

# Mixed Effects Models

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- ▶ Days: day 0 is normal sleep baseline (interval, i.e. Numeric)
- ▶ Subject: numbered (categorical, non ordinal, i.e. Factor)

## In R

```
> library(lme4)
```

```
## Loading required package: Matrix
```

```
## Loading required package: Rcpp
```

```
> data(sleepstudy)
```

```
> library(lattice)
```

```
> str(sleepstudy)
```

```
## 'data.frame': 180 obs. of 3 variables:
```

```
## $ Reaction: num 250 259 251 321 357 ...
```

```
## $ Days : num 0 1 2 3 4 5 6 7 8 9 ...
```

```
## $ Subject : Factor w/ 18 levels "308","309","310",...: 1
```

# A quick warning

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- ▶ timeo danaos et dona ferentes!
- ▶ Relax, it'll be okay.



Back to basics

# Statistical Primitives

Basic methods all possible (if tedious) to calculate by hand:

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- ▶ (correlations)

# General linear model

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- ▶  $t$ -test between groups thus fully equivalent to coefficient tests in linear regression
- ▶ ANOVA ( $F$ -test) thus fully equivalent to test  $F$ -test for overall model fit in linear regression

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  - ▶ traditionally separate  $F_1$  (ANOVA by subjects) and  $F_2$  (ANOVA by items) to model crossed random effects
  - ▶ Clark (1973): combine these two tests into a single measure
- ▶ ANOVA sensitive to unbalanced designs and empty cells

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# Impasse

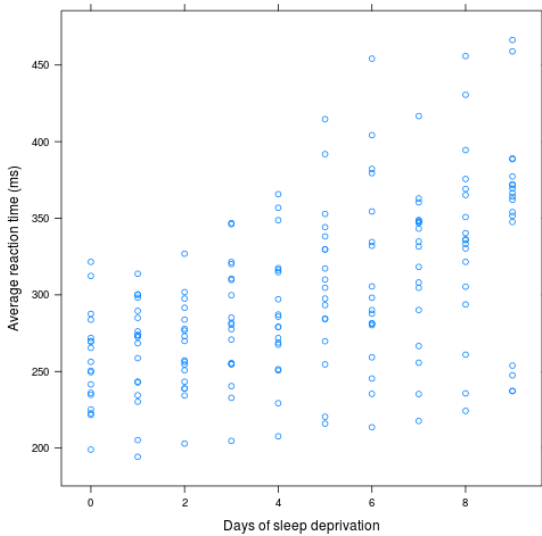
- ▶ all the usual stuff can be expressed as a variant of (generalized) linear regression
- ▶ except repeated measures ANOVA
- ▶ so we have a choice
  - ▶ a detailed, full model with lots of subjects and items
  - ▶ or
  - ▶ ramming everything into a factorial model

What happens if we use linear regression on repeated measures data?



# Linear Regression

```
> # simple scatter plot  
> sleep.xy <- xyplot(Reaction ~ Days, data=sleepstudy,  
+                   xlab = "Days of sleep deprivation",  
+                   ylab = "Average reaction time (ms)")
```



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- ▶  $\text{outcome} = (\text{model}) + \text{error}$

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- ▶ ...

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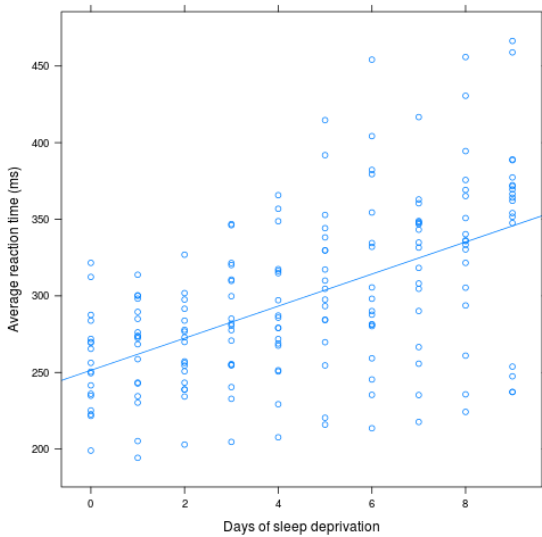
- ▶ Fit a line to observed data with magic and matrices:
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  - ▶  $Y = \beta_3 X + \beta_2 X + \beta_1 X + \beta_0 + \epsilon$
  - ▶ ...
- ▶ R has this built in:
  - > `sleep.lm <- lm(Reaction~Days,data=sleepstudy)`

## Fit a line

- ▶ Fit a line to observed data with magic and matrices:
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  - ▶  $Y = \beta_2 X + \beta_1 X + \beta_0 + \epsilon$
  - ▶  $Y = \beta_3 X + \beta_2 X + \beta_1 X + \beta_0 + \epsilon$
  - ▶ ...
- ▶ R has this built in:
  - > `sleep.lm <- lm(Reaction~Days,data=sleepstudy)`
- ▶ additional predictors with + (no interaction) or \* (interaction)

## Add a regression line with lattice graphics

```
> # p for points, r for regression  
> sleep.xy <- update(sleep.xy, type=c("p", "r"))
```



## Model summary

```
> summary(sleep.lm)

##
## Call:
## lm(formula = Reaction ~ Days, data = sleepstudy)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -110.85  -27.48    1.55   26.14  139.95
##
## Coefficients:
##              Estimate Std. Error t value Pr(>|t|)
## (Intercept)   251.41      6.61    38.03 < 2e-16 ***
## Days          10.47      1.24     8.45 9.9e-15 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 47.7 on 178 degrees of freedom
## Multiple R-squared:  0.286, Adjusted R-squared:  0.282
## F-statistic: 71.5 on 1 and 178 DF, p-value: 9.89e-15
```

Not a great fit!



## Sidebar: ANOVA

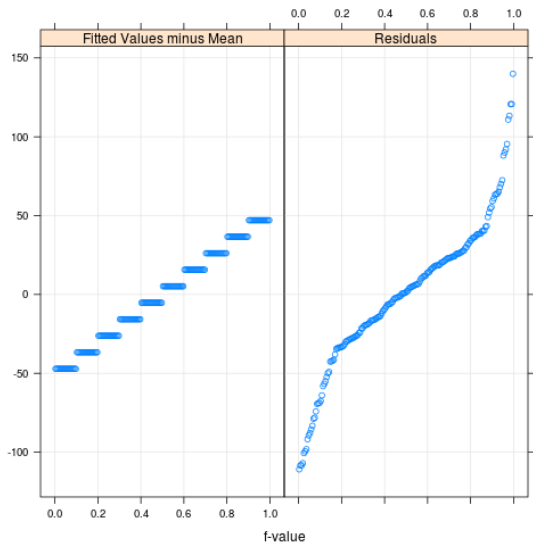
```
> anova(sleep.lm)

## Analysis of Variance Table
##
## Response: Reaction
##           Df Sum Sq Mean Sq F value Pr(>F)
## Days       1 162703  162703    71.5 9.9e-15 ***
## Residuals 178 405252    2277
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

But still not a great fit!

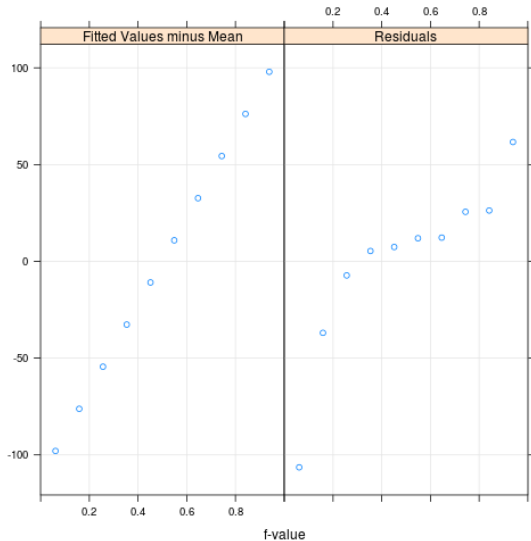
# Residuals for all data

```
> rfs(sleep.lm)
```



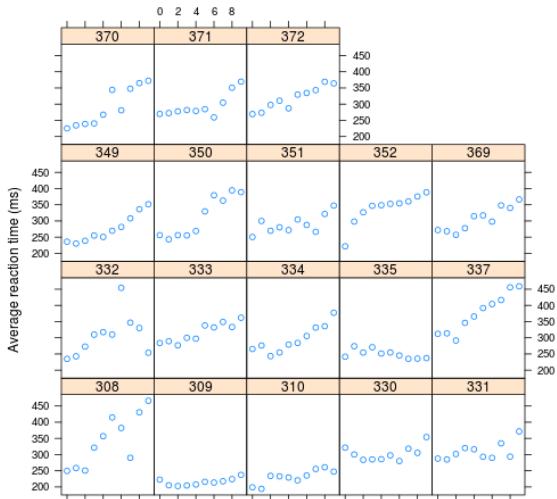
# Residuals for a single subject

```
> sleep.lm.vp1 <- lm(Reaction ~ Days,  
+ data=sleepstudy[sleepstudy$Subject=="308",])  
> rfs(sleep.lm.vp1)
```



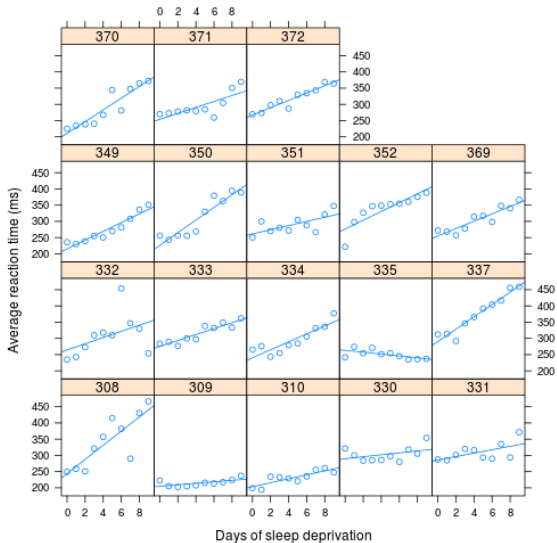
# Models for single subjects

```
> sleep.xy.bysubj <- xyplot(Reaction ~ Days|Subject,  
+ data=sleepstudy,  
+ xlab = "Days of sleep deprivation",  
+ ylab = "Average reaction time (ms)"  
> sleep.xy.bysubj
```



# With regression lines

```
> sleep.xy.bysubj <- update(sleep.xy.bysubj, type=c("p", "r"))  
> sleep.xy.bysubj
```



What do repeated measures actually do to the data?

# Variance and Repeated Measures

- ▶ inter- and intra- variance



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- ▶ random jitter from our choice of sample population

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# Variance and Repeated Measures

- ▶ inter- and intra- variance
- ▶ random jitter from our choice of sample population
- ▶ each subject fulfills a certain “condition”, but random error pro instance of the condition
- ▶ similar idea for item analysis in linguistic designs

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Subjects as fixed effect?

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Subjects as fixed effect?

- ▶ only when we want to make intrasample predictions
- ▶ i.e. sample==population
- ▶ fixed means known variance / manipulation
- ▶ fixed-effects: directed, preferably “exhaustive” manipulation

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## Subjects as random effects?

- ▶ “random” means unknown variance
- ▶ error term is a random effect
- ▶ correction for the error resulting from our particular choice of sample
- ▶ correction per grouping for slope and intercept possible
- ▶ error term per grouping!

# Mixed Effects Models

- ▶ “Mixed” because both fixed random effects are used

```
> sleep.lmer <- lmer(Reaction ~ Days + (1|Subject),  
+                   data=sleepstudy)
```

```
> ?formula
```

# Mixed Effects Models

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- ▶ Same basic formula syntax `dep ~ indep | group`

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- ▶ Same basic formula syntax `dep ~ indep | group`
- ▶ additional `(indep|group)` terms for random effects

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> sleep.lmer <- lmer(Reaction ~ Days + (1|Subject),  
+                   data=sleepstudy)
```

- ▶ More info [here](#), [here](#), and [here](#)

```
> ?formula
```



# Model Summary

```
> summary(sleep.lmer)

## Linear mixed model fit by REML ['lmerMod']
## Formula: Reaction ~ Days + (1 | Subject)
## Data: sleepstudy
##
## REML criterion at convergence: 1786
##
## Scaled residuals:
##   Min      1Q  Median      3Q      Max
## -3.226 -0.553  0.011  0.519  4.251
##
## Random effects:
##   Groups   Name      Variance Std.Dev.
##   Subject (Intercept) 1378     37.1
##   Residual              960     31.0
## Number of obs: 180, groups: Subject, 18
##
## Fixed effects:
##              Estimate Std. Error t value
## (Intercept)  251.405     9.747    25.8
## Days         10.467     0.804    13.0
##
## Correlation of Fixed Effects:
##      (Intr)
## Days -0.371
```

# Model Summary I

```
## Linear mixed model fit by REML ['lmerMod']
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##
## Random effects:
##   Groups      Name                Variance Std.Dev.
##   Subject  (Intercept)           1378      37.1
##   Residual                                960      31.0
## Number of obs: 180, groups:  Subject, 18
##
```

## Model Summary II

## Fixed effects:

##	Estimate	Std. Error	t value
## (Intercept)	251.405	9.747	25.8
## Days	10.467	0.804	13.0

##

## Correlation of Fixed Effects:

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## Days	-0.371

# Fixed effect structure

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- ▶ Package lmerTest
- ▶ Package languageR
- ▶ Package LMERConvenienceFunctions
- ▶ Package lmtest

# Models

```
> sleep.lmer <- update(sleep.lmer, REML=FALSE)
> null <- update(sleep.lmer, . ~ (1|Subject))
```

# Likelihood-ratio test via model comparison

```
> # can only be used for nested models!  
> anova(null,sleep.lmer)  
  
## Data: sleepstudy  
## Models:  
## null: Reaction ~ (1 | Subject)  
## sleep.lmer: Reaction ~ Days + (1 | Subject)  
##           Df  AIC  BIC logLik deviance Chisq Chi Df Pr(>Chisq)  
## null           3 1917 1926  -955    1911  
## sleep.lmer    4 1802 1815  -897    1794   116     1 <2e-16 ***  
## ---  
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

# Random effect structure

- ▶ combine by-subject and by-item analyses in one step

# Random effect structure

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- ▶ cf. Clark (1973)

## Random effect structure

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# Random effect structure

- ▶ Early idea: build up from minimal structure until improvements don't bring you anything on ANOVA (R. Baayen, Davidson, and Bates 2008)
- ▶ New idea: Use the most complicated random effects structure possible (Barr et al. 2013)

# Random effect structure

Possible random effect structures for ONE fixed factor:

1. Intercepts only by random factor:  
(1 | random.factor)

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`(1 | random.factor)`
2. Slopes only by random factor:  
`(0 + fixed.factor | random.factor)`

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`(1 | random.factor)`
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`(0 + fixed.factor | random.factor)`
3. Intercepts and slopes by random factor:  
`(1 + fixed.factor | random.factor)`
4. Intercept and slope, separately, by random factor:  
`(1 | random.factor) + (0 + fixed.factor | random.factor)`

# Models

```
> sleep.lmer.slopes <- update(sleep.lmer,  
+                             . ~ Days + (1+Days|Subject))  
> sleep.lmer.slopes.int <- update(sleep.lmer,  
+                                 . ~ Days + (0+Days|Subject))
```

# Comparing Models

```
> # can only be used for nested models!  
> anova(sleep.lmer,sleep.lmer.slopes, sleep.lmer.slopes.int)  
  
## Data: sleepstudy  
## Models:  
## sleep.lmer: Reaction ~ Days + (1 | Subject)  
## sleep.lmer.slopes.int: Reaction ~ Days + (0 + Days | Subject)  
## sleep.lmer.slopes: Reaction ~ Days + (1 + Days | Subject)  
##  
##          Df  AIC  BIC logLik deviance Chisq Chi Df Pr(>Chisq)  
## sleep.lmer          4 1802 1815  -897    1794  
## sleep.lmer.slopes.int 4 1782 1795  -887    1774  20.0    0 < 2e-16  
## sleep.lmer.slopes    6 1764 1783  -876    1752  22.1    2  1.6e-05  
##  
## sleep.lmer  
## sleep.lmer.slopes.int ***  
## sleep.lmer.slopes      ***  
## ---  
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

# Judging Fit

- ▶ `anova()` function for `lmer()` provided for convenience and parallel to `lm()`



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- ▶ Use `REML=FALSE` when comparing models!

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  - ▶ hard to determine what a significant difference is
  - ▶ tips on AIC
- ▶ Use `REML=FALSE` when comparing models!
- ▶ More advanced techniques for testing in package `pbkrtest`



relationship to AN(C)OVA

## Relationship to ANOVA

- ▶ `ezANOVA()` depends on `aov()` which depends on `lm()`

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- ▶ MEMs work at an individual trial level and can accommodate empty cells and unbalanced designs!
- ▶ `anova()` can be used on individual `lm()`s and `lmer()`s to produce more traditional ANOVA-style output



# ANOVA

```
anova(sleep.lmer.slopes)
```

```
## Analysis of Variance Table
```

```
##      Df Sum Sq Mean Sq F value
```

```
## Days  1  31798   31798    48.5
```

# ANOVA

```
library(car)
Anova(sleep.lmer.slopes)

## Analysis of Deviance Table (Type II Wald chisquare tests)
##
## Response: Reaction
##      Chisq Df Pr(>Chisq)
## Days  48.5  1  3.2e-12 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Wait, where are the  $p$ -values?

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- ▶ so treat  $t$ -values as  $z$  values, which are significant at  $\alpha = 0.05$  when  $|z| > 2$  (cf. R. Baayen, Davidson, and Bates 2008)
- ▶ forget  $p$ -values and traditional notions of significance behind!



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- ▶ this is the foundation of the new statistics (cf. Cummings 2014)
- ▶ after all, “the goal is precision”

When things go wrong

## Possible warning messages

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- ▶ Singular: perfect multicollinearity (at least one variable is linear combination of the others)
- ▶ Not positive definite: matrix not greater than “zero”; too much correlation / collinearity, not enough data

# Generalized linear mixed models (GLMMs)

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- ▶ mixed effects: `glmer()`

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- ▶ Poisson: count ~ continuous
- ▶ (inverse.gaussian, quasi, quasibinomial, quasipoisson)

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  - ▶ performance (cf. Vanrullen 2011)
  - ▶ anomaly detection

## Even more advanced extensions

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- ▶ General additive models extend linear models to arbitrary smooth functions
- ▶ Variants also available for mixed effects: generalized additive mixed models, implemented in R with the `gamm4` package

That's it, but I've added a bunch of further reading after this slide. . .

## (More) References I

- ▶ FAQ from the mailing list (lots to absorb at first, but a good place to keep going back to)





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