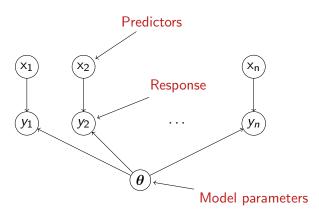
Mixed-effects (a.k.a. multi-level, or hierarchical) models

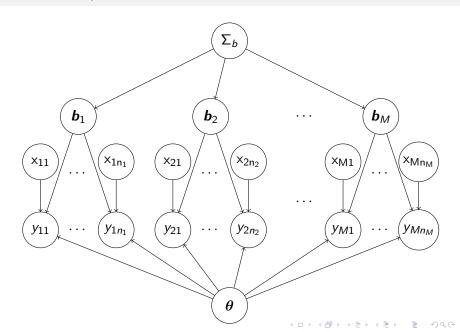
Roger Levy

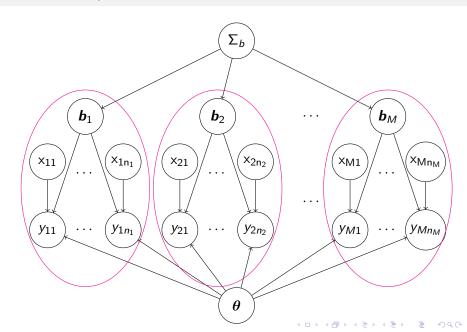
Massachusetts Institute of Technology

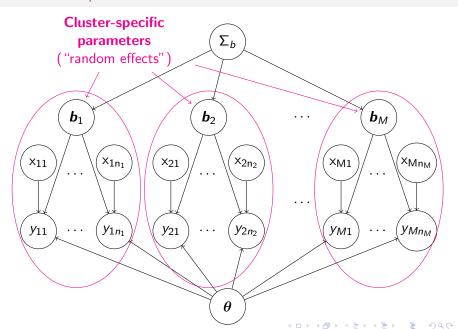
March 12, 2025

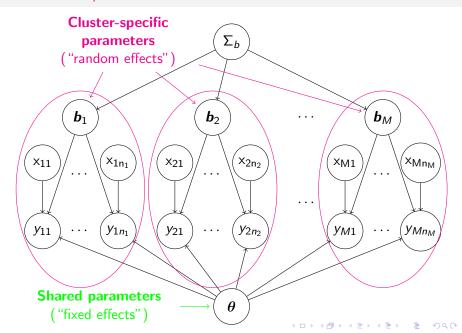
The non-hierarchical GLM picture:

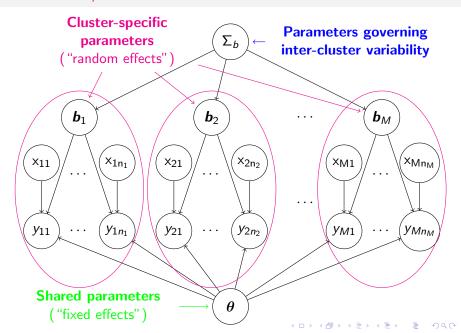












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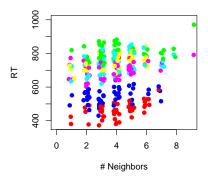
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- One beauty of multi-level models is that you can simulate trial-level data
- ► This is invaluable for achieving deeper understanding of both your analysis and your data

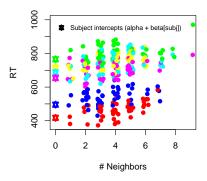
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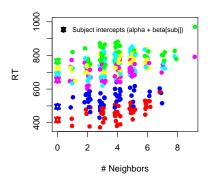
```
## simulate some data
> sigma.b <- 125
                        # inter-subject variation larger than
> sigma.e <- 40
                        # intra-subject, inter-trial variation
> alpha <- 500
> beta <- 12
> M <- 6
                                     # number of participants
> n < -50
                                     # trials per participant
> b <- rnorm(M, 0, sigma.b)</pre>
                                     # individual differences
> nneighbors <- rpois(M*n,3) + 1</pre>
                                     # generate num. neighbors
> subj <- rep(1:M,n)
> RT <- alpha + beta * nneighbors + # simulate RTs!
    b[subj] + rnorm(M*n,0,sigma.e)
                                         4D + 4B + 4B + B + 900
```



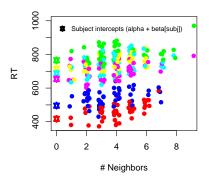
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- Participant-level clustering is easily visible
- ► This reflects the fact that inter-participant variation (125ms) is larger than inter-trial variation (40ms)
- And the effects of neighborhood density are also visible



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- ▶ We have data and we need to infer a model
 - ▶ Specifically, the "fixed-effect" parameters α , β , and σ_{ϵ} , plus the parameter governing inter-subject variation, σ_b
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- Fortunately, we can use the same principles as before to do this:
 - ► The principle of maximum likelihood
 - Or Bayesian inference

```
\sim N(0, \sigma_b) Noise \sim N(0, \sigma_\epsilon)
RT_{ii} = \alpha + \beta x_{ii} + 
> m <- lmer(time ~ neighbors.centered +
    (1 | participant), dat, REML=F)
> print(m,corr=F)
[...]
Random effects:
               Name
                          Variance Std.Dev.
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 Residual
                              19240.5 138.710
Number of obs: 1760, groups: participant, 44
Fixed effects:
                       Estimate Std. Error t value
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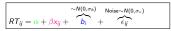
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► What about the participants' idiosyncracies themselves—the *b_i*?

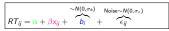
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$$P(b_i|\widehat{\alpha},\widehat{\beta},\widehat{\sigma}_b,\widehat{\sigma}_\epsilon)$$

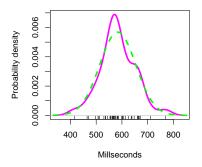


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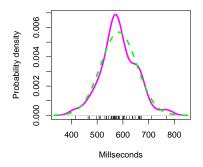
► The BLUPS are the conditional modes of *b_i*—the choices that maximize the above probability

► The BLUP participant-specific "average" RTs for this dataset are black lines on the base of this graph



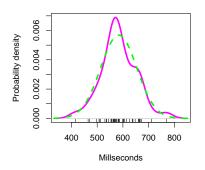
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- ► The solid line is a guess at their distribution
- ► The dotted line is the distribution predicted by the model for the population from which the participants are drawn
- ► Reasonably close correspondence



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Random effects:

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The results of the lmer() fit are saying that the maximum-likelihood estimate of the covariance matrix Σ_b governing participant-specific variability is

$$\widehat{\Sigma_b} = \begin{pmatrix} 70.20 & -0.3097 \\ -0.3097 & 4.41 \end{pmatrix}$$

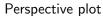


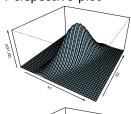
Visualizing some multivariate normal distributions:

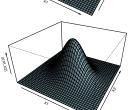
Covariance matrix Perspective plot Contour plot

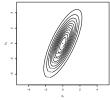
$$\Sigma_b = \begin{pmatrix} 1 & 0.75 \\ 0.75 & 4 \end{pmatrix}$$

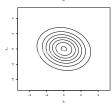
$$\Sigma_b = \begin{pmatrix} 2.5 & -0.13 \\ -0.13 & 2 \end{pmatrix}$$



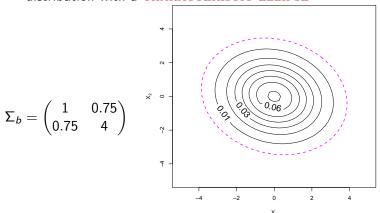




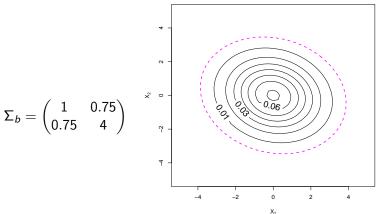




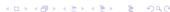
► In 2D we often visually summarize a multivariate normal distribution with a CHARACTERISTIC ELLIPSE

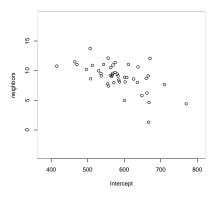


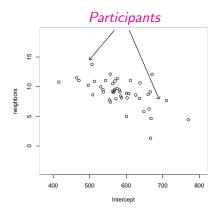
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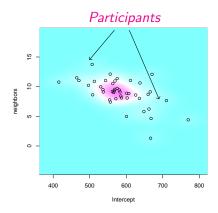


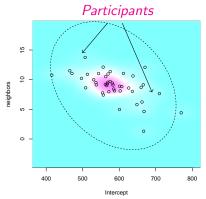
► This ellipse contains a certain proportion (here & conventionally, 95%) of the probability mass for the distribution in question



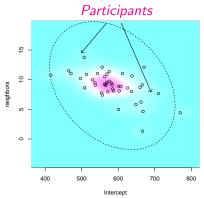








► Correlation visible in participant-specific BLUPs



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- Participants who were faster overall also tend to be more affected by neighborhood density

$$\widehat{\Sigma_b} = \begin{pmatrix} 70.20 & -0.3097 \\ -0.3097 & 4.41 \end{pmatrix}$$



Bayesian inference for multilevel models

$$P(\{\beta_i\}, \sigma_b, \sigma_\epsilon | Y) = \underbrace{\frac{\text{Likelihood}}{P(Y | \{\beta_i\}, \sigma_b, \sigma_\epsilon)} \underbrace{P(\{\beta_i\}, \sigma_b, \sigma_\epsilon)}_{P(Y)}}_{P(Y)} P(\{\beta_i\}, \sigma_b, \sigma_\epsilon)}$$

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Bayesian inference for multilevel models

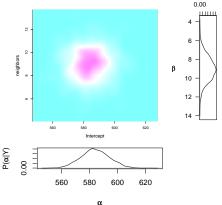
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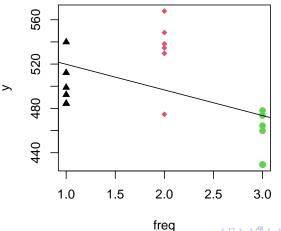
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- There are two situations:
 - 1. When the (average) value that a fixed effect takes *varies* across clusters
 - 2. When the value that a fixed effect takes *varies within some or all clusters*

Predictors varying between clusters

Hypothetical relationship observed for three words:



Predictors varying between clusters



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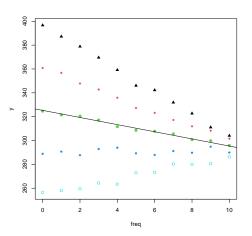
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- Our model will wind up answering the question of whether there is a systematic trend across words for frequency sensitivity, above and beyond idiosyncratic variation among

Predictors varying within clusters

Hypothetical frequency-based responses for five different individual participants:



Predictors varying within clusters I



► It looks like we have good evidence for frequency-sensitivity of the response

Predictors varying within clusters II

- ► Classic question: above and beyond idiosyncratic sensitivities of different individuals to context-driven predictability, are predictable words in general named faster than unpredictable words?
- In mixed-effects models, this implies a need for a random by-speaker slope in our null-hypothesis model
- Inferences about the fixed effect will wind up meaning, is there a systematic effect of word frequency, above and beyond idiosyncratic speaker-specific sensitivities to word frequency?

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```
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- ▶ In the formula syntax of R's 1me4 package:

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- ► Finally, models differing in random effects structure alone can in principle be compared with likelihood-ratio tests
 - However, these results can be either conservative or anti-conservative, so take them with a grain of salt



Results for the nonword-recognition experiment

```
dat$X <- dat$neighbors
m2 <- lmer(time ~ X + (1 + X | participant) + (1|target), dat, REML=F)
## Warning in checkConv(attr(opt, "derivs"), opt$par, ctrl =
control$checkConv, : Model failed to converge with max|grad| =
0.0021026 (tol = 0.002, component 1)
print(m2,corr=F)
## Linear mixed model fit by maximum likelihood ['lmerMod']
## Formula: time ~ X + (1 + X | participant) + (1 | target)
    Data: dat
##
## AIC BIC logLik deviance df.resid
## 22451.93 22490.24 -11218.96 22437.93
                                          1753
## Random effects:
## Groups Name Std.Dev. Corr
## participant (Intercept) 76.140
              X 4.803 -0.46
##
## target (Intercept) 25.485
## Residual 135.763
## Number of obs: 1760, groups: participant, 44; target, 40
## Fixed Effects:
```

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- ► This position has been widely (though not universally) accepted by the field, and we continue to advocate for it

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 - ... were generally arrogant and rude.
- ► The question of theoretical interest for our data is whether the processing penalty induced by disambiguation of the RC attachment would show up immediately (before potentially biasing semantic content of the RC shows up).

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- We'll abbreviate the type of verb (implicit causality or not) the V factor and the RC's attachment level (high or low) the A factor
- ► These factors are crossed in the experiment, and both within-subject. We **sum-code** the two factors to give the main effects the desired interpretation.

Results of a maximal LME fit:

```
## Linear mixed model fit by maximum likelihood ['lmerMod']
## Formula: rt ~ V * A + (V * A | subj) + (V * A | item)
##
     Data: d
##
               BIC logLik deviance df.resid
##
       AIC
  12527.4 12648.0 -6238.7 12477.4
##
## Scaled residuals:
##
      Min 1Q Median 3Q
                                   Max
## -2.0464 -0.5641 -0.1500 0.2521 6.0545
##
## Random effects:
## Groups Name Variance Std.Dev. Corr
   subj (Intercept) 16769.20 129.496
           V
                      476.13 21.820 -0.82
##
                       24.39 4.939 -0.94 0.96
##
          V : A
                      12433.81 111.507 -0.48 0.90 0.75
##
  item (Intercept) 1520.66 38.996
##
           V
                 2055.65 45.339 -0.80
##
                   1663.59 40.787 0.06 -0.64
##
           V · A
                   5531.05 74.371 0.04 0.56 -1.00
##
                      38497.55 196.208
## Residual
## Number of obs: 919, groups: subj, 55; item, 20
##
## Fixed effects:
             Estimate Std. Error t value
## (Intercept) 470.4938 20.5949 22.845
          -33.7621 16.8373 -2.005
## V
            -0.1967 15.9802 -0.012
## A
## V · A
           -85.0056 34.4913 -2.465
## optimizer (nloptwrap) convergence code: 0 (OK)
## boundary (gingular) fit, goo beln(liggingular)
```

Likelihood-ratio-based hypothesis testing for a fixed effect:

Error: bad 'data': object 'd' not found

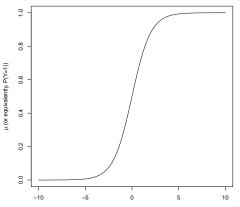
```
rt.lmer.null <- lmer(rt ~ V + A + ( V*A | subj) + ( V*A | item),
    data=d,REML=F)

print(anova(rt.lmer.full,rt.lmer.null))
## Error in h(simpleError(msg, call)): error in evaluating the
argument 'x' in selecting a method for function 'print': object
'rt.lmer.null' not found</pre>
```

```
##
## Attaching package: 'bayesm'
## The following object is masked from
'package:brms':
##
## rdirichlet
```

Recall the inverse logit function that we used for logistic regression:

$$\mu = \frac{e^{\eta}}{1 + e^{\eta}}$$



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$$P(Y=y|\mu) = egin{cases} \mu & y=1 \ 1-\mu & y=0 \ 0 & ext{otherwise} \end{cases}$$

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And linear predictor

$$\eta = X\beta + Zb$$

where b is multivariate-normal distributed:

$$b \sim \textit{N}(0, \Sigma_b)$$



References I

- Barr, D. J., Levy, R., Scheepers, C., and Tily, H. J. (2013). Random effects structure for confirmatory hypothesis testing: Keep it maximal. *Journal of Memory and Language*, 68(3):255–278.
- Bicknell, K., Elman, J. L., Hare, M., McRae, K., and Kutas, M. (2010). Effects of event knowledge in processing verbal arguments. *Journal of Memory and Language*, 63:489–505.
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- Rohde, H., Levy, R., and Kehler, A. (2011). Anticipating explanations in relative clause processing. *Cognition*, 118(3):339–358.

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- Note that so-called "p_{MCMC}" is NOT a p-value in the Neyman-Pearson sense!
- Weakness, both in practice and in principle: the alternative hypothesis is never actually used (except indirectly in determining optimal acceptance and rejection regions)

$$\frac{P(H_0|D)}{P(H_1|D)} = \frac{P(D|H_0)}{P(D|H_1)} \frac{P(H_0)}{P(H_1)}$$

► Alternative: Bayesian hypothesis testing, which is symmetric:

$$\frac{P(H_0|D)}{P(H_1|D)} = \frac{P(D|H_0)}{P(D|H_1)} \frac{P(H_0)}{P(H_1)}$$

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- Technically, such a measure doesn't need to be a true
 Neyman-Pearson p-value (p_{MCMC} falls into this category)

