone; for education, conditional on age being present in the model; for the interaction, conditional on both (it is identical to the result we got above for drop1).

We would get the same result (for the third time) if we explicitly dropped the interaction and did an anova test between the two models:

```
fit3 <- update(fit2,.~.-age:education)</pre>
anova(fit3,fit2,test="Chisq")
## Analysis of Deviance Table
##
## Model 1: cbind(using, notUsing) ~ age + education + wantsMore + age:wantsMore +
       education:wantsMore
## Model 2: cbind(using, notUsing) ~ (age + education + wantsMore)^2
##
    Resid. Df Resid. Dev Df Deviance Pr(>Chi)
            6
## 1
                10.8240
## 2
            3
                  2.4415 3 8.3826 0.03873 *
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```

There are three ways to proceed with this analysis:

- if we are not really interested in the interaction at all we could split the data into two sets and analyze the low-education and high-education data separately;
- we could drop the interaction (i.e. assume that age and education

do not interact);

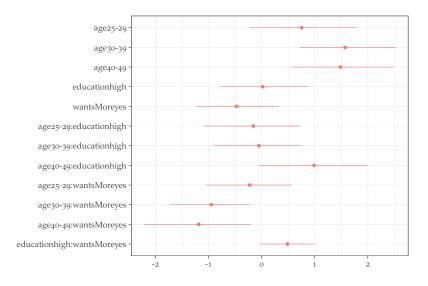
• we could set sum-to-zero contrasts and do "type III" analyses, estimating the average effect of age across education levels and vice versa.

Method 2: drop the interaction. We've already fitted the model (fit2), now we just have to look at it:

Once we have gotten rid of the interaction, the effect of education appears significant:

```
drop1(fit2,test="Chisq")
## Single term deletions
##
## Model:
## cbind(using, notUsing) ~ (age + education + wantsMore)^2
                     Df Deviance
                                    AIC
                                            LRT Pr(>Chi)
##
## <none>
                          2.4415 99.949
                      3 10.8240 102.332 8.3826 0.03873 *
## age:education
## age:wantsMore
                      3 13.7639 105.272 11.3224 0.01010 *
## education:wantsMore 1 5.7983 101.306 3.3568 0.06693 .
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```

dwplot(fit2)



```
options(contrasts=c("contr.sum","contr.poly"))
fit1S <- glm(cbind(using,notUsing)~age*education,family=binomial,</pre>
            data=cuse)
```

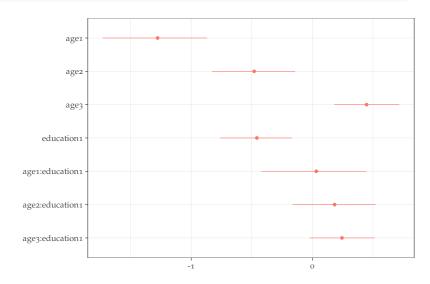
The main effects parameters now represent averages: for example, age1 represents the difference between age (< 25) and age (25 --29) across both education levels ...

```
summary(fit1S)
##
## Call:
## glm(formula = cbind(using, notUsing) ~ age * education, family = binomial,
##
      data = cuse)
##
## Deviance Residuals:
             1Q Median
##
     Min
                              3Q
                                     Max
## -3.7849 -1.0158 -0.0602 1.3826
                                   3.4456
##
## Coefficients:
##
                Estimate Std. Error z value Pr(>|z|)
## (Intercept)
               -0.76622 0.07301 -10.495 < 2e-16 ***
## age1
                ## age2
## age3
                0.30631 0.09369
                                   3.269 0.00108 **
## education1
                -0.22172
                        0.07301 -3.037 0.00239 **
## age1:education1 0.02205 0.14936 0.148 0.88264
## age2:education1 0.12561
                         0.11930 1.053 0.29238
## age3:education1 0.16727
                           0.09369 1.785 0.07420 .
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
## (Dispersion parameter for binomial family taken to be 1)
##
##
      Null deviance: 165.772 on 15 degrees of freedom
## Residual deviance: 73.033 on 8 degrees of freedom
## AIC: 160.54
##
## Number of Fisher Scoring iterations: 4
```

We can use the .~. trick to get drop1 to test all the terms in the model (not just the marginal ones):

```
drop1(fit1S,test="Chisq",.~.)
## Single term deletions
##
## Model:
## cbind(using, notUsing) ~ age * education
              Df Deviance AIC
##
                                    LRT Pr(>Chi)
                    73.033 160.54
## <none>
## age
                3 152.440 233.95 79.407 < 2.2e-16 ***
## education 1 82.804 168.31 9.771 0.001773 **
## age:education 3 80.418 161.93 7.385 0.060575 .
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```

dwplot(fit1S)



confint gives us confidence intervals (95% by default) on the individual parameters. confint.default constructs Wald intervals; confint constructs likelihood profile intervals, which are more accurate — but you can see in this case that they're hardly different:

```
confint.default(fit1S)
##
                         2.5 %
                                     97.5 %
## (Intercept)
                   -0.90931508 -0.62313297
## age1
                   -1.16738905 -0.58192265
## age2
                   -0.56288205 -0.09524232
## age3
                    0.12268409 0.48993966
```

```
## education1 -0.36480902 -0.07862691
## age1:education1 -0.27068499 0.31478141
## age2:education1 -0.10820972 0.35943001
## age3:education1 -0.01635686 0.35089870
confint(fit1S)
## Waiting for profiling to be done...
                        2.5 %
                                   97.5 %
##
## (Intercept) -0.91214655 -0.62513376
## age1
                  -1.18500335 -0.59507053
## age2
                 -0.56679298 -0.09800231
                 0.12356065 0.49130391
## age3
## education1
                 -0.36825983 -0.08135303
## age1:education1 -0.28756291 0.30273336
## age2:education1 -0.11127414 0.35760550
## age3:education1 -0.01476205 0.35293583
```

They're likely to be most different for small data sets and data sets with small numbers of samples per observation: although this data set is only 16 rows, it represents a sample of 1607 total individuals (sum(cuse\$tot)).

pseudo-R² measures

The UCLA statistics site has a very nice description of pseudo- R^2 measures.

- fraction of variance explained
- model improvement
- fraction of deviance explained: (dev(null)-dev(model))/dev(null) ("McFadden"):

```
with(g1,1-deviance/null.deviance)
## [1] 0.7973723
```

• correlation ("Efron"):

```
cor(lizards$gfrac,predict(g1,type="response"))^2
## [1] 0.6350147
```

• Cox and Snell: average deviance explained

$$1 - (L(\text{null})/L(\text{full}))^{2/n}$$

(i.e. look at proportion on the likelihood scale, not the log-likelihood scale)

• Nagelkerke: Cox and Snell, adjusted to max=1

```
descr::LogRegR2(g1)
## Chi2
                       55.89726
## Df
                       5
## Sig.
                       8.532219e-11
## Cox and Snell Index 0.911991
## Nagelkerke Index
                       0.9574288
## McFadden's R2
                0.7973723
```

References

Bliss, C. I. (1935). The calculation of the dosage-mortality curve. Annals of Applied Biology 22(1), 134-167.

Dobson, A. J. and A. Barnett (2008, May). An Introduction to Generalized Linear Models, Third Edition (3 ed.). Chapman and Hall/CRC.

McCullagh, P. and J. A. Nelder (1989). Generalized Linear Models. London: Chapman and Hall.