

Kurs Bio144:

Datenanalyse in der Biologie

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Week 3: Multiple linear regression

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Overview (todo: check)

- Checking the assumptions of linear regression
- Tukey-Anscombe diagram, QQ-plot
- Multiple predictors x_1, x_2, \dots, x_m
- R^2 in multiple linear regression
- t -tests, F -tests and p -values
- Binary and factor covariates

Course material covered today

- Chapters 3.1, 3.2a-q of *Lineare Regression*
- Chapters 4.1 4.2f, 4.3a-e of *Lineare Regression*
- Chapter 11.2 in *Statistische Datenanalyse*

Recap of last week I

- The linear regression model for the data $\mathbf{y} = (y_1, \dots, y_n)$ given $\mathbf{x} = (x_1, \dots, x_n)$ is

$$y_i = \alpha + \beta x_i + e_i, \quad e_i \sim N(0, \sigma_e^2) \text{ independent.}$$

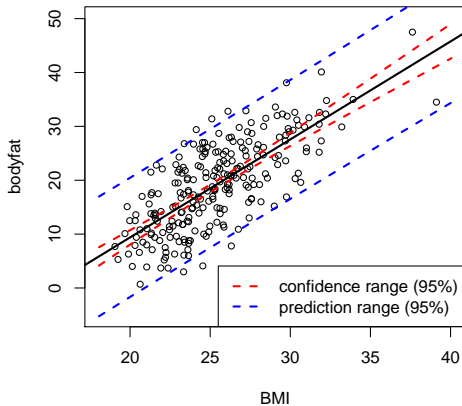
- Estimate the parameters α , β and σ_e^2 by least squares.
- The estimated parameters $\hat{\alpha}$, $\hat{\beta}$ contain **uncertainty** and are normally distributed around the true values.
- Use the knowledge about the distribution to formulate **statistical tests**, such as: Is $\beta = 0$?
- All this is done automatically by R:

```
> summary(r.bodyfat)$coef
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	-26.984368	2.7689004	-9.745518	3.921511e-19
bmi	1.818778	0.1083411	16.787522	2.063854e-42

Recap of last week II

- Confidence and prediction ranges:

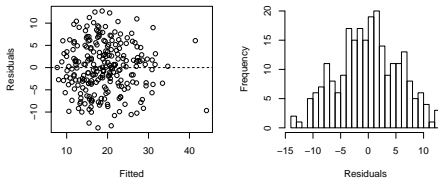


Recap of last week III

Remember: The assumption in linear regression is that the residuals follow a $N(0, \sigma_e^2)$ distribution, implying that :

- a) The expected value of e_i is 0: $E(e_i) = 0$.
- b) All e_i have the same variance: $\text{Var}(e_i) = \sigma_e^2$.
- c) The e_i are normally distributed.
- d) The e_i are independent of each other.

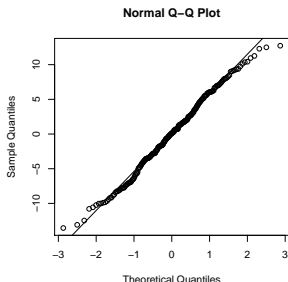
We started to do some residual analysis using the **Tukey-Anscombe plot** and the **Histogram** of the residuals R_i .



Another useful diagnostic plot: The QQ-plot

Usually, not the histogram of the residuals is plotted, but the so-called **quantile-quantile** (QQ) plot. The quantiles of the observed distribution are plotted against the quantiles of the respective theoretical (normal) distribution:

```
> qqnorm(r.bodyfat$residuals)
> qqline(r.bodyfat$residuals)
```

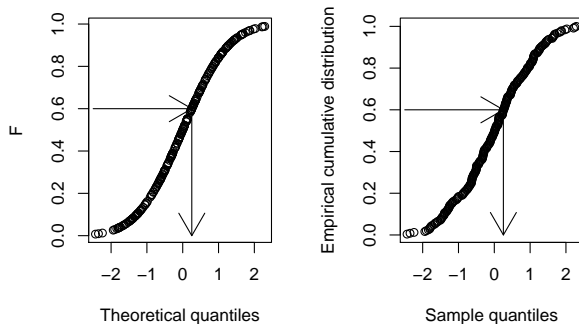


If the points lie approximately on a straight line, the data is fairly normally distributed.

This is often “tested” by eye, and needs some experience.

Please read “Quantil-Quantil-Diagramme”, Chapter 11.2., p.258-261, in “Statistische Datenanalyse” by W. Stahel (Mat183 literature).

It gives a very nice and intuitive description of QQ diagrams!



The idea is that, for each observed point, theoretical quantiles are plotted against the sample quantiles.

Multiple linear regression

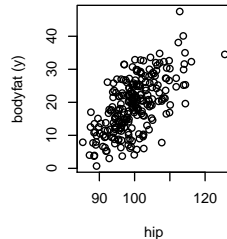
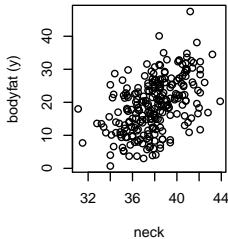
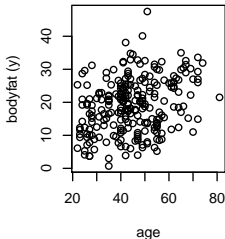
Bodyfat example

We have so far modelled bodyfat in dependence of bmi, that is:

$$(\text{bodyfat})_i = \alpha + \beta \cdot \text{bmi}_i + e_i.$$

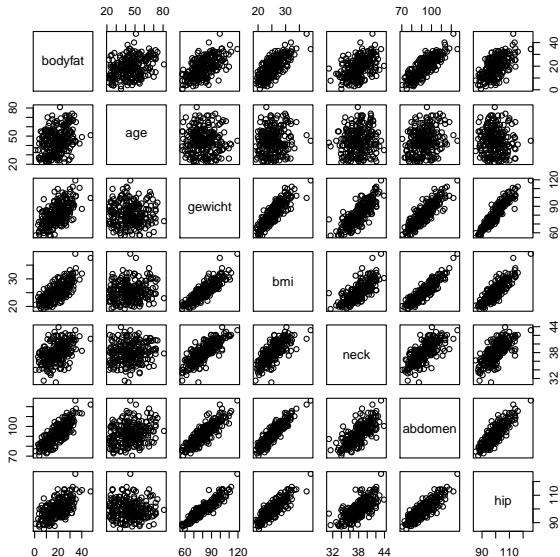
However, other predictors might also be relevant for an accurate prediction of bodyfat.

Examples: Age, neck fat (Nackenfalte), hip circumference, abdomen circumference etc.



Or again the pairs plot:

```
> pairs(d.bodyfat)
```



Multiple linear regression model

The idea is simple: just **extend the linear model by additional predictors**.

- Given several influence factors $x_i^{(1)}, \dots, x_i^{(m)}$, the straightforward extension of the simple linear model is

$$y_i = \beta_0 + \beta_1 x_i^{(1)} + \beta_2 x_i^{(2)} + \dots + \beta_m x_i^{(m)} + e_i$$

with $e_i \sim N(0, \sigma_e^2)$.

- The parameters of this model are $\beta = (\beta_0, \beta_1, \dots, \beta_m)$ and σ_e^2 .

The components of β are again estimated using the **least squares** method. Basically, the idea is (again) to minimize

$$\sum_{i=1}^n r_i^2$$

with

$$r_i = y_i - (\beta_0 + \beta_1 x_i^{(1)} + \beta_2 x_i^{(2)} + \dots + \beta_m x_i^{(m)})$$

It is a bit more complicated than for simple linear regression, see Sections 3.3 and 3.4 of the Stahel script.

Some **linear algebra** is needed to understand these sections, but we do not look into this for the moment. (It will come later in week 6.)

Multiple linear regression for bodyfat

Let us regress the proportion (%) of bodyfat (from last week) on the predictors **bmi** and **age** simultaneously. The model thus is given as

$$\begin{aligned}(bodyfat)_i &= \beta_0 + \beta_1 \cdot bmi_i + \beta_2 \cdot age_i + e_i, \\ \text{with } e_i &\sim N(0, \sigma_e^2) .\end{aligned}$$

Before we estimate the parameters, let us ask the questions that we intend to answer:

- 1 Does the **ensemble** of all covariates explain a relevant part of the variability of the response?
- 2 If yes, which influence variables are good predictors of bodyfat?
- 3 How good is the overall model fit?

Multiple linear regression with R

Let's now fit the model with R, and quickly glance at the output:

```
> r.bodyfatM <- lm(bodyfat ~ bmi + age ,d.bodyfat)
> summary(r.bodyfatM)
```

Call:

```
lm(formula = bodyfat ~ bmi + age, data = d.bodyfat)
```

Residuals:

Min	1Q	Median	3Q	Max
-12.0415	-3.8725	-0.1237	3.9193	12.6599

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	-31.25451	2.78973	-11.203	< 2e-16 ***
bmi	1.75257	0.10449	16.773	< 2e-16 ***
age	0.13268	0.02732	4.857	2.15e-06 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

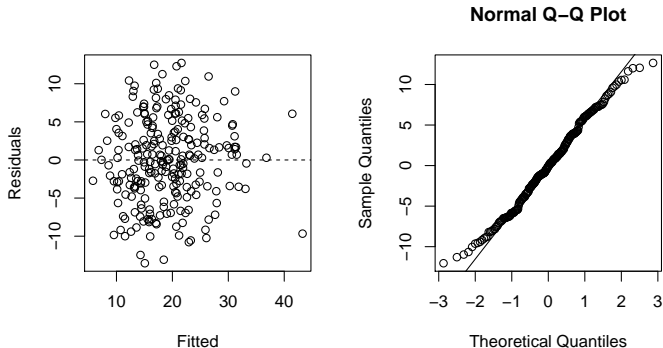
Residual standard error: 5.329 on 240 degrees of freedom

Multiple R-squared: 0.5803, Adjusted R-squared: 0.5768

F-statistic: 165.9 on 2 and 240 DF, p-value: < 2.2e-16

Model checking

Before we look at the results, we have to check if the modelling assumptions are fulfilled:



This seems ok, so continue with answering questions 1-3.

Question 1: Does the model have some explanatory power?

To answer question 1, we need to perform a so-called F -test. The results of the test are displayed in the final line of the regression summary. Here, it says:

F-statistic: 165.9 on 2 and 240 DF, p-value: < 2.2e-16

So apparently (and we already suspected that) the model has some explanatory power.

*The F -statistic and -test is briefly recaptured in 3.1.f) of the Stahel script, but see also Mat183 chapter 6.2.5. It uses the fact that

$$\frac{SSQ^{(R)}/m}{SSQ^{(E)}/(n-p)} \sim F_{m,n-p}$$

follows an F -distribution ($\text{df}()$ in R) with m and $(n-p)$ degrees of freedom, where m are the number of variables, n the number of data points, p the number of β -parameters (typically $m+1$). $SSQ^{(E)} = \sum_{i=1}^n R_i^2$ is the squared sum of the residuals, and $SSQ^{(R)} = SSQ^{(Y)} - SSQ^{(E)}$ with $SSQ^{(Y)} = \sum_{i=1}^n (y_i - \bar{y})^2$.

Question 2: Which variables influence the response?

```
> summary(r.bodyfatM)$coef
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	-31.2545057	2.78973238	-11.203406	1.039096e-23
bmi	1.7525705	0.10448723	16.773060	2.600646e-42
age	0.1326767	0.02731582	4.857137	2.149482e-06

To answer this question, again look at the t -tests, for which the p -values are given in the final column. Each p -value refers to the test for the null hypothesis $\beta_0^{(j)} = 0$ for covariate $x^{(j)}$.

As in simple linear regression, the T -statistic for the j -th covariate is calculated as

$$T_j = \frac{\hat{\beta}_j - \beta_{j0}}{se^{(\beta_j)}} \underbrace{=}_{\text{if } \beta_{j0}=0} \frac{\hat{\beta}_j}{se^{(\beta_j)}}, \quad (1)$$

with $se^{(\beta_j)}$ given in the second column of the regression output.

The distribution of this statistic is $T_j \sim t_{n-p}$.

Therefore: A “small” p -value indicates that the variable is relevant in the model.

Here, we have

- $p < 0.001$ for bmi
- $p < 0.001$ for age

Thus both, bmi and age seem to have some predictive power for bodyfat.

Again, a 95% CI for β_j can be calculated with $[\hat{\beta} - c \cdot \sigma^{(\beta)}; \hat{\beta} + c \cdot \sigma^{(\beta)}]$, where c is the 97.5% quantile of the t -distribution with $n - p$ degrees of freedom (compare to slide 38 of last week). With R:

```
> confint(r.bodyfatM)
```

	2.5 %	97.5 %
(Intercept)	-36.7499929	-25.7590185
bmi	1.5467413	1.9583996
age	0.0788673	0.1864861

!However!:

The p -value and T -statistics should only be used as a **rough guide** for the “significance” of the coefficients.

For illustration, let us extend the model a bit more, including also neck, hip and abdomen:

```
> r.bodyfatM2 <- lm(bodyfat ~ bmi + age + neck + hip + abdomen,d.bodyfat)
> summary(r.bodyfatM2)$coef
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	-7.74964673	7.29830233	-1.0618424	2.893881e-01
bmi	0.42647368	0.23132902	1.8435805	6.649276e-02
age	0.01457356	0.02782994	0.5236649	6.010010e-01
neck	-0.80206081	0.19096606	-4.2000177	3.779800e-05
hip	-0.31764315	0.10751209	-2.9544876	3.447492e-03
abdomen	0.83909391	0.08417902	9.9679702	9.035870e-20

It is now much **less clear** what the influences of age ($p = 0.60$) and bmi ($p = 0.06$) are.

Basically, the problem is that the variables in the model are correlated and therefore explain similar aspects of % bodyfat.

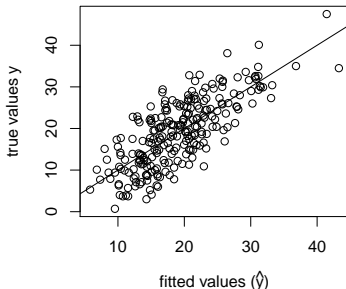
This problem is at the heart of many confusions of regression analysis, and we will talk about such issues later in the course (weeks 7/8).

Question 3: How good is the overall model fit?

To answer this question, we can look at the **multiple R^2** (see Stahel 3.1.h). It is a generalized version of R^2 for simple linear regression:

R^2 for multiple linear regression is defined as the squared correlation between (y_1, \dots, y_n) and $(\hat{y}_1, \dots, \hat{y}_n)$, where the \hat{y} are the fitted values

$$\hat{y}_i = \hat{\beta}_0 + \hat{\beta}_1 x^{(1)} + \dots + \hat{\beta}_m x^{(m)}$$



R^2 is also called the *coefficient of determination* or “Bestimmtheitsmass”, because it measures the proportion of the response's variability that is explained by the ensemble of all covariates:

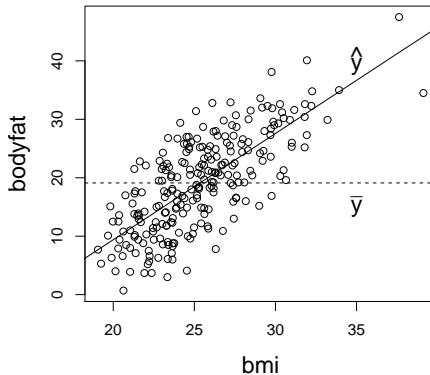
$$R^2 = SSQ^{(R)} / SSQ^{(Y)} = 1 - SSQ^{(E)} / SSQ^{(Y)}$$

Remembering that

total variability = explained variability + residual variability

$$\sum_{i=1}^n (y_i - \bar{y})^2 = \sum_{i=1}^n (\hat{y}_i - \bar{y})^2 + \sum_{i=1}^n (y_i - \hat{y}_i)^2$$

$$SSQ^{(Y)} = SSQ^{(R)} + SSQ^{(E)}$$



Let us look at the R^2 s from the three bodyfat models

(model 1: $y \sim bmi$

model 2: $y \sim bmi + age$

model 3: $y \sim bmi + age + neck + hip + abdomen$):

[1] 0.5390391

[1] 0.5802956

[1] 0.718497

The models thus explain 54 %, 58 % and 72 % of the total variability of y .

It thus *seems* that larger models are “better”. However, R^2 does always increase when new variables are included, but this does not mean that the model is more reasonable.

Model selection is a topic that will be treated in more detail later in this course (week 7).

Adjusted R^2

When the sample size n is small with respect to the number of variables m included in the model, an **adjusted** R^2 gives a better (or “fairer”, i.e. unbiased) estimation of the actual variability that is explained by the covariates:

$$R_a^2 = 1 - (1 - R^2) \frac{n - 1}{n - m - 1}$$

Why R_a^2 ? Todo: explain briefly

Interpretation of the coefficients

Apart from model checking and thinking about questions 1-3, it is probably even **more important to understand what you see**. Look at the output and ask yourself:

What does the regression output actually mean?

	Coefficient	95%-confidence interval	<i>p</i> -value
Intercept	-31.25	from -36.75 to -25.76	< 0.0001
bmi	1.75	from 1.55 to 1.96	< 0.0001
age	0.13	from 0.08 to 0.19	< 0.0001

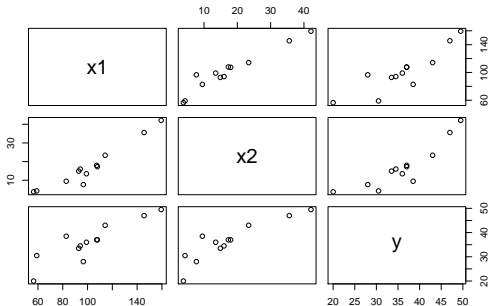
Table : Parameter estimates of model 3.

Task in teams: Interpret the coefficients, 95% CIs and *p*-values.

Example: Catheter Data

Catheter length (y) for heart surgeries depending on two characteristic variables $x^{(1)}$ and $x^{(2)}$ of the patients. Aim: estimate y from $x^{(1)}$ and $x^{(2)}$ ($n = 12$). Again look at the data first:

```
> pairs(d.cath)
```



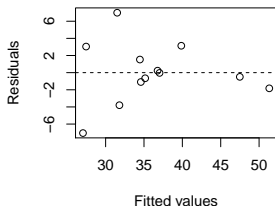
Note that $x^{(1)}$ and $x^{(2)}$ are highly correlated!

Regression results:

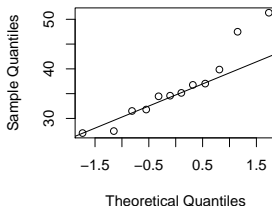
	Coefficient	95%-confidence interval	<i>p</i> -value
Intercept	21.09	from 1.25 to 40.93	0.04
x1	0.077	from -0.25 to 0.40	0.61
x2	0.43	from -0.41 to 1.26	0.28

with $R^2 = 0.81$, $R_a^2 = 0.76$, *p*-value of the *F*-test $p = 0.0006$, and diagnostic residual plots

Tukey–Anscombe



Normal Q–Q Plot



- 1 Are the modelling assumptions fulfilled?
- 2 Does the model have some predictive power?
- 3 Which variable(s) influence(s) the response?
- 4 How good is the overall fit of the model?
- 5 Interpretation of the results?

To understand what is going on, the regression results of y on $x^{(1)}$ and $x^{(2)}$ alone may be useful:

	Coefficient	95%-confidence interval	p -value
Intercept	12.13	from 2.66 to 21.59	0.017
x_1	0.24	from 0.15 to 0.33	0.0002

	Coefficient	95%-confidence interval	p -value
Intercept	25.63	from 21.16 to 30.09	< 0.0001
x_2	0.62	from 0.40 to 0.83	< 0.0001

Binary covariates

So far, the covariates x were always continuous.

However, in our regression models there are **no restrictions assumed with respect to the x variables**.

One very frequent data type of covariates are **binary** variables, that is, variables that can only attain values 0 or 1.

See section 3.2c of the Stahel script:

If the binary variable x is the only variable in the model $y_i = \beta_0 + \beta_1 x_i + e_i$, the model has only two predicted outcomes (plus error):

$$y_i = \begin{cases} \beta_0 + e_i & \text{if } x_i = 0 \\ \beta_0 + \beta_1 + e_i & \text{if } x_i = 1 \end{cases}$$

Example: Smoking variable in Hg Study

For the 59 mothers in the Hg study, check if their smoking status (0=no,1=yes) influences the Hg-concentration in the urin.

We fit the following linear regression model:

$$\log(Hg_{urin})_i = \beta_0 + \beta_1 \cdot x_i^{(1)} + \beta_2 \cdot x_i^{(2)} + \beta_3 \cdot x_i^{(3)} + e_i ,$$

Where

- $\log(Hg_{urin})_i$ is the urine mercury concentration.
- $x^{(1)}$ is the binary smoking indicator (0/1), denoted as **dummy variable**.
- $x^{(2)}$ the number of amalgam fillings.
- $x^{(3)}$ the monthly number of marine fish meals.

(Remember from week 1 that the log of Hg concentrations is needed to obtain useful distributions.)

The results table is given as follows:

	Coefficient	95%-confidence interval	p-value
Intercept	-1.01	from -1.22 to -0.80	< 0.0001
smoking	0.22	from -0.06 to 0.50	0.12
amalgam	0.092	from 0.05 to 0.14	0.0001
fish	0.032	from 0.01 to 0.06	0.015

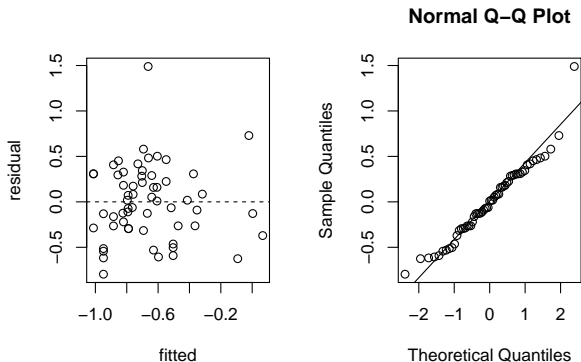
There is some weak ($p = 0.12$) indication that smokers have a slightly increased Hg concentration in their body.

In principle, we have now – at the same time – fitted **two models**: one for smokers and one for non-smokers, assuming that the slopes of the other covariates are the same for both groups.

$$\text{Smokers: } y_i = -1.01 + 0.22 \cdot \text{smoking}_i + 0.092 \cdot \text{amalgam}_i + 0.032 \cdot \text{fish}_i + e_i$$

$$\text{Non-smokers: } y_i = -1.01 + 0.092 \cdot \text{amalgam}_i + 0.032 \cdot \text{fish}_i + e_i$$

...and for completeness, again a short check of the modelling assumptions:



It seems ok, apart from one point that could be categorized as an outlier. We ignore it for the moment.

Factor covariates

Some covariates indicate a **category**, for instance the species of an animal or a plant. This type of covariate is called a **factor**. The trick is to convert a factor with k levels (for instance 3 species) into k dummy variables $x_i^{(j)}$ with

$$x_i^{(j)} = \begin{cases} 1, & \text{if the } i\text{th observation belongs to group } j. \\ 0, & \text{otherwise.} \end{cases}$$

Each of the covariates $x^{(1)}, \dots, x^{(k)}$ can then be included as a binary variable in the model

$$y_i = \beta_0 + \beta_1 x^{(1)} + \dots + \beta_k x^{(k)} + e_i .$$

However: one of the k categories must be selected as a *reference category* and is *not included in the model*, because the model is otherwise not identifiable!! Typically: $\beta_1 = 0$

The model thus discriminates between the factor levels, such that (assuming $\beta_1 = 0$)

$$\hat{y}_i = \begin{cases} \beta_0, & \text{if } x_i^{(1)} = 1 \\ \beta_0 + \beta_2, & \text{if } x_i^{(2)} = 1 \\ \dots & \\ \beta_0 + \beta_k, & \text{if } x_i^{(k)} = 1 \end{cases} .$$

Please also consult Stahel 3.2e.

Example: Earthworm study

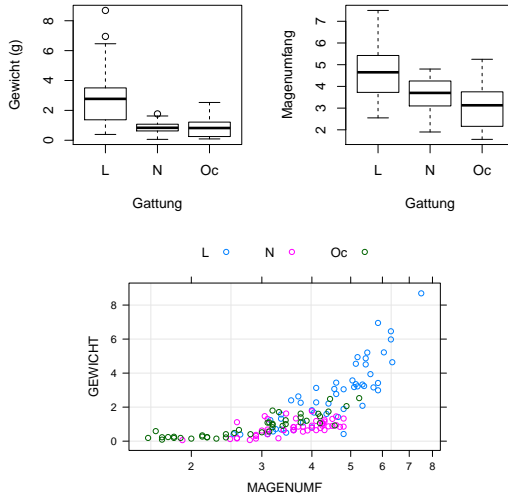
(Angelika von Förster und Burgi Liebst)

Die Dachse im Sihlwald ernähren sich zu einem grossen Prozentsatz von Regenwürmern. Ein Teil des Muskelmagens der Regenwürmer wird während der Passage durch den Dachsdarm nicht verdaut und mit dem Kot ausgeschieden. Wenn man aus der Grösse des Muskelmagenteilchens auf das Gewicht des Regenwurms schliessen kann, ist die Energiemenge berechenbar, die der Dachs aufgenommen hat.

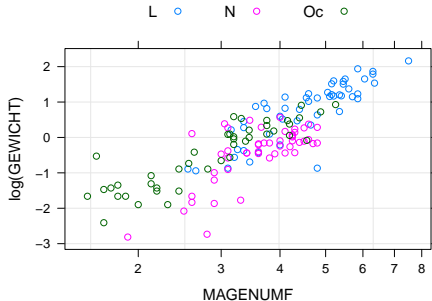
Frage: Besteht eine Beziehung zwischen dem Umfang des Muskelmagenteilchens und dem Gewicht des Regenwurms?

Data set with three species (Lumbricus, Nicodrilus, Octolasion), weight, stomachic circumference (Magenumfang).

Data inspection suggests that the three species have different weight and stomach sizes:



However, data inspection also suggests that there is not really a linear relationship between weight and stomach size. Therefore, log-transform weight, and it looks much better:



Formulate a model (in short notation)

$\log(\text{Gewicht}) \sim \text{Magenumfang} + \text{Gattung}.$

Fitting it is simple in R:

```
> r.lm <- lm(log(GEWICHT) ~ MAGENUMF + Gattung,d.wurm)
> worm.coef <- summary(r.lm)$coef
```

But make sure that **Gattung** is stored as a factor in R (check by `str(d.wurm)`)!

This leads to

	Coefficient	95%-confidence interval	p-value
Intercept	-2.54	from -2.97 to -2.10	< 0.0001
MAGENUMF	0.71	from 0.62 to 0.80	< 0.0001
GattungN	-0.52	from -0.73 to -0.30	< 0.0001
GattungOc	-0.091	from -0.34 to 0.16	0.48

Why is Gattung Lumbricus (L) not in the results table?

Answer: L was chosen as the “reference category”, thus $\beta_L = 0$.

We have now actually fitted **three** models, one model for each species:

Lumbricus: $\hat{y}_i = -2.54 + 0.71 \cdot \text{MAGENUMF}$

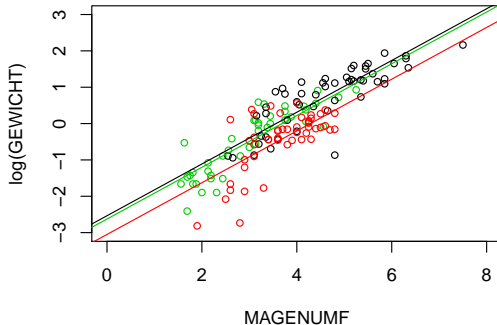
Nicodrilus: $\hat{y}_i = -2.54 + -0.52 + 0.71 \cdot \text{MAGENUMF}$

Octolasion: $\hat{y}_i = -2.54 + -0.09 + 0.71 \cdot \text{MAGENUMF}$

Important: The p -values of the worm species are not very meaningful. They belong to tests that compare the actual level with the reference level. However, the question is whether the species variable has an effect in total.

Plotting the earthworms results

All species have the same slope (this is a modelling assumption), but different intercepts:



F-test to compare models

F When a factor covariate with k levels is in the model, it occupies $k - 1$ parameters. Therefore, the t -test needs to be replaced by the F -test:

F-Test zum Vergleich von Modellen. Die Frage sei, ob die q Koeffizienten $\beta_{j_1}, \beta_{j_2}, \dots, \beta_{j_q}$ in einem linearen Regressionsmodell gleich null sein könnten.

- Nullhypothese: $\beta_{j_1} = 0$ und $\beta_{j_2} = 0$ und ... und $\beta_{j_q} = 0$
- Teststatistik:

$$T = \frac{(\text{SSQ}^{(E)*} - \text{SSQ}^{(E)})/q}{\text{SSQ}^{(E)}/(n-p)};$$

$\text{SSQ}^{(E)*}$ ist die Quadratsumme des Fehlers im „kleinen“ Modell, die man aus einer Regression mit den verbleibenden $m - q$ X -Variablen erhält, und p die Anzahl Koeffizienten im „grossen“ Modell ($= m + 1$, falls das Modell einen Achsenabschnitt enthält, $= m$ sonst).

- Verteilung von T unter der Nullhypothese: $T \sim \mathcal{F}_{q, n-p}$, F -Verteilung mit q und $n - p$ Freiheitsgraden.

Der Test heisst F -Test zum Vergleich von Modellen. Allerdings kann nur ein kleineres Modell mit einem grösseren verglichen werden, in dem alle X -Variablen des kleinen wieder vorkommen, also mit einem „umfassenderen“ Modell. Der früher besprochene F -Test für das gesamte Modell (3.1.e) ist ein Spezialfall: das „kleine“ Modell besteht dort nur aus dem Achsenabschnitt β_0 .

Remember: $F_{1, n-p} = t_{n-p}^2$

F-test for the earthworms

Question: Is the “Gattung” covariate relevant in the model, that is, do the models differ for the three species?

There exists a function (ANOVA) in R that does the F -test for categorical variables:

```
> anova(r.lm)
```

Analysis of Variance Table

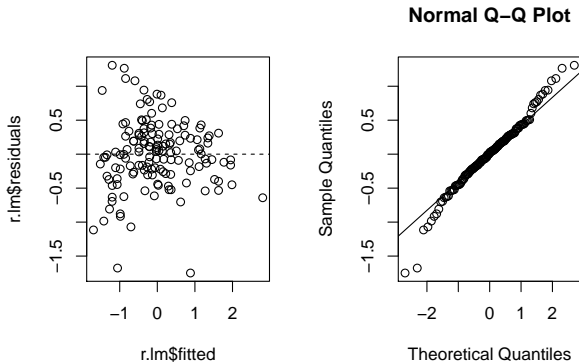
Response: log(GEWICHT)

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
MAGENUMF	1	104.866	104.866	409.69	< 2.2e-16 ***
Gattung	2	7.177	3.589	14.02	2.842e-06 ***
Residuals	139	35.579	0.256		

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Thus: there seems to be some difference in the regression models for the three species.

Again: checking modelling assumptions



Summary

Todo