Logistic Regression

Logistic regression

- When response variable is measured/counted, regression can work well.
- But what if response is yes/no, lived/died, success/failure?
- Model *probability* of success.
- Probability must be between 0 and 1; need method that ensures this.
- *Logistic regression* does this. In R, is a *generalized linear model* with binomial "family":

```
glm(y ~ x, family="binomial")
```

- Begin with simplest case.

Packages

```
library(MASS)
library(tidyverse)
library(marginaleffects)
library(broom)
library(nnet)
library(conflicted)
conflict_prefer("select", "dplyr")
conflict_prefer("filter", "dplyr")
conflict_prefer("rename", "dplyr")
conflict_prefer("summarize", "dplyr")
```

The rats, part 1

- Rats given dose of some poison; either live or die:

```
dose status
0 lived
1 died
2 lived
3 lived
4 died
5 died
```

Read in:

```
my_url <- "http://ritsokiguess.site/datafiles/rat.txt"
rats <- read_delim(my_url, " ")
rats
```

```
# A tibble: 6 x 2
   dose status
  <dbl> <chr>
1     0 lived
2     1 died
3     2 lived
4     3 lived
5     4 died
6     5 died
```

Basic logistic regression

- Make response into a factor first:

```
rats2 <- rats %>% mutate(status = factor(status))
```

- then fit model:

```
status.1 <- glm(status ~ dose, family = "binomial", data = rats2
```

Output

```
summary(status.1)
```

```
Call:
glm(formula = status ~ dose, family = "binomial", data = rats2)

Coefficients:
            Estimate Std. Error z value Pr(>|z|)
(Intercept)   1.6841     1.7979   0.937    0.349
dose         -0.6736     0.6140  -1.097    0.273

(Dispersion parameter for binomial family taken to be 1)

    Null deviance: 8.3178  on 5  degrees of freedom
Residual deviance: 6.7728  on 4  degrees of freedom
AIC: 10.773

Number of Fisher Scoring iterations: 4
```

Interpreting the output

- Like (multiple) regression, get tests of significance of individual x's
- Here not significant (only 6 observations).
- "Slope" for dose is negative, meaning that as dose increases, probability of event modelled (survival) decreases.

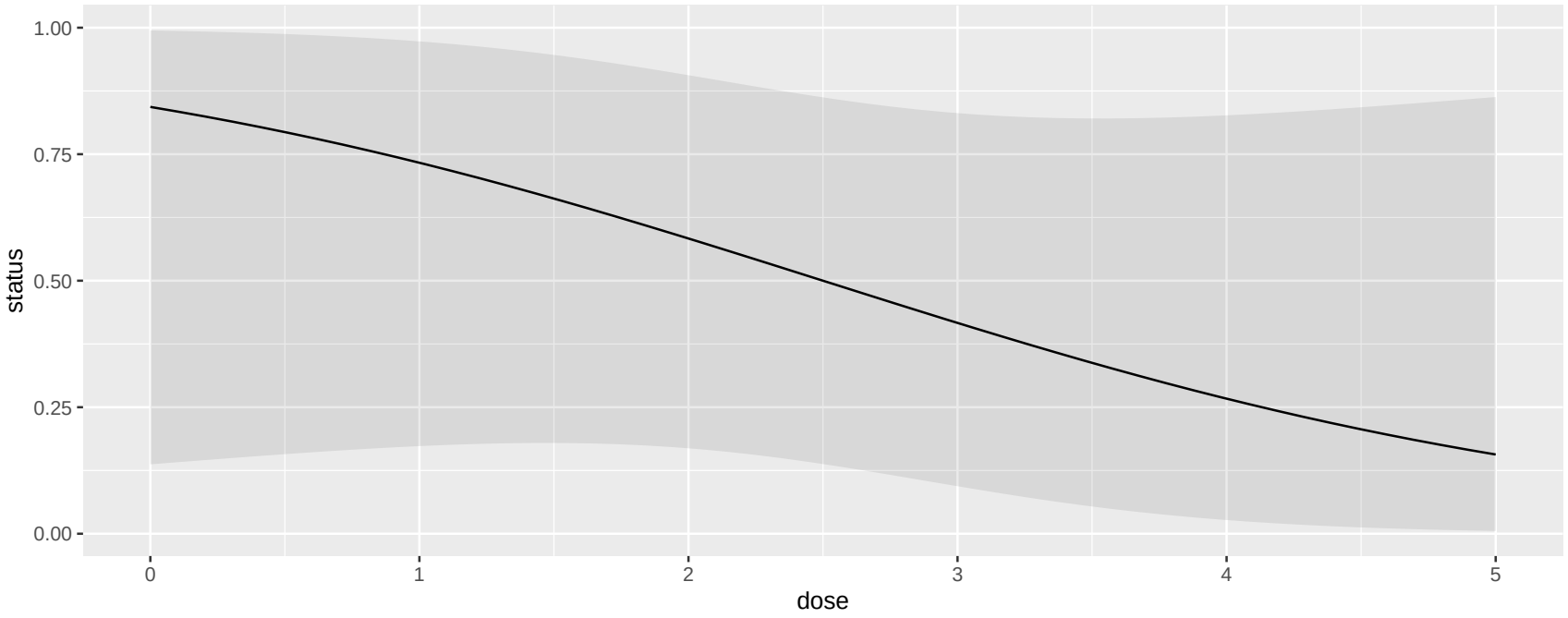
Output part 2: predicted survival probs

```
cbind(predictions(status.1)) %>%
  select(dose, estimate)
```

```
  dose  estimate
1    0 0.8434490
2    1 0.7331122
3    2 0.5834187
4    3 0.4165813
5    4 0.2668878
6    5 0.1565510
```

On a graph

```
plot_cap(status.1, condition = "dose")
```



The rats, more

- More realistic: more rats at each dose (say 10).
- Listing each rat on one line makes a big data file.
- Use format below: dose, number of survivals, number of deaths.

```
dose lived died
0    10    0
1     7    3
2     6    4
3     4    6
4     2    8
5     1    9
```

- 6 lines of data correspond to 60 actual rats.
- Saved in `rat2.txt`.

These data

```
my_url <- "http://ritsokiguess.site/datafiles/rat2.txt"
rat2 <- read_delim(my_url, " ")
rat2
```

```
# A tibble: 6 x 3
   dose lived  died
  <dbl> <dbl> <dbl>
1     0    10     0
2     1     7     3
3     2     6     4
4     3     4     6
5     4     2     8
6     5     1     9
```

Create response matrix:

- Each row contains *multiple* observations.
- Create *two-column* response:
  - #survivals in first column,
  - #deaths in second.

```
response <- with(rat2, cbind(lived, died))
response
```

```
     lived died
[1,]    10    0
[2,]     7    3
[3,]     6    4
[4,]     4    6
[5,]     2    8
[6,]     1    9
```

- Response is R `matrix`:

```
class(response)
```

```
[1] "matrix" "array"
```

Fit logistic regression

- using response you just made:

```
rat2.1 <- glm(response ~ dose,
  family = "binomial",
  data = rat2
)
```

Output

```
summary(rat2.1)
```

```
Call:
glm(formula = response ~ dose, family = "binomial", data = rat2)

Coefficients:
            Estimate Std. Error z value Pr(>|z|)
(Intercept)   2.3619     0.6719   3.515 0.000439 ***
dose         -0.9448     0.2351  -4.018 5.87e-05 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

    Null deviance: 27.530  on 5  degrees of freedom
Residual deviance:  2.474  on 4  degrees of freedom
AIC: 18.94

Number of Fisher Scoring iterations: 4
```

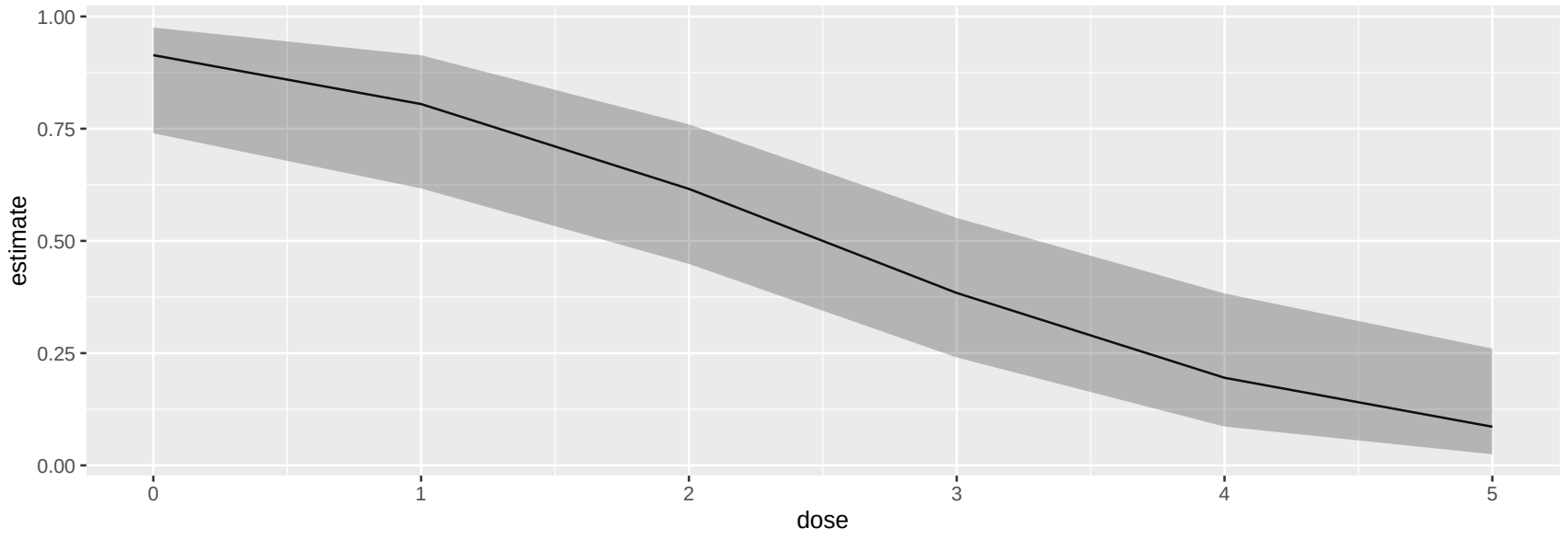
Predicted survival probs

```
new <- datagrid(model = rat2.1, dose = 0:5)
cbind(predictions(rat2.1, newdata = new))
```

```
  rowid  estimate      p.value   s.value   conf.low conf.hi
1     1 0.9138762 0.0004389651 11.153606 0.73983042 0.97536
2     2 0.8048905 0.0031438277  8.313262 0.61695841 0.91353
3     3 0.6159474 0.1721141940  2.538562 0.44876099 0.75959
4     4 0.3840526 0.1721142921  2.538561 0.24040837 0.55123
5     5 0.1951095 0.0031438384  8.313257 0.08646093 0.38304
6     6 0.0861238 0.0004389668 11.153600 0.02463288 0.26016
```

On a picture

```
cbind(predictions(rat2.1, newdata = new)) %>%
  select(estimate, conf.low, conf.high, dose) %>%
  ggplot(aes(x = dose, y = estimate,
             ymin = conf.low, ymax = conf.high)) +
    geom_line() + geom_ribbon(alpha = 0.3)
```



Comments

- Significant effect of dose.
- Effect of larger dose is to *decrease* survival probability ("slope" negative; also see in decreasing predictions.)
- Confidence intervals around prediction narrower (more data).

Multiple logistic regression

- With more than one x, works much like multiple regression.
- Example: study of patients with blood poisoning severe enough to warrant surgery. Relate survival to other potential risk factors.
- Variables, 1=present, 0=absent:
  - survival (death from sepsis=1), response
  - shock
  - malnutrition
  - alcoholism
  - age (as numerical variable)
  - bowel infarction
- See what relates to death.

Read in data

```
my_url <-
  "http://ritsokiguess.site/datafiles/sepsis.txt"
sepsis <- read_delim(my_url, " ")
```

Make sure categoricals really are

```
sepsis %>%
  mutate(across(-age, \(x) factor(x))) -> sepsis
```

The data (some)

```
sepsis
```

```
# A tibble: 106 x 6
   death shock malnut alcohol   age bowelinf
   <fct> <fct> <fct>  <fct>    <dbl> <fct>
 1 0     0     0      0           56 0
 2 0     0     0      0           80 0
 3 0     0     0      0           61 0
 4 0     0     0      0           26 0
 5 0     0     0      0           53 0
 6 1     0     1      0           87 0
 7 0     0     0      0           21 0
 8 1     0     0      1           69 0
 9 0     0     0      0           57 0
10 0     0     1      0           76 0
# i 96 more rows
```

Fit model

```
sepsis.1 <- glm(death ~ shock + malnut + alcohol + age +
  bowelinf,
family = "binomial",
data = sepsis
)
```

Output part 1

```
tidy(sepsis.1)
```

```
# A tibble: 6 x 5
  term         estimate std.error statistic  p.value
  <chr>           <dbl>     <dbl>     <dbl>    <dbl>
1 (Intercept)  -9.75       2.54       -3.84 0.000124
2 shock1        3.67       1.16        3.15 0.00161
3 malnut1       1.22       0.728       1.67 0.0948
4 alcohol1      3.35       0.982       3.42 0.000635
5 age           0.0922     0.0303      3.04 0.00237
6 bowelinf1     2.80       1.16        2.40 0.0162
```

- All P-values fairly small
- but `malnut` not significant: remove.

Removing malnut

```
sepsis.2 <- update(sepsis.1, . ~ . - malnut)
tidy(sepsis.2)
```

```
# A tibble: 5 x 5
  term         estimate std.error statistic  p.value
  <chr>           <dbl>     <dbl>     <dbl>    <dbl>
1 (Intercept)  -8.89       2.32       -3.84 0.000124
2 shock1        3.70       1.10        3.35 0.000797
3 alcohol1      3.19       0.917       3.47 0.000514
4 age           0.0898     0.0292      3.07 0.00211
5 bowelinf1     2.39       1.07        2.23 0.0260
```

- Everything significant now.

Comments

- Most of the original x's helped predict death. Only `malnut` seemed not to add anything.
- Removed `malnut` and tried again.
- Everything remaining is significant (though `bowelinf` actually became *less* significant).
- All coefficients are *positive*, so having any of the risk factors (or being older) *increases* risk of death.

Predictions from model without "malnut"

- A few (rows of original dataframe) chosen "at random":

```
sepsis %>% slice(c(4, 1, 2, 11, 32)) -> new
new
```

```
# A tibble: 5 x 6
  death shock malnut alcohol   age bowelinf
  <fct> <fct> <fct>  <fct>   <dbl> <fct>
1 0     0     0      0          26 0
2 0     0     0      0          56 0
3 0     0     0      0          80 0
4 1     0     0      1          66 1
5 1     0     0      1          49 0
```

```
cbind(predictions(sepsis.2, newdata = new)) %>%
  select(estimate, conf.low, conf.high, shock:bowelinf)
```

```
     estimate     conf.low  conf.high shock malnut alcohol age bowelinf
1 0.001415347 6.272642e-05 0.03103047     0      0       0  26        0
2 0.020552383 4.102504e-03 0.09656596     0      0       0  56        0
3 0.153416834 5.606838e-02 0.35603441     0      0       0  80        0
4 0.931290137 5.490986e-01 0.99341482     0      0       1  66        1
5 0.213000997 7.639063e-02 0.46967947     0      0       1  49        0
```

Comments

- Survival chances pretty good if no risk factors, though decreasing with age.
- Having more than one risk factor reduces survival chances dramatically.
- Usually good job of predicting survival; sometimes death predicted to survive.

Another way to assess effects

of age:

```
new <- datagrid(model = sepsis.2, age = seq(30, 70, 10))
new
```

```
  death shock alcohol bowelinf age
1     0     0       0        0  30
2     0     0       0        0  40
3     0     0       0        0  50
4     0     0       0        0  60
5     0     0       0        0  70
```

Assessing age effect

```
cbind(predictions(sepsis.2, newdata = new)) %>%
  select(estimate, shock:age)
```

```
      estimate shock alcohol bowelinf age
1 0.002026053     0       0        0  30
2 0.004960283     0       0        0  40
3 0.012092515     0       0        0  50
4 0.029179226     0       0        0  60
5 0.068729752     0       0        0  70
```

Assessing shock effect

```
new <- datagrid(shock = c(0, 1), model = sepsis.2)
new
```

```
  death alcohol      age bowelinf shock
1     0       0 51.28302        0     0
2     0       0 51.28302        0     1
```

```
cbind(predictions(sepsis.2, newdata = new)) %>%
  select(estimate, death:shock)
```

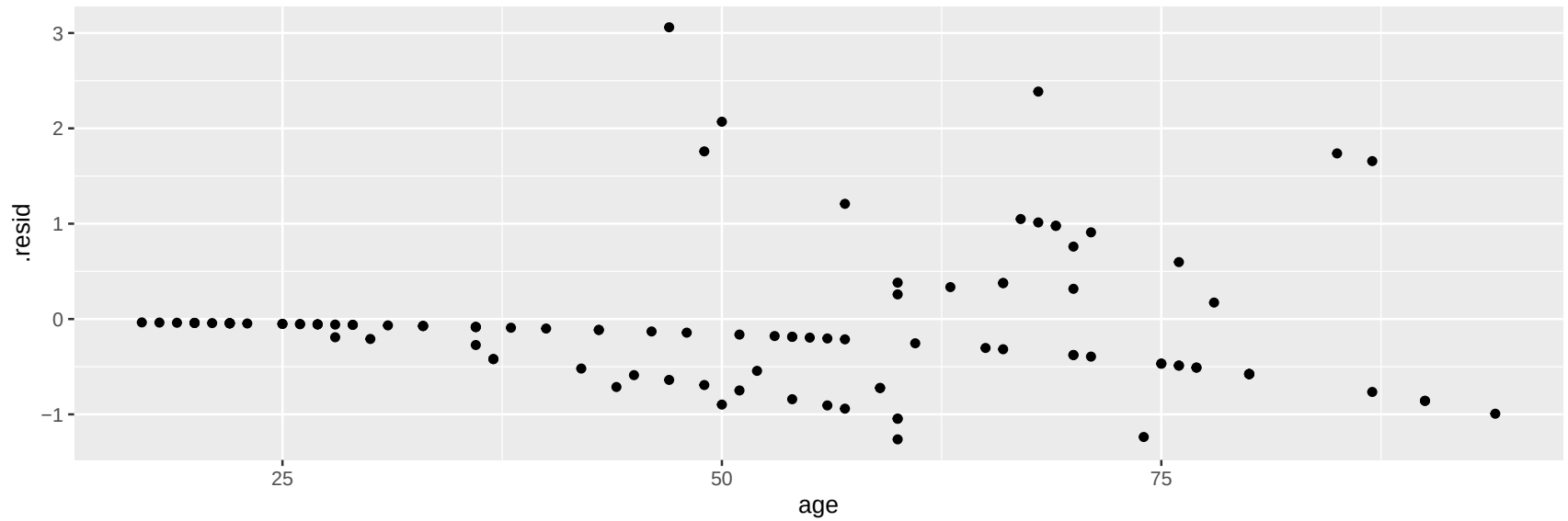
```
    estimate death alcohol      age bowelinf shock
1 0.01354973     0       0 51.28302        0     0
2 0.35742607     0       0 51.28302        0     1
```

Assessing proportionality of odds for age

- An assumption we made is that log-odds of survival depends linearly on age.
- Hard to get your head around, but basic idea is that survival chances go continuously up (or down) with age, instead of (for example) going up and then down.
- In this case, seems reasonable, but should check:

Residuals vs. age

```
sepsis.2 %>% augment(sepsis) %>%
  ggplot(aes(x = age, y = .resid)) +
  geom_point()
```



Comments

- No apparent problems overall.
- Confusing “line” across: no risk factors, survived.

Probability and odds

- For probability p, odds is $p/(1-p)$:

Prob.	Odds	log-odds	in words
0.5	$0.5/0.5 = 1/1 = 1.00$	0.00	"even money"
0.1	$0.1/0.9 = 1/9 = 0.11$	-2.20	"9 to 1"
0.4	$0.4/0.6 = 1/1.5 = 0.67$	-0.41	"1.5 to 1"
0.8	$0.8/0.2 = 4/1 = 4.00$	1.39	"4 to 1 on"

- Gamblers use odds: if you win at 9 to 1 odds, get original stake back plus 9 times the stake.
- Probability has to be between 0 and 1
- Odds between 0 and infinity
- *Log*-odds can be anything: any log-odds corresponds to valid probability.

Odds ratio

- Suppose 90 of 100 men drank wine last week, but only 20 of 100 women.
- Prob of man drinking wine $90/100 = 0.9$, woman $20/100 = 0.2$.
- Odds of man drinking wine $0.9/0.1 = 9$, woman $0.2/0.8 = 0.25$.
- Ratio of odds is $9/0.25 = 36$.
- Way of quantifying difference between men and women: "odds of drinking wine 36 times larger for males than females' '.

Sepsis data again

- Recall prediction of probability of death from risk factors:

```
sepsis.2.tidy <- tidy(sepsis.2)
sepsis.2.tidy
```

```
# A tibble: 5 x 5
  term         estimate std.error statistic  p.value
  <chr>           <dbl>     <dbl>     <dbl>    <dbl>
1 (Intercept)  -8.89       2.32       -3.84 0.000124
2 shock1        3.70       1.10        3.35 0.000797
3 alcohol1      3.19       0.917       3.47 0.000514
4 age           0.0898     0.0292      3.07 0.00211
5 bowelinf1     2.39       1.07        2.23 0.0260
```

- Slopes in column `estimate`.

Multiplying the odds

- Can interpret slopes by taking "exp" of them. We ignore intercept.

```
sepsis.2.tidy %>%
  mutate(exp_coeff=exp(estimate)) %>%
  select(term, exp_coeff)
```

```
# A tibble: 5 x 2
  term        exp_coeff
  <chr>           <dbl>
1 (Intercept)  0.000137
2 shock1      40.5
3 alcohol1    24.2
4 age          1.09
5 bowelinf1   10.9
```

Interpretation

```
# A tibble: 5 x 2
  term         exp_coeff
  <chr>            <dbl>
1 (Intercept)  0.000137
2 shock1      40.5
3 alcohol1    24.2
4 age          1.09
5 bowelinf1   10.9
```

- These say "how much do you *multiply* odds of death by for increase of 1 in corresponding risk factor?" Or, what is odds ratio for that factor being 1 (present) vs. 0 (absent)?
- Eg. being alcoholic vs. not increases odds of death by 24 times
- One year older multiplies odds by about 1.1 times. Over 40 years, about $1.09^{40} = 31$ times.

Odds ratio and relative risk

- **Relative risk** is ratio of probabilities.
- Above: 90 of 100 men (0.9) drank wine, 20 of 100 women (0.2).
- Relative risk 0.9/0.2=4.5. (odds ratio was 36).
- When probabilities small, relative risk and odds ratio similar.
- Eg. prob of man having disease 0.02, woman 0.01.
- Relative risk $0.02/0.01 = 2$.

Odds ratio vs. relative risk

- Odds for men and for women:

```
(od1 <- 0.02 / 0.98) # men
```

```
[1] 0.02040816
```

```
(od2 <- 0.01 / 0.99) # women
```

```
[1] 0.01010101
```

- Odds ratio

```
od1 / od2
```

```
[1] 2.020408
```

- Very close to relative risk of 2.

More than 2 response categories

- With 2 response categories, model the probability of one, and prob of other is one minus that. So doesn't matter which category you model.
- With more than 2 categories, have to think more carefully about the categories: are they
- *ordered*: you can put them in a natural order (like low, medium, high)
- *nominal*: ordering the categories doesn't make sense (like red, green, blue).
- R handles both kinds of response; learn how.

Ordinal response: the miners

- Model probability of being in given category *or lower*.
- Example: coal-miners often suffer disease pneumoconiosis. Likelihood of disease believed to be greater among miners who have worked longer.
- Severity of disease measured on categorical scale: none, moderate, severe.

Miners data

- Data are frequencies:

Exposure	None	Moderate	Severe
5.8	98	0	0
15.0	51	2	1
21.5	34	6	3
27.5	35	5	8
33.5	32	10	9
39.5	23	7	8
46.0	12	6	10
51.5	4	2	5

Reading the data

Data in aligned columns with more than one space between, so:

```
my_url <- "http://ritsokiguess.site/datafiles/miners-tab.txt"
freqs <- read_table(my_url)
```

The data

```
freqs
```

```
# A tibble: 8 x 4
  Exposure  None Moderate Severe
     <dbl> <dbl>    <dbl>  <dbl>
1      5.8    98        0      0
2     15      51        2      1
3     21.5    34        6      3
4     27.5    35        5      8
5     33.5    32       10      9
6     39.5    23        7      8
7     46      12        6     10
8     51.5     4        2      5
```

Tidying

```
freqs %>%
  pivot_longer(-Exposure, names_to = "Severity", values_to
  mutate(Severity = fct_inorder(Severity)) -> miners
```

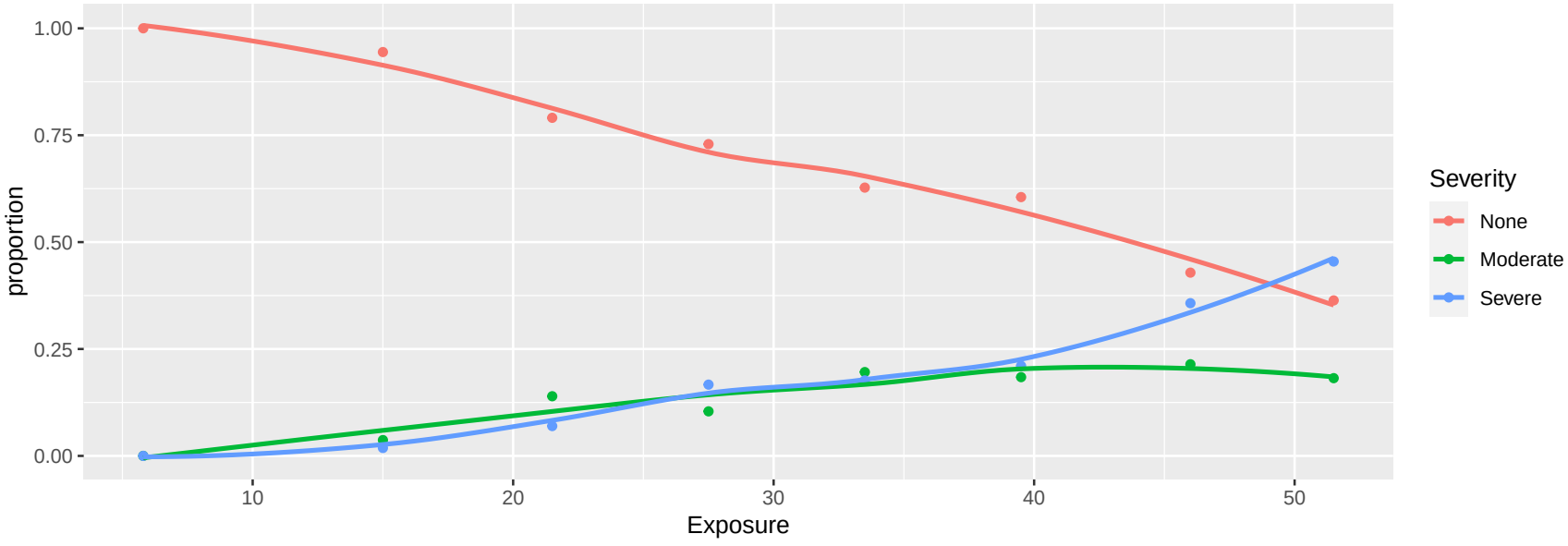
Result

```
miners
```

```
# A tibble: 24 x 3
   Exposure Severity  Freq
      <dbl> <fct>    <dbl>
 1      5.8 None        98
 2      5.8 Moderate     0
 3      5.8 Severe       0
 4     15   None        51
 5     15   Moderate     2
 6     15   Severe       1
 7     21.5 None        34
 8     21.5 Moderate     6
 9     21.5 Severe       3
10     27.5 None        35
# i 14 more rows
```

Plot proportions against exposure

```
miners %>%
  group_by(Exposure) %>%
  mutate(proportion = Freq / sum(Freq)) -> prop
ggplot(prop, aes(x = Exposure, y = proportion,
                 colour = Severity)) +
  geom_point() + geom_smooth(se = F)
```



Reminder of data setup

```
miners
```

```
# A tibble: 24 x 3
   Exposure Severity  Freq
      <dbl> <fct>    <dbl>
 1      5.8 None        98
 2      5.8 Moderate     0
 3      5.8 Severe       0
 4     15   None        51
 5     15   Moderate     2
 6     15   Severe       1
 7     21.5 None        34
 8     21.5 Moderate     6
 9     21.5 Severe       3
10     27.5 None        35
# i 14 more rows
```

Fitting ordered logistic model

Use function `polr` from package `MASS`. Like `glm`.

```
sev.1 <- polr(Severity ~ Exposure,
  weights = Freq,
  data = miners
)
```

Output: not very illuminating

```r
sev.1 <- polr(Severity ~ Exposure,
  weights = Freq,
  data = miners,
  Hess = TRUE
)
```

```r
summary(sev.1)
```

```
Call:
polr(formula = Severity ~ Exposure, data = miners, weights = Freq,
    Hess = TRUE)

Coefficients:
          Value Std. Error t value
Exposure 0.0959    0.01194   8.034

Intercepts:
                Value   Std. Error t value
None|Moderate    3.9558  0.4097     9.6558
Moderate|Severe  4.8690  0.4411    11.0383

Residual Deviance: 416.9188
AIC: 422.9188
```

Does exposure have an effect?

Fit model without Exposure, and compare using anova. Note 1 for model with just intercept:

```
sev.0 <- polr(Severity ~ 1, weights = Freq, data = miners)
anova(sev.0, sev.1)
```

```
Likelihood ratio tests of ordinal regression models

Response: Severity
     Model Resid. df Resid. Dev   Test    Df LR stat.
1        1       369   505.1621                      
2 Exposure       368   416.9188 1 vs 2     1 88.24324
  Pr(Chi)
1        
2       0
```

Exposure definitely has effect on severity of disease.

Another way

- What (if anything) can we drop from model with `exposure`?

```
drop1(sev.1, test = "Chisq")
```

```
Single term deletions

Model:
Severity ~ Exposure
         Df    AIC    LRT  Pr(>Chi)
<none>      422.92
Exposure  1 509.16 88.243 < 2.2e-16 ***
---
Signif. codes:
0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

- Nothing. Exposure definitely has effect.

Predicted probabilities 1/2

```
freqs %>% select(Exposure) -> new
new
```

```
# A tibble: 8 x 1
  Exposure
     <dbl>
1      5.8
2     15
3     21.5
4     27.5
5     33.5
6     39.5
7     46
8     51.5
```

Predicted probabilities 2/2

```
cbind(predictions(sev.1, newdata = new)) %>%
  select(group, estimate, Exposure) %>%
  pivot_wider(names_from = group, values_from = estimate)
```

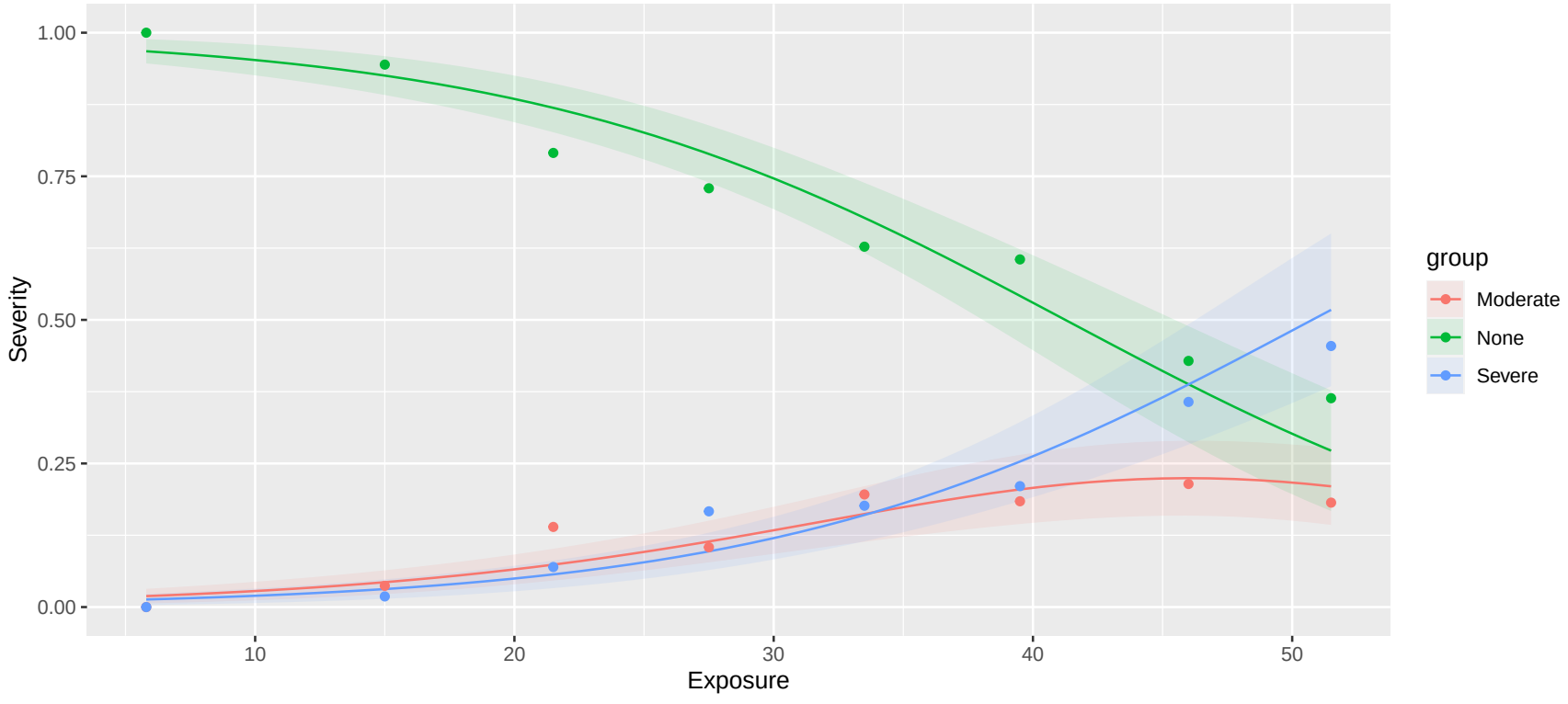
```
# A tibble: 8 x 4
  Exposure  None Moderate Severe
     <dbl> <dbl>    <dbl>  <dbl>
1      5.8 0.968   0.0191 0.0132
2     15   0.925   0.0433 0.0314
3     21.5 0.869   0.0739 0.0569
4     27.5 0.789   0.114  0.0969
5     33.5 0.678   0.162  0.160
6     39.5 0.542   0.205  0.253
7     46   0.388   0.224  0.388
8     51.5 0.272   0.210  0.517
```

Plot of predicted probabilities

```
plot_cap(model = sev.1, condition = c("Exposure", "group"),
         type = "probs") +
  geom_point(data = prop, aes(x = Exposure, y = proportion,
                              colour = Severity)) -> ggg
```

The graph

ggg



Comments

- Model appears to match data well enough.
- As exposure goes up, prob of None goes down, Severe goes up (sharply for high exposure).
- So more exposure means worse disease.

Unordered responses

- With unordered (nominal) responses, can use *generalized logit*.
- Example: 735 people, record age and sex (male 0, female 1), which of 3 brands of some product preferred.
- Data in `mlogit.csv` separated by commas (so `read_csv` will work):

```
my_url <- "http://ritsokiguess.site/datafiles/mlogit.csv"
brandpref <- read_csv(my_url)
```

The data (some)

```
brandpref
```

```
# A tibble: 735 x 3
   brand   sex   age
   <dbl> <dbl> <dbl>
 1     1     0    24
 2     1     0    26
 3     1     0    26
 4     1     1    27
 5     1     1    27
 6     3     1    27
 7     1     0    27
 8     1     0    27
 9     1     1    27
10     1     0    27
# i 725 more rows
```

Bashing into shape

- `sex` and `brand` not meaningful as numbers, so turn into factors:

```
brandpref %>%
  mutate(sex = ifelse(sex == 1, "female", "male"),
         sex = factor(sex),
         brand = factor(brand)
         ) -> brandpref
```

Fitting model

- We use `multinom` from package `nnet`. Works like `polr`.

```
brands.1 <- multinom(brand ~ age + sex, data = brandpref)
```

```
# weights:  12 (6 variable)
initial  value 807.480032
iter  10 value 702.990572
final  value 702.970704
converged
```

Can we drop anything?

- Unfortunately `drop1` seems not to work:

```
drop1(brands.1, test = "Chisq", trace = 0)
```

```
trying - age
```

```
Error in if (trace) {: argument is not interpretable as log
```

- So, fall back on fitting model without what you want to test, and comparing using `anova`.

Do age/sex help predict brand? 1/3

Fit models without each of age and sex:

```
brands.2 <- multinom(brand ~ age, data = brandpref)
```

```
# weights:  9 (4 variable)
initial  value 807.480032
iter  10 value 706.796323
iter  10 value 706.796322
final  value 706.796322
converged
```

```
brands.3 <- multinom(brand ~ sex, data = brandpref)
```

```
# weights:  9 (4 variable)
initial  value 807.480032
final  value 791.861266
converged
```

Do age/sex help predict brand? 2/3

```
anova(brands.2, brands.1)
```

```
Likelihood ratio tests of Multinomial Models

Response: brand
      Model Resid. df Resid. Dev   Test    Df LR stat.
1       age      1466   1413.593
2 age + sex      1464   1405.941 1 vs 2     2 7.651236
     Pr(Chi)
1
2 0.02180496
```

```
anova(brands.3, brands.1)
```

```
Likelihood ratio tests of Multinomial Models

Response: brand
      Model Resid. df Resid. Dev   Test    Df LR stat.
1       sex      1466   1583.723
2 age + sex      1464   1405.941 1 vs 2     2 177.7811
  Pr(Chi)
1
2       0
```

Do age/sex help predict brand? 3/3

- age definitely significant (second `anova`)
- `sex` significant also (first `anova`), though P-value less dramatic
- Keep both.
- Expect to see a large effect of `age`, and a smaller one of `sex`.

Another way to build model

- Start from model with everything and run `step`:

```
step(brands.1, trace = 0)
```

```
trying - age
trying - sex

Call:
multinom(formula = brand ~ age + sex)

Coefficients:
  (Intercept)       age    sexmale
2   -11.25127 0.3682202 -0.5237736
3   -22.25571 0.6859149 -0.4658215

Residual Deviance: 1405.941
AIC: 1417.941
```

- Final model contains both age and sex so neither could be removed.

Making predictions

Find age 5-number summary, and the two sexes:

```
summary(brandpref)
```

```
 brand        sex             age
 1:207   female:466   Min.   :24.0
 2:307   male  :269   1st Qu.:32.0
 3:221                Median :32.0
                      Mean   :32.9
                      3rd Qu.:34.0
                      Max.   :38.0
```

Space the ages out a bit for prediction (see over).

Combinations

```
new <- datagrid(age = c(24, 30, 33, 35, 38),
                sex = c("female", "male"), model = brands.1
new
```

```
   brand age    sex
1      2  24 female
2      2  24   male
3      2  30 female
4      2  30   male
5      2  33 female
6      2  33   male
7      2  35 female
8      2  35   male
9      2  38 female
10     2  38   male
```

The predictions

```
cbind(predictions(brands.1, newdata = new)) %>%
  select(group, estimate, age, sex) %>%
  pivot_wider(names_from = group, values_from = estimate)
```

```
# A tibble: 10 x 5
     age sex        `1`    `2`      `3`
   <dbl> <fct>    <dbl>  <dbl>    <dbl>
 1    24 female 0.915   0.0819 0.00279
 2    24 male   0.948   0.0502 0.00181
 3    30 female 0.500   0.407  0.0933
 4    30 male   0.625   0.302  0.0732
 5    33 female 0.203   0.500  0.297
 6    33 male   0.296   0.432  0.272
 7    35 female 0.0840  0.432  0.484
 8    35 male   0.131   0.397  0.472
 9    38 female 0.0162  0.252  0.732
10    38 male   0.0260  0.239  0.735
```

Comments

- Young males prefer brand 1, but older males prefer brand 3.
- Females similar, but like brand 1 less and brand 2 more.
- A clear `brand` effect, but the `sex` effect is less clear.

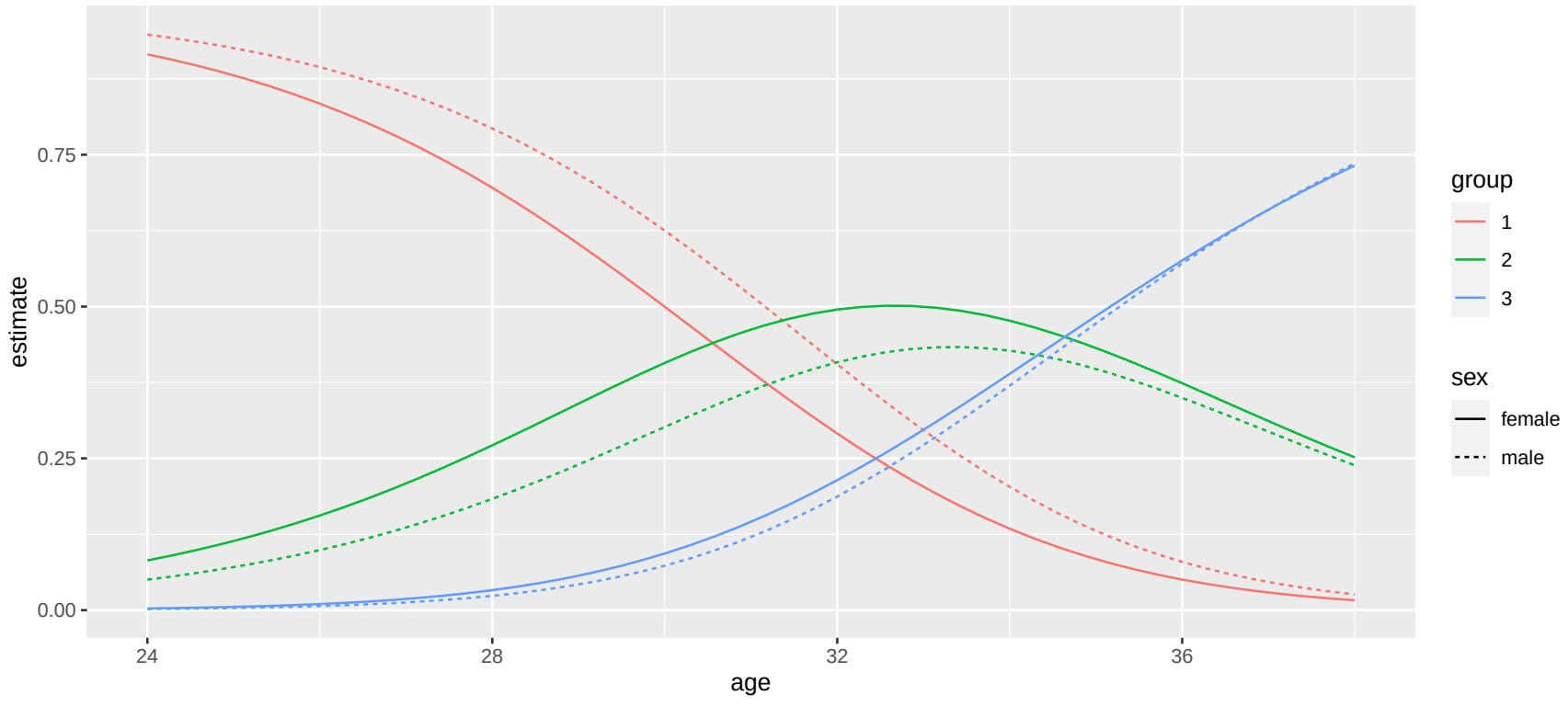
Making a plot

- `plot_cap` doesn't quite work
- so don't draw, edit, *then* make graph:

```
plot_cap(brands.1, condition = c("age", "brand", "sex"),
         type = "probs", draw = FALSE)  %>%
  ggplot(aes(x = age, y = estimate, colour = group,
             linetype = sex)) +
  geom_line() -> g
```

The graph

```
g
```


0.75
0.50
0.25
0.00
estimate
24
28
32
36
age
group
1
2
3
sex
female
male

Digesting the plot

- Brand vs. age: younger people (of both genders) prefer brand 1, but older people (of both genders) prefer brand 3. (Explains significant age effect.)
- Brand vs. sex: females (solid) like brand 1 less than males (dashed), like brand 2 more (for all ages).
- Not much brand difference between genders (solid and dashed lines of same colours close), but enough to be significant.
- Model didn't include interaction, so modelled effect of gender on brand same for each age, modelled effect of age same for each gender. (See also later.)

Alternative data format

Summarize all people of same brand preference, same sex, same age on one line of data file with frequency on end:

```
brandpref
```

```
# A tibble: 735 x 3
   brand sex      age
   <fct> <fct>  <dbl>
 1 1     male      24
 2 1     male      26
 3 1     male      26
 4 1     female    27
 5 1     female    27
 6 3     female    27
 7 1     male      27
 8 1     male      27
 9 1     female    27
10 1     male      27
# i 725 more rows
```

Getting alternative data format

```
brandpref %>%
  group_by(age, sex, brand) %>%
  summarize(Freq = n()) %>%
  ungroup() -> b
b
```

```
# A tibble: 65 x 4
     age sex    brand  Freq
   <dbl> <fct>  <fct> <int>
 1    24 male   1         1
 2    26 male   1         2
 3    27 female 1         4
 4    27 female 3         1
 5    27 male   1         4
 6    28 female 1         6
 7    28 female 2         2
 8    28 female 3         1
 9    28 male   1         4
10    28 male   3         2
```

Fitting models, almost the same

- Just have to remember `weights` to incorporate frequencies.
- Otherwise `multinom` assumes you have just 1 obs on each line!
- Again turn (numerical) `sex` and `brand` into factors:

```
b %>%
  mutate(sex = factor(sex)) %>%
  mutate(brand = factor(brand)) -> bf
b.1 <- multinom(brand ~ age + sex, data = bf, weights = Freq)
b.2 <- multinom(brand ~ age, data = bf, weights = Freq)
```

P-value for sex identical

```
anova(b.2, b.1)
```

```
Likelihood ratio tests of Multinomial Models

Response: brand
      Model Resid. df Resid. Dev   Test    Df LR stat.
1       age       126   1413.593                      
2 age + sex       124   1405.941 1 vs 2     2 7.651236
     Pr(Chi)
1           
2 0.02180496
```

Same P-value as before, so we haven't changed anything important.

Trying interaction between age and gender

```
brands.4 <- update(brands.1, . ~ . + age:sex)
```

```
# weights:  15 (8 variable)
initial  value 807.480032
iter  10 value 703.191146
iter  20 value 702.572260
iter  30 value 702.570900
iter  30 value 702.570893
iter  30 value 702.570893
final  value 702.570893
converged
```

```
anova(brands.1, brands.4)
```

```
Likelihood ratio tests of Multinomial Models

Response: brand
                Model Resid. df Resid. Dev   Test    Df
1           age + sex      1464   1405.941
2 age + sex + age:sex      1462   1405.142 1 vs 2     2
   LR stat.   Pr(Chi)
1
2 0.7996223 0.6704466
```

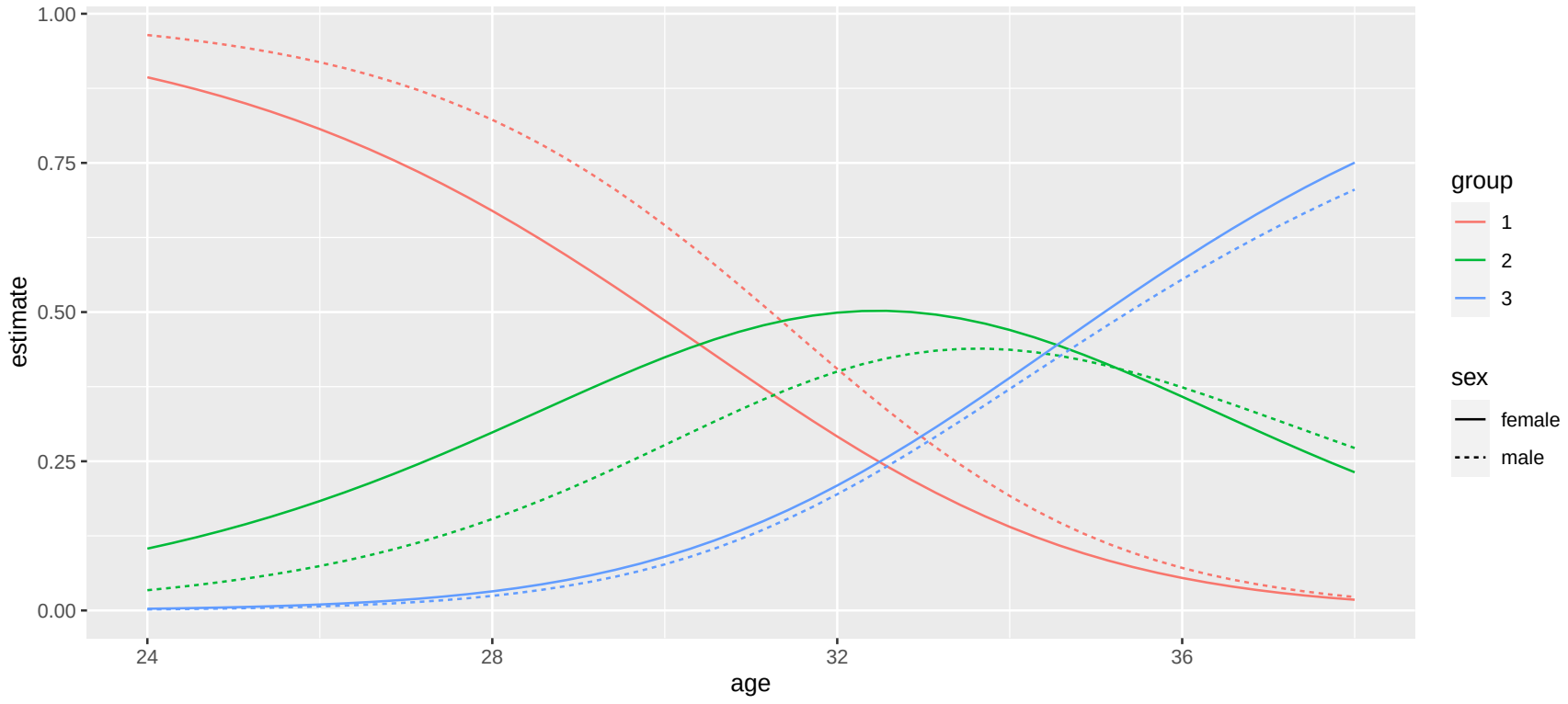
- No evidence that effect of age on brand preference differs for the two genders.

Make graph again

```
plot_cap(brands.4, condition = c("age", "brand", "sex"),
         type = "probs", draw = FALSE)  %>%
  ggplot(aes(x = age, y = estimate, colour = group,
             linetype = sex)) +
  geom_line() -> g4
```

Not much difference in the graph

g4



Compare model without interaction

```
g
```