

ASTA

The ASTA team

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1 Contingency tables

1.1 A contingency table

- We return to the dataset `popularKids`, where we study **association** between 2 **factors**: `Goals` and `Urban.Rural`.
- Based on a sample we make a cross tabulation of the factors and we get a so-called **contingency table** (krydstabel).

```
popKids <- read.delim("https://asta.math.aau.dk/datasets?file=PopularKids.dat")
library(mosaic)
tab <- tally(~Urban.Rural + Goals, data = popKids, margins = TRUE)
tab
```

```
##           Goals
## Urban.Rural Grades Popular Sports Total
##   Rural          57      50      42   149
##   Suburban       87      42      22   151
##   Urban          103      49      26   178
##   Total          247     141      90   478
```

1.1.1 A conditional distribution

- Another representation of data is the percent-wise distribution of `Goals` for each level of `Urban.Rural`, i.e. the sum in each row of the table is 100 (up to rounding):

```
tab <- tally(~Urban.Rural + Goals, data = popKids)
addmargins(round(100 * prop.table(tab, 1)),margin = 2)
```

```
##           Goals
## Urban.Rural Grades Popular Sports Sum
##   Rural          38      34      28  100
##   Suburban       58      28      15  101
##   Urban          58      28      15  101
```

- Here we will talk about the **conditional distribution** of `Goals` given `Urban.Rural`.
- An important question could be:
 - Are the goals of the kids different when they come from urban, suburban or rural areas? I.e. are the rows in the table significantly different?
- There is (almost) no difference between urban and suburban, but it looks like rural is different.

2 Independence

2.1 Independence

- Recall, that two factors are **independent**, when there is no difference between the population's distributions of one factor given the levels of the other factor.
- Otherwise the factors are said to be **dependent**.

- If we e.g. have the following conditional **population distributions** of Goals given Urban.Rural:

```
##           Goals
## Urban.Rural Grades Popular Sports
##   Rural      500      300   200
##   Suburban   500      300   200
##   Urban     500      300   200
```

- Then the factors Goals and Urban.Rural are independent.
- We take a sample and “measure” the factors F_1 and F_2 . E.g. Goals and Urban.Rural for a random child.
- The hypothesis of interest today is:

$$H_0 : F_1 \text{ and } F_2 \text{ are independent, } H_a : F_1 \text{ and } F_2 \text{ are dependent.}$$

2.2 The Chi-squared test for independence

- The relative frequencies in the sample gives an estimate of the unconditional distribution of Goals:

```
n <- margin.table(tab)
pctGoals <- round(100 * margin.table(tab, 2)/n, 1)
pctGoals
```

```
## Goals
## Grades Popular Sports
##   51.7   29.5   18.8
```

- If we assume independence, then this is also a guess of the conditional distributions of Goals given Urban.Rural.
- The corresponding expected counts in the sample are then:

```
##           Goals
## Urban.Rural Grades      Popular      Sports      Sum
##   Rural      77.0 (51.7%)  44.0 (29.5%)  28.1 (18.8%) 149.0 (100%)
##   Suburban   78.0 (51.7%)  44.5 (29.5%)  28.4 (18.8%) 151.0 (100%)
##   Urban     92.0 (51.7%)  52.5 (29.5%)  33.5 (18.8%) 178.0 (100%)
##   Sum       247.0 (51.7%) 141.0 (29.5%)  90.0 (18.8%) 478.0 (100%)
```

2.3 Calculation of expected table

```
pctexptab
```

```
##           Goals
## Urban.Rural Grades      Popular      Sports      Sum
##   Rural      77.0 (51.7%)  44.0 (29.5%)  28.1 (18.8%) 149.0 (100%)
##   Suburban   78.0 (51.7%)  44.5 (29.5%)  28.4 (18.8%) 151.0 (100%)
##   Urban     92.0 (51.7%)  52.5 (29.5%)  33.5 (18.8%) 178.0 (100%)
##   Sum       247.0 (51.7%) 141.0 (29.5%)  90.0 (18.8%) 478.0 (100%)
```

- We note that
 - The relative frequency for a given column is columnTotal divided by tableTotal. For example **Grades**, which is $\frac{247}{478} = 51.7\%$.
 - The expected value in a given cell in the table is then the cell’s relative column frequency multiplied by the cell’s rowTotal. For example **Rural** and **Grades**: $149 \times 51.7\% = 77.0$.
- This can be summarized to:
 - The expected value in a cell is the product of the cell’s rowTotal and columnTotal divided by tableTotal.

2.4 Chi-squared (χ^2) test statistic

- We have an **observed table**:

```
tab
```

```
##           Goals
## Urban.Rural Grades Popular Sports
##   Rural      57      50      42
##   Suburban   87      42      22
##   Urban     103      49      26
```

- And an **expected table**, if H_0 is true:

```
##           Goals
## Urban.Rural Grades Popular Sports Sum
##   Rural      77.0  44.0  28.1 149.0
##   Suburban   78.0  44.5  28.4 151.0
##   Urban      92.0  52.5  33.5 178.0
##   Sum       247.0 141.0  90.0 478.0
```

- If these tables are “far from each other”, then we reject H_0 . We want to measure the distance via the Chi-squared test statistic:
 - $X^2 = \sum \frac{(f_o - f_e)^2}{f_e}$: Sum over all cells in the table
 - f_o is the frequency in a cell in the observed table
 - f_e is the corresponding frequency in the expected table.

- We have:

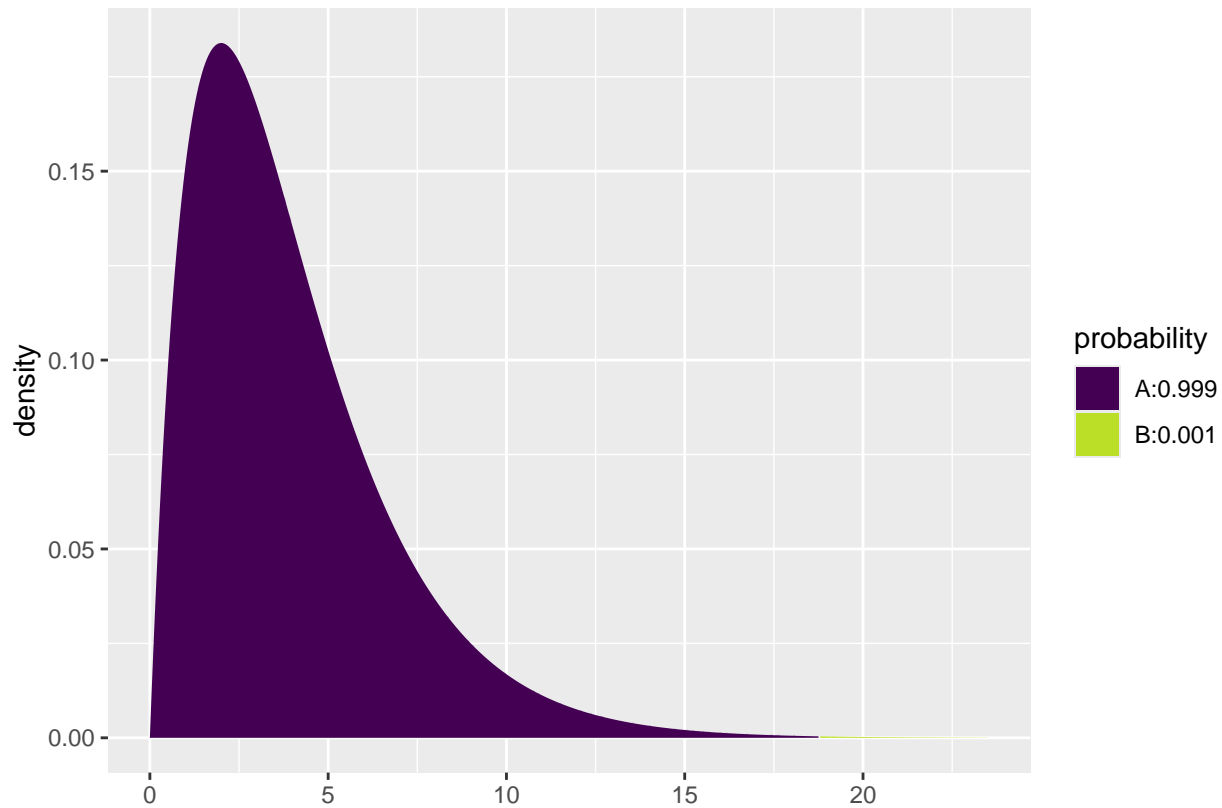
$$X_{obs}^2 = \frac{(57 - 77)^2}{77} + \dots + \frac{(26 - 33.5)^2}{33.5} = 18.8$$

- Is this a large distance??

2.5 χ^2 -test template.

- We want to test the hypothesis H_0 of independence in a table with r rows and c columns:
 - We take a sample and calculate X_{obs}^2 - the observed value of the test statistic.
 - p-value: Assume H_0 is true. What is then the chance of obtaining a larger X^2 than X_{obs}^2 , if we repeat the experiment?
- This can be approximated by the χ^2 -**distribution** with $df = (r - 1)(c - 1)$ degrees of freedom.
- For **Goals** and **Urban.Rural** we have $r = c = 3$, i.e. $df = 4$ and $X_{obs}^2 = 18.8$, so the p-value is:

```
1 - pdist("chisq", 18.8, df = 4)
```



```
## [1] 0.0008603303
```

- There is clearly a significant association between Goals and Urban.Rural.

2.6 The function `chisq.test`.

- All of the above calculations can be obtained by the function `chisq.test`.

```
tab <- tally(~ Urban.Rural + Goals, data = popKids)
testStat <- chisq.test(tab, correct = FALSE)
testStat
```

```
##
## Pearson's Chi-squared test
##
## data:  tab
## X-squared = 18.828, df = 4, p-value = 0.0008497
```

```
testStat$expected
```

```
##           Goals
## Urban.Rural  Grades  Popular  Sports
##   Rural    76.99372 43.95188 28.05439
##   Suburban 78.02720 44.54184 28.43096
##   Urban   91.97908 52.50628 33.51464
```

-
- The frequency data can also be put directly into a matrix.

```
data <- c(57, 87, 103, 50, 42, 49, 42, 22, 26)
tab <- matrix(data, nrow = 3, ncol = 3)
```

```
row.names(tab) <- c("Rural", "Suburban", "Urban")
colnames(tab) <- c("Grades", "Popular", "Sports")
tab
```

```
##           Grades Popular Sports
## Rural           57      50    42
## Suburban        87      42    22
## Urban          103      49    26
```

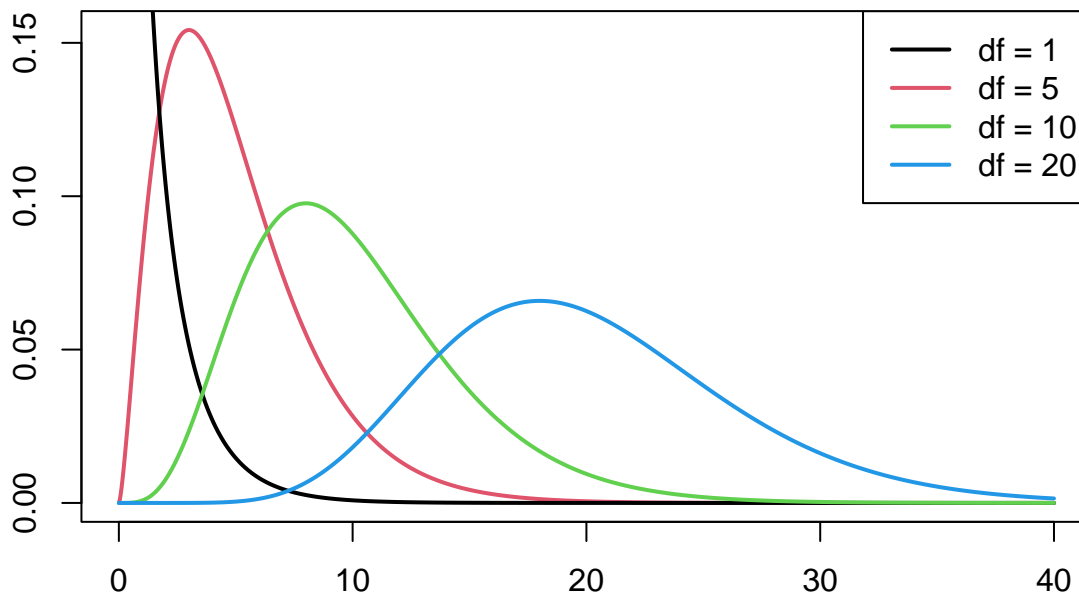
```
chisq.test(tab)
```

```
##
## Pearson's Chi-squared test
##
## data:  tab
## X-squared = 18.828, df = 4, p-value = 0.0008497
```

3 The χ^2 -distribution

3.1 The χ^2 -distribution

- The χ^2 -distribution with df degrees of freedom:
 - Is never negative.
 - Has mean $\mu = df$
 - Has standard deviation $\sigma = \sqrt{2df}$
 - Is skewed to the right, but approaches a normal distribution when df grows.



4 Agresti - Summary

4.1 Summary

- For the the Chi-squared statistic, X^2 , to be appropriate we require that the expected values have to be $f_e \geq 5$.
- Now we can summarize the ingredients in the Chi-squared test for independence.

TABLE 8.5: The Five Parts of the Chi-Squared Test of Independence

-
1. Assumptions: Two categorical variables, random sampling, $f_e \geq 5$ in all cells
 2. Hypotheses: H_0 : Statistical independence of variables
 H_a : Statistical dependence of variables
 3. Test statistic: $\chi^2 = \sum \frac{(f_o - f_e)^2}{f_e}$, where $f_e = \frac{(\text{Row total})(\text{Column total})}{\text{Total sample size}}$
 4. P -value: $P =$ right-tail probability above observed χ^2 value,
for chi-squared distribution with $df = (r - 1)(c - 1)$
 5. Conclusion: Report P -value
If decision needed, reject H_0 at α -level if $P \leq \alpha$
-

5 Standardized residuals

5.1 Residual analysis

- If we reject the hypothesis of independence it can be of interest to identify the significant deviations.
- In a given cell in the table, $f_o - f_e$ is the deviation between data and the expected values under the null hypothesis.
- We assume that $f_e \geq 5$.
- If H_0 is true, then the standard error of $f_o - f_e$ is given by

$$se = \sqrt{f_e(1 - \text{rowProportion})(1 - \text{columnProportion})}$$

- The corresponding z -score

$$z = \frac{f_o - f_e}{se}$$

should in 95% of the cells be between ± 2 . Values above 3 or below -3 should not appear.

- In popKids table cell **Rural** and **Grade** we got $f_e = 77.0$ and $f_o = 57$. Here $\text{columnProportion} = 51.7\%$ and $\text{rowProportion} = 149/478 = 31.2\%$.
- We can then calculate

$$z = \frac{57 - 77}{\sqrt{77(1 - 0.517)(1 - 0.312)}} = -3.95$$

- Compared to the null hypothesis there are way too few rural kids who find grades important.
- In summary: The standardized residuals allow for cell-by-cell (f_e vs f_o) comparison.

5.2 Residual analysis in R

- In R we can extract the standardized residuals from the output of `chisq.test`:

```
tab <- tally(~ Urban.Rural + Goals, data = popKids)
testStat <- chisq.test(tab, correct = FALSE)
testStat$stdres
```

```
##           Goals
## Urban.Rural  Grades  Popular  Sports
## Rural      -3.9508449  1.3096235  3.5225004
## Suburban    1.7666608 -0.5484075 -1.6185210
## Urban       2.0865780 -0.7274327 -1.8186224
```

5.3 Why not just use two-way ANOVA ?

- number of persons in different categories are *not* normally distributed
- variance typically larger the larger expected frequency
- underlying data are discrete (for each person, which column and row category does person belong to)
- these discrete variables are naturally modelled in terms of probabilities for different categories
- therefore hypothesis of independence becomes natural null hypothesis
- it is possible to model table frequencies as dependent variable using a regression model but then we need the framework of *generalized linear models* (see last slides)

Contingency table:

- *counts* of how many individuals fall within different categories for two (or more) categorical variables

Two-way ANOVA:

- a number of individuals/objects/... available for each combination of two categorical variables
- next a continuous variable is measured for each individual or object (this becomes the response variable)

6 Models for table data in R

6.1 Example

- We will study the dataset `HairEyeColor`.

```
HairEyeColor <- read.delim("https://asta.math.aau.dk/datasets?file=HairEyeColor.txt")
head(HairEyeColor)
```

```
##   Hair  Eye  Sex Freq
## 1 Black Brown Male   32
## 2 Brown Brown Male   53
## 3  Red Brown Male   10
## 4 Blond Brown Male    3
## 5 Black  Blue Male   11
## 6 Brown  Blue Male   50
```

- Data is organized such that the variable `Freq` gives the frequency of each combination of the factors `Hair`, `Eye` and `Sex`.
- For example: 32 observations are men with black hair and brown eyes.
- We are interested in the association between eye color and hair color ignoring the sex
- We aggregate data, so we have a table with frequencies for each combination of `Hair` and `Eye`.

```
HairEye <- aggregate(Freq ~ Eye + Hair, FUN = sum, data = HairEyeColor)
HairEye
```

```
##      Eye  Hair  Freq
## 1  Blue Black   20
## 2  Brown Black   68
## 3  Green Black    5
## 4  Hazel Black   15
## 5  Blue  Blond   94
## 6  Brown  Blond    7
## 7  Green  Blond   16
## 8  Hazel  Blond   10
## 9  Blue  Brown   84
## 10 Brown  Brown  119
## 11 Green  Brown   29
## 12 Hazel  Brown   54
```



```
## 13 Blue Red 17
## 14 Brown Red 26
## 15 Green Red 14
## 16 Hazel Red 14
```

6.2 Model specification

- We can write down a model for (the logarithm of) the expected frequencies by using dummy variables z_{e1}, z_{e2}, z_{e3} and z_{h1}, z_{h2}, z_{h3}
- To denote the different levels of **Eye** and **Hair** (the reference level has all dummy variables equal to 0):

$$\log(f_e) = \alpha + \beta_{e1}z_{e1} + \beta_{e2}z_{e2} + \beta_{e3}z_{e3} + \beta_{h1}z_{h1} + \beta_{h2}z_{h2} + \beta_{h3}z_{h3}.$$

- Note that we haven't included an interaction term, which in this case implies, that we assume independence between **Eye** and **Hair** in the model.
- Since our response variable now is a count it is no longer a linear model (`lm`) as we have been used to (linear regression).
- Instead it is a so-called generalized linear model and the relevant R command is `glm`.

6.3 Model specification in R

```
model <- glm(Freq ~ Hair + Eye, family = poisson, data = HairEye)
```

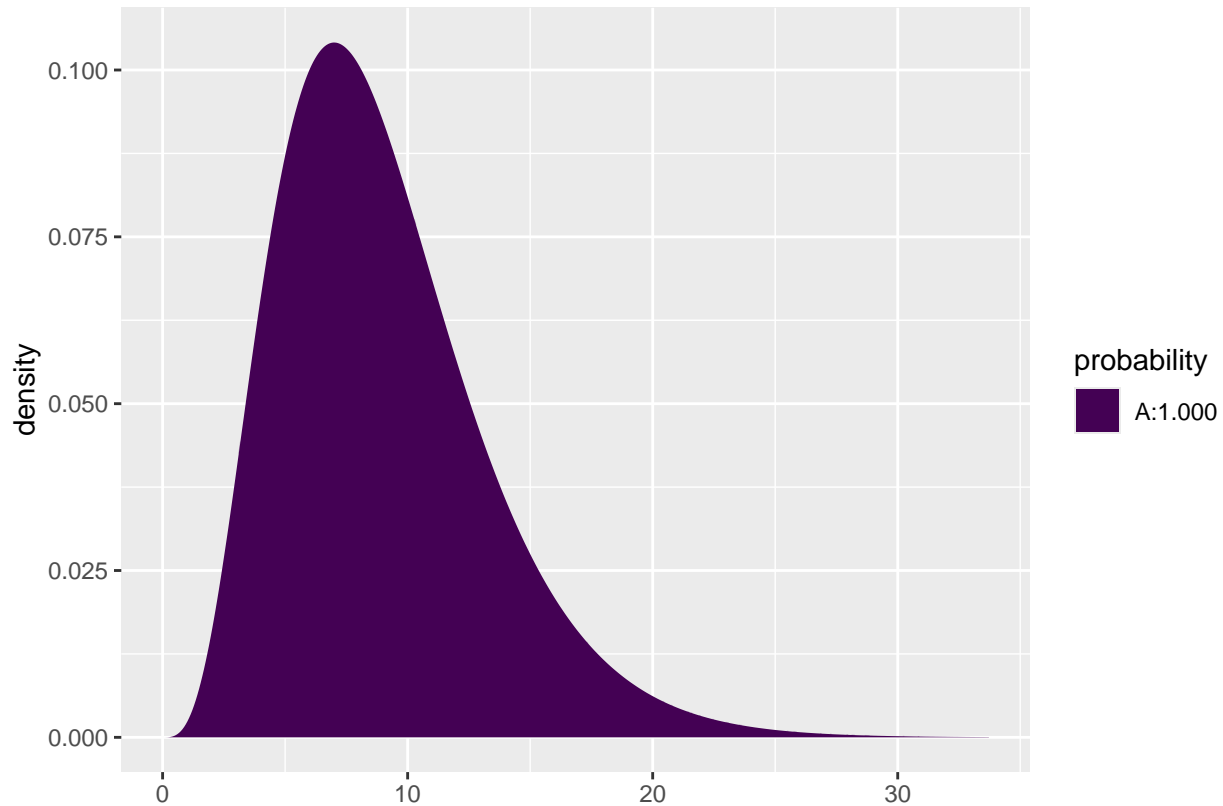
- The argument `family = poisson` ensures that R knows that data should be interpreted as discrete counts and not a continuous variable.

```
summary(model)
```

```
##
## Call:
## glm(formula = Freq ~ Hair + Eye, family = poisson, data = HairEye)
##
## Coefficients:
##             Estimate Std. Error z value Pr(>|z|)
## (Intercept)  3.66926    0.11055  33.191 < 2e-16 ***
## HairBlond    0.16206    0.13089   1.238  0.21569
## HairBrown    0.97386    0.11294   8.623 < 2e-16 ***
## HairRed     -0.41945    0.15279  -2.745  0.00604 **
## EyeBrown     0.02299    0.09590   0.240  0.81054
## EyeGreen    -1.21175    0.14239  -8.510 < 2e-16 ***
## EyeHazel    -0.83804    0.12411  -6.752 1.46e-11 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for poisson family taken to be 1)
##
## Null deviance: 453.31 on 15 degrees of freedom
## Residual deviance: 146.44 on 9 degrees of freedom
## AIC: 241.04
##
## Number of Fisher Scoring iterations: 5
```

- A value of $X^2 = 146.44$ with $df = 9$ shows that there is very clear significance and we reject the null hypothesis of independence between hair and eye color.

```
1 - pdist("chisq", 146.44, df = 9)
```



```
## [1] 0
```

6.4 Expected values and standardized residuals

- We also want to look at expected values and standardized (studentized) residuals.
- The null hypothesis predicts $e^{3.67+0.02} = 40.1$ with brown eyes and black hair, but we have observed 68.
- This is significantly too many, since the standardized residual is 5.86.
- The null hypothesis predicts 47.2 with brown eyes and blond hair, but we have seen 7. This is significantly too few, since the standardized residual is -9.42.

```
HairEye$fitted <- fitted(model)
HairEye$resid <- rstudent(model)
HairEye
```

```
##      Eye Hair Freq fitted resid
## 1  Blue Black   20  39.22 -4.492
## 2  Brown Black   68  40.14  5.856
## 3  Green Black    5  11.68 -2.508
## 4  Hazel Black   15  16.97 -0.583
## 5   Blue Blond   94  46.12  9.368
## 6  Brown Blond    7  47.20 -9.423
## 7  Green Blond   16  13.73  0.719
## 8  Hazel Blond   10  19.95 -2.936
## 9   Blue Brown   84 103.87 -3.437
## 10 Brown Brown  119 106.28  2.151
## 11 Green Brown   29  30.92 -0.511
## 12 Hazel Brown   54  44.93  2.023
```

```
## 13 Blue Red 17 25.79 -2.399
## 14 Brown Red 26 26.39 -0.101
## 15 Green Red 14 7.68 2.368
## 16 Hazel Red 14 11.15 0.961
```

7 Introduction to logistic regression

7.1 Binary response

- We consider a binary response y with outcome 1 or 0. This might be a code indicating whether a person is able or unable to perform a given task.
- Furthermore, we are given an explanatory variable x , which is numeric, e.g. age.
- We shall study models for

$$P(y = 1 | x)$$

i.e. the probability that a person of age x is able to complete the task.

- We shall see methods for determining whether or not age actually influences the probability, i.e. is y independent of x ?

7.2 A linear model

$$P(y = 1 | x) = \alpha + \beta x$$

is simple, but often inappropriate. If β is positive and x sufficiently large, then the probability exceeds 1.

8 Simple logistic regression

8.1 Logistic model

Instead we consider the **odds** that the person is able to complete the task

$$\text{Odds}(y = 1 | x) = \frac{P(y = 1 | x)}{P(y = 0 | x)} = \frac{P(y = 1 | x)}{1 - P(y = 1 | x)}$$

which can have any positive value.

The **logistic model** is defined as:

$$\text{logit}(P(y = 1 | x)) = \log(\text{Odds}(y = 1 | x)) = \alpha + \beta x$$

The function $\text{logit}(p) = \log\left(\frac{p}{1-p}\right)$ - i.e. **log of odds** - is termed **the logistic transformation**.

Remark that log odds can be any number, where zero corresponds to $P(y = 1 | x) = 0.5$. Solving $\alpha + \beta x = 0$ shows that at age $x_0 = -\alpha/\beta$ you have fifty-fifty chance of solving the task.

8.2 Logistic transformation

- The function `logit()` (remember to load `mosaic` first) can be used to calculate the logistic transformation:

```
p <- seq(0.1, 0.9, by = 0.2)
p
```

```
## [1] 0.1 0.3 0.5 0.7 0.9
```

```
l <- logit(p)
l
```

```
## [1] -2.197 -0.847 0.000 0.847 2.197
```

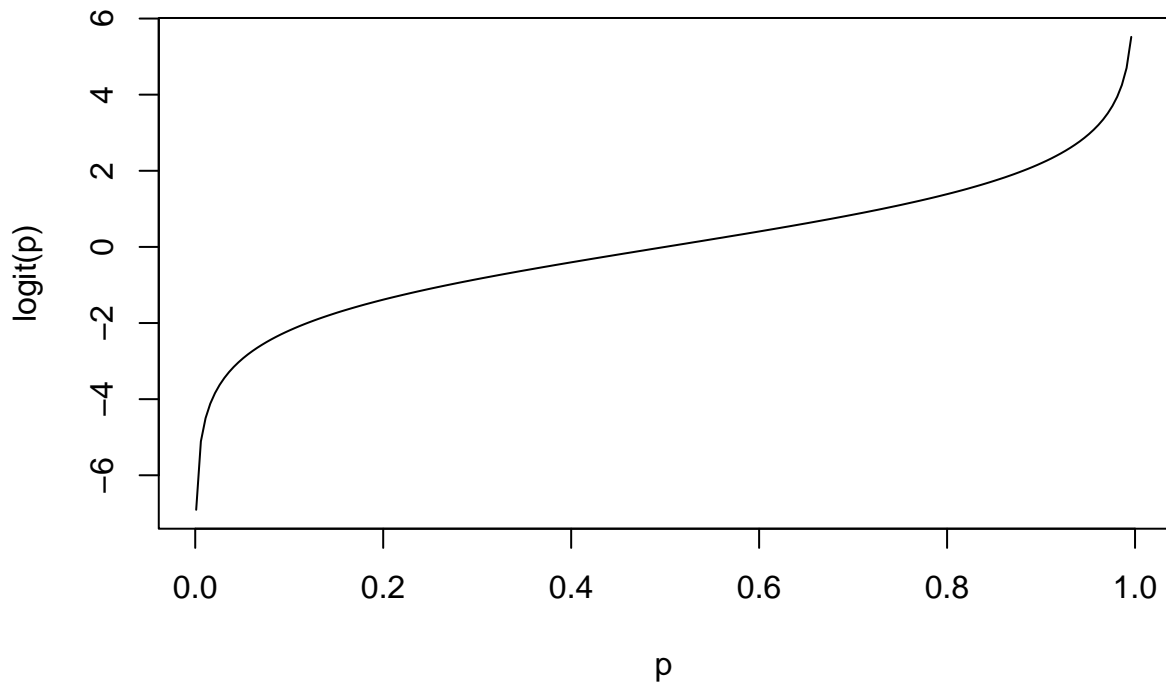
- The inverse logistic transformation `ilogit()` applied to the transformed values can recover the original probabilities:

```
ilogit(1)
```

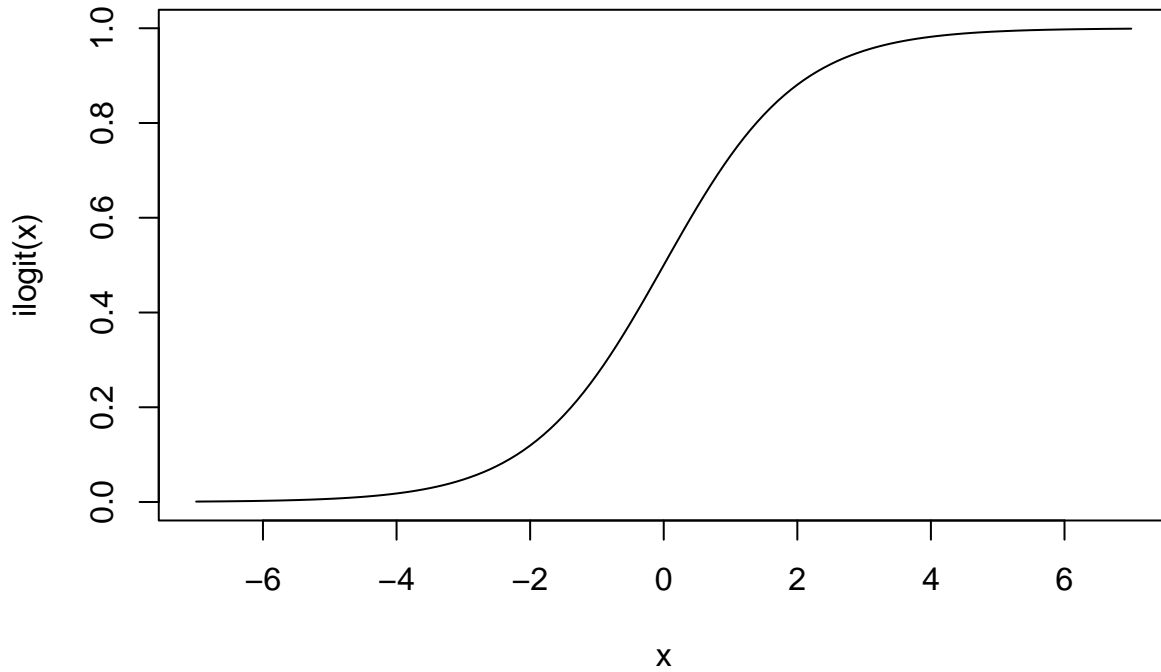
```
## [1] 0.1 0.3 0.5 0.7 0.9
```

Plot of logistic function and inverse logistic

```
p=seq(0.001,0.999,by=0.005)  
plot(p,logit(p),type="l")
```



```
x=seq(-7,7,by= 0.1)  
plot(x,ilogit(x),type="l")
```



8.3 Odds-ratio

Interpretation of β :

What happens to odds, if we increase age by 1 year?

Consider the so-called **odds-ratio**:

$$\frac{\text{Odds}(y = 1 | x + 1)}{\text{Odds}(y = 1 | x)} = \frac{\exp(\alpha + \beta(x + 1))}{\exp(\alpha + \beta x)} = \exp(\beta)$$

where we see, that $\exp(\beta)$ equals the odds for age $x + 1$ relative to odds at age x .

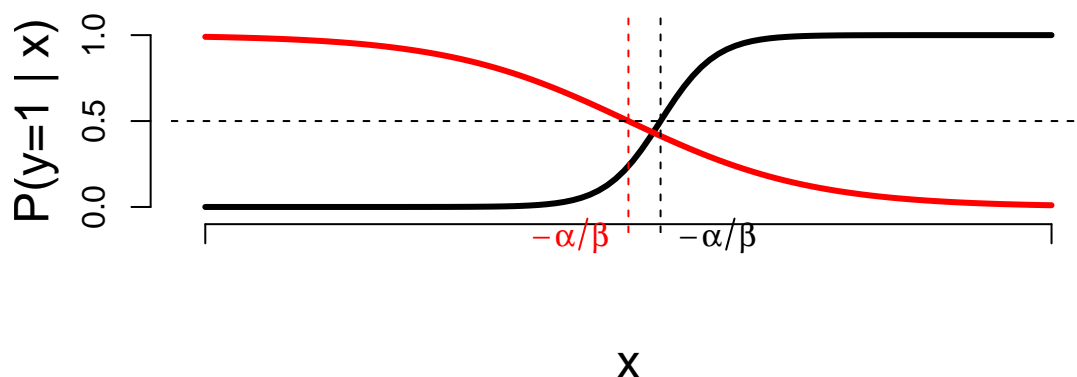
This means that when age increase by 1 year, then the relative change

$$\frac{\exp(\alpha + \beta(x + 1)) - \exp(\alpha + \beta x)}{\exp(\alpha + \beta x)}$$

in odds is given by $100(\exp(\beta) - 1)\%$.

8.4 Simple logistic regression

Logistic curves



Examples of logistic curves for $P(y = 1|x)$. The black curve has a positive β -value ($=10$), whereas the red has a negative β ($=-3$).

In addition we note that:

- Increasing the absolute value of β yields a steeper curve.
- When $P(y = 1|x) = \frac{1}{2}$ then logit is zero, i.e. $\alpha + \beta x = 0$.

This means that at age $x = -\frac{\alpha}{\beta}$ you have 50% chance to perform the task.

8.5 Example: Credit card data

We shall investigate if income is a good predictor of whether or not you have a credit card.

- Data structure: For each level of income, we let n denote the number of persons with that income, and $credit$ how many of these that carries a credit card.

```
creInc <- read.csv("https://asta.math.aau.dk/datasets?file=income-credit.csv")
```

```
head(creInc)
```

```
##   Income  n credit
## 1     12  1     0
## 2     13  1     0
## 3     14  8     2
## 4     15 14     2
## 5     16  9     0
## 6     17  8     2
```

8.6 Example: Fitting the model

```
modelFit <- glm(cbind(credit,n-credit) ~ Income, data = creInc, family = binomial)
```

- `cbind` gives a matrix with two column vectors: `credit` and `n-credit`, where the latter is the vector counting the number of persons without a credit card.
- The response has the form `cbind(credit,n-credit)`.
- We need to use the function `glm` (generalized linear model).

- The argument `family=binomial` tells the function that the data has binomial variation. Leaving out this argument will lead R to believe that data follows a normal distribution - as with `lm`.
- The function `coef` extracts the coefficients (estimates of parameters) from the model summary:

```
coef(summary(modelFit))

##           Estimate Std. Error z value Pr(>|z|)
## (Intercept)  -3.518     0.7103  -4.95 7.33e-07
## Income         0.105     0.0262   4.03 5.58e-05
```

8.7 Test of no effect

```
coef(summary(modelFit))

##           Estimate Std. Error z value Pr(>|z|)
## (Intercept)  -3.518     0.7103  -4.95 7.33e-07
## Income         0.105     0.0262   4.03 5.58e-05
```

Our model for dependence of odds of having a credit card related to income(x) is

$$\text{logit}(x) = \alpha + \beta x$$

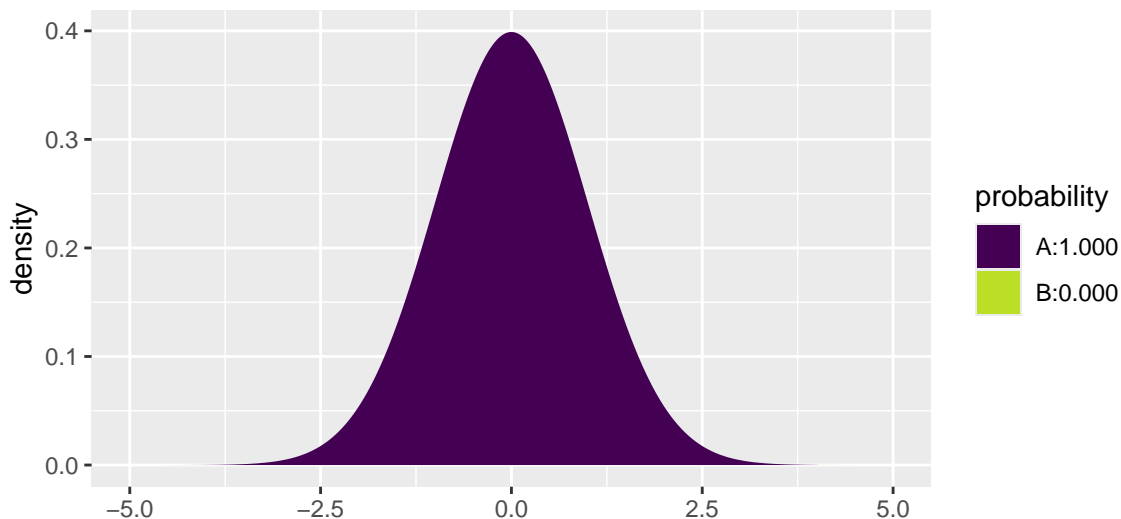
The hypothesis of no relation between income and ability to obtain a credit card corresponds to

$$H_0: \beta = 0$$

with the alternative $\beta \neq 0$. Inspecting the summary reveals that $\hat{\beta} = 0.1054$ is more than 4 standard errors away from zero.

With a z-score equal to 4.03 we get the tail probability

```
ptail <- 2*(1-pdist("norm",4.03,xlim=c(-5,5)))
```



```
ptail
## [1] 5.58e-05
```

Which is very significant - as reflected by the p-value.

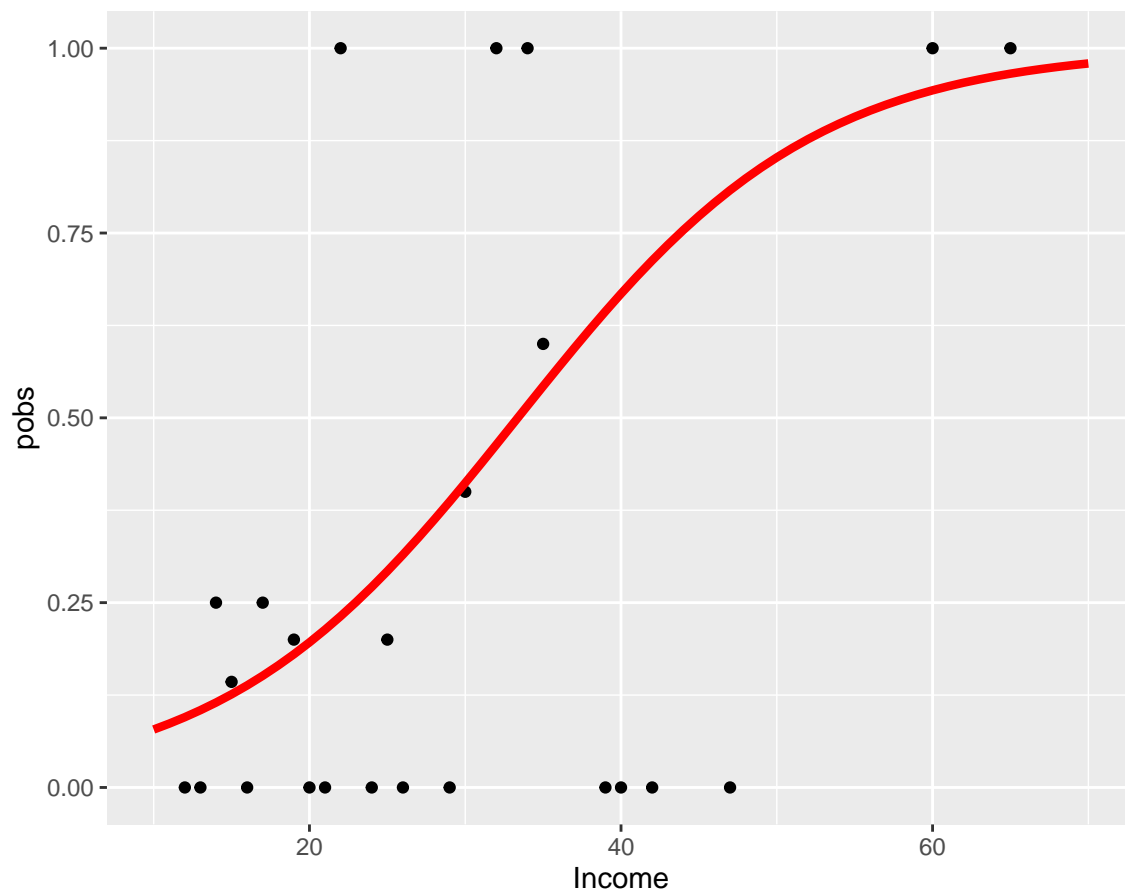
8.8 Confidence interval for odds ratio

From the summary:

- $\hat{\beta} = 0.10541$ and hence $\exp(\hat{\beta}) - 1 = 0.11$. If income increases by 1000 euro, then odds increases by 11%.
- Standard error on $\hat{\beta}$ is 0.02616 and hence a 95% confidence interval for log-odds ratio is $\hat{\beta} \pm 1.96 \times 0.02616 = (0.054; 0, 157)$.
- Corresponding interval for odds ratio: $\exp((0.054; 0, 157)) = (1.056; 1.170)$,
i.e. the increase in odds is - with confidence 95% - between 5.6% and 17%.

8.9 Plot of model predictions against actual data

Expected (red line) and observed (black dots) probabilities



- Tendency is fairly clear and very significant.
- Due to low sample size at some income levels, the deviations are quite large.

9 Multiple logistic regression

9.1 Several numeric predictors

We generalize the model to the case, where we have k predictors x_1, x_2, \dots, x_k . Where some might be dummies for a factor.

$$\text{logit}(P(y = 1 | x_1, x_2, \dots, x_k)) = \alpha + \beta_1 x_1 + \dots + \beta_k x_k$$

Interpretation of β -values is unaltered: If we fix x_2, \dots, x_k and increase x_1 by one unit, then the relative change in odds is given by $\exp(\beta_1) - 1$.

9.2 Example

Wisconsin Breast Cancer Database covers 683 observations of 10 variables in relation to examining tumors in the breast.

- Nine clinical variables with a score between 0 and 10.
- The binary variable `Class` with levels `benign/malignant`.
- By default R orders the levels lexicographically and chooses the first level as reference ($y = 0$). Hence `benign` is reference, and we model odds of `malignant`.

We shall work with only 4 of the predictors, where two of these have been discretized.

```
BC <- read.table("https://asta.math.aau.dk/datasets?file=BC0.dat",header=TRUE)
head(BC)
```

```
##   nuclei cromatin Size.low Size.medium Shape.low   Class
## 1      1         3     TRUE      FALSE     TRUE   benign
## 2     10         3    FALSE     TRUE     FALSE   benign
## 3      2         3     TRUE     FALSE     TRUE   benign
## 4      4         3    FALSE     FALSE    FALSE   benign
## 5      1         3     TRUE     FALSE     TRUE   benign
## 6     10         9    FALSE     FALSE    FALSE malignant
```

9.3 Global test of no effects

First we fit the model `mainEffects` with main effect of all predictors - remember the notation `~ .` for all predictors. Then we fit the model `noEffects` with no predictors.

```
mainEffects <- glm(factor(Class)~., data=BC, family=binomial)
noEffects <- glm(factor(Class)~1, data=BC, family=binomial)
```

First we want to test, whether there is any effect of the predictors, i.e the null hypothesis

$$H_0 : \beta_1 = \beta_2 = \beta_3 = \beta_4 = \beta_5 = 0$$

9.4 Example

Similarly to `lm` we can use the function `anova` to compare `mainEffects` and `noEffects`. Only difference is that we need to tell the function that the test is a chi-square test and not an F-test.

```
anova(noEffects, mainEffects, test="Chisq")
```

```
## Analysis of Deviance Table
##
## Model 1: factor(Class) ~ 1
## Model 2: factor(Class) ~ nuclei + cromatin + Size.low + Size.medium +
##   Shape.low
##   Resid. Df Resid. Dev Df Deviance Pr(>Chi)
## 1      682      884
## 2      677      135  5      749  <2e-16 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

`mainEffects` is a much better model.

The test statistic is the Deviance (749.29), which should be small.

It is evaluated in a chi-square with 5 (the number of parameters equal to zero under the nul hypothesis) degrees of freedom.

The 95%-critical value for the $\chi^2(5)$ distribution is 11.07 and the p-value is in practice zero.

9.5 Test of influence of a given predictor

```
round(coef(summary(mainEffects)),4)
```

```
##           Estimate Std. Error z value Pr(>|z|)
## (Intercept)   -0.709     0.8570  -0.827   0.4080
## nuclei         0.440     0.0823   5.348   0.0000
## cromatin       0.506     0.1444   3.503   0.0005
## Size.lowTRUE  -3.615     0.8081  -4.474   0.0000
## Size.mediumTRUE -2.377     0.7188  -3.307   0.0009
## Shape.lowTRUE  -2.149     0.6054  -3.550   0.0004
```

For each predictor p can we test the hypothesis:

$$H_0 : \beta_p = 0$$

- Looking at the z-values, there is a clear effect of all 5 predictors. Which of course is also supported by the p-values.

9.6 Prediction and classification

```
BC$pred <- round(predict(mainEffects,type="response"),3)
```

- We add the column `pred` to our dataframe `BC`.
- `pred` is the final model's estimate of the probability of malignant.

```
head(BC[,c("Class", "pred")])
```

```
##      Class  pred
## 1  benign 0.011
## 2  benign 0.945
## 3  benign 0.017
## 4  benign 0.929
## 5  benign 0.011
## 6 malignant 1.000
```

Not good for patients 2 and 4.

We may classify by `round(BC$pred)`:

- 0 to denote benign (probability `BC$pred` less than 0.5)
- 1 to denote malignant (probability `BC$pred` more than 0.5)

```
tally(~ Class + round(pred), data = BC)
```

```
##           round(pred)
## Class              0  1
##  benign          433  11
##  malignant        11 228
```

22 patients are misclassified.

```
sort(BC$pred[BC$Class=="malignant"])[1:5]
```

```
## [1] 0.035 0.037 0.089 0.190 0.205
```

There is a malignant woman with a predicted probability of malignancy, which is only 3.5%.

If we assign all women with predicted probability of malignancy above 5% to further investigation, then we only miss two malignant.

```
tally(~ Class + I(pred>.05), data = BC)
```

```
##           I(pred > 0.05)
## Class      TRUE FALSE
##  benign      50   394
##  malignant  237     2
```

The expense is that the number of false positive increases from 11 to 50.

```
tally(~ Class + I(pred>.1), data = BC)
```

```
##           I(pred > 0.1)
## Class      TRUE FALSE
##  benign      27   417
##  malignant  236     3
```

- If we instead set the alarm to 10%, then the number of false positives decreases from 50 to 27.
- But at the expense of 3 false negative.