# Session 5.1: Hierarchical models: longitudinal data

Spatial and Spatio-Temporal Bayesian Models with R-INLA, Imperial College

#### Learning Objectives

After this session you should be able to:

- Specify hierarchical models for longitudinal data
- Distinguish between random intercept and random slope models and recognise when each is more appropriate
- Be able to run the above models in R-INLA

The topics treated in this lecture are covered in Chapter 4 of Gómez-Rubio (2020).

#### Outline

There is huge scope for elaborating the basic hierarchical models discussed in the previous lecture to reflect additional structure and complexity in the data, e.g.

- Adding covariates at different levels of the hierarchy
- Adding further levels to the hierarchy (patients within wards within hospitals, pupils within schools within local authorities, . . .)
- Adding non-nested (cross-classified) levels (patients within GPs crossed with hospitals, . . .)
- Repeated observations on some/all units (longitudinal data we will see it in this lecture)
- Modelling temporal or spatial structure in data, . . . (we will see it from next week)

#### Outline

- 1. What are longitudinal data
- 2. Example: antidepressant clinical trial
- 3. Model specification
- 4. Interpretation

# What are longitudinal data

### What are longitudinal data?

- Arise in studies where individual (or units) are measured repeatedly over time
- For a given individual, observations over time will be typically dependent
- Longitudinal data can arise in various forms:
- continuous or discrete response; discrete response can be binary/binomial, categorical or counts
- equally spaced or irregularly spaced
- same or different time points for each individual
- with or without missing data
- ullet many or few time points, T
- $\bullet$  many or few individuals or units, n

## Analysing longitudinal data

- There are many different ways to analyse longitudinal data
- This is a very big field, so we have to be selective
- The key feature of longitudinal data is the need to account for the dependence structure of the data
- Two common methods:
  - random effects (hierarchical) models
  - autoregressive models
- Here, we will focus on random effects models

Example: sleep study

### Sleep study

- Belenky, Wesensten, Thorne, Thomas, Sing, Redmond, Russo, and Balkin (2003) describes a study of reaction time in patients under sleep deprivation up to 10 days.
- 18 subjects followed for 10 days
- Subjects rated on average reaction time (in ms) for different activities at each measurement

```
> library(lme4)
> data(sleepstudy)
> head(sleepstudy)

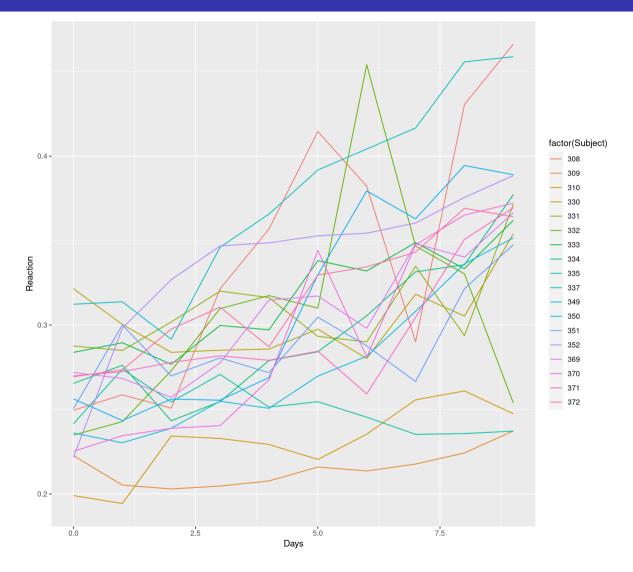
Reaction Days Subject
1 249.5600 0 308
```

```
Reaction Days Subject
1 249.5600 0 308
2 258.7047 1 308
3 250.8006 2 308
4 321.4398 3 308
5 356.8519 4 308
6 414.6901 5 308
```

• Reaction time will be rescaled by dividing by 1000 to have the reaction time in seconds

```
> sleepstudy$Reaction <- sleepstudy$Reaction / 1000</pre>
```

# Sleep Example: data



#### Sleep Example: objective

- Study objective: is the length of sleep deprivation a determinant of reaction time?
- The variables we will use are:
  - -y: Reaction time (in s)
  - -t: Days
- For simplicity we will
  - assume a linear relationship
- The models we will consider are:
  - a non-hierarchical model (standard linear regression) (LM)
  - a hierarchical model with random intercepts (LMM)
  - a hierarchical model with random intercepts and random slopes (LMM2)

# Model specification

### Sleep Example: a Bayesian (non-hierarchical) linear model (LM)

• Specification: 1. probability distribution for responses:

$$y_{it} \sim ext{Normal}(\mu_{it}, \sigma^2)$$

- $-y_{it}$  = the reaction time for individual i on day  $t(\text{days }0,\ldots,9)$
- linear predictor:  $\mu_{it} = \alpha + \beta t$
- -t = the day of the measurement
- 2. In this model no account is taken of the repeated structure (observations are nested within individuals)
- 3. Assume vague priors for all parameters:

$$lpha,eta\sim ext{Normal}(0,10000)$$

## Sleep Example: a Bayesian hierarchical linear model

• Modify LM to allow a separate intercept for each individual:

$$egin{aligned} y_{it} &\sim ext{Normal}(\mu_{it}, \sigma^2) \ \mu_{it} &= rac{oldsymbol{lpha_i}}{2} + eta t \end{aligned}$$

We are assuming that *conditionally* on  $lpha_i$  ,  $\{y_{it}, t=0,\dots,9\}$  are independent

• Assume that all the  $\{\alpha_i\}$  follow a *common* prior distribution, e.g.

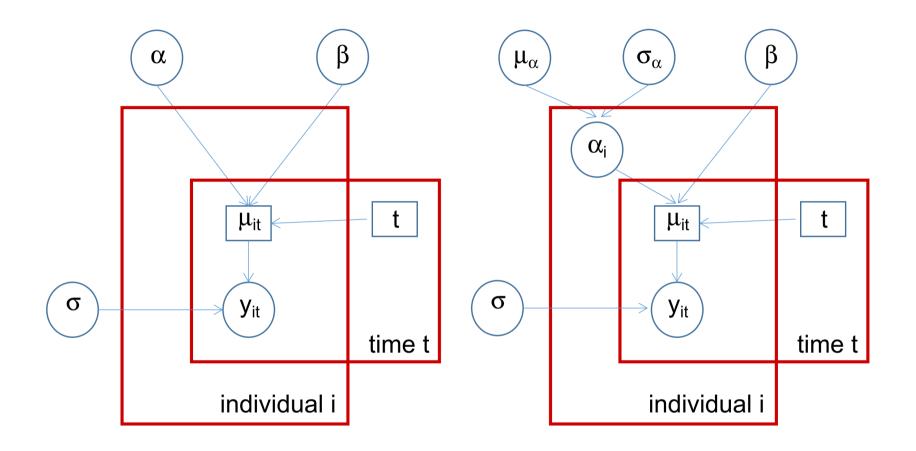
$$lpha_i \sim ext{Normal}(\mu_lpha, \sigma_lpha^2) \quad i = 1, \dots, 246$$

- Here we are assuming exchangeability between all the individuals
- We may then assume vague priors for the *hyperparameters* of the population distribution:

$$\mu_{lpha}{\sim} ext{Normal}(0, 10000) \ rac{1}{\sigma_{lpha}}{\sim} ext{Gamma}(1, 0.001)$$

• This is an example of a *Hierarchical LM* or *Linear Mixed Model (LMM)* or *Random Intercepts* model

## Comparing the two models



(left): LM (right): LMM - random intercept

#### HAMD example: long vs wide format

• In R-INLA is extremely easy to work with longitudinal data, as long as the dataset is in *long format* 

	Long format		Wide f	Wide format	
	Reaction	Days	Subject		
1	0.2495600	0	308		
2	0.2587047	1	308		
3	0.2508006	2	308		
4	0.3214398	3	308		
5	0.3568519	4	308		
6	0.4146901	5	308		
7	0.3822038	6	308		
8	0.2901486	7	308		
9	0.4305853	8	308		
10	0.4663535	9	308		

• Note that a score is present only if the corresponding subject has been observed at that time point

#### HAMD example: long vs wide format

• In R-INLA is extremely easy to work with longitudinal data, as long as the dataset is in *long format* 

```
Wide format
      Long format
   Subject Reaction.0 Reaction.1 Reaction.2 Reaction.3 Reaction.4
       308
            0.2495600
                        0.2587047
                                    0.2508006
                                                0.3214398
                                                           0.3568519
       309
            0.2227339
                        0.2052658
                                    0.2029778
                                                0.2047070
                                                           0.2077161
21
            0.1990539
       310
                        0.1943322
                                    0.2343200
                                                0.2328416
                                                           0.2293074
            0.3215426
                                                0.2851330
31
       330
                        0.3004002
                                    0.2838565
                                                           0.2857973
            0.2876079
41
       331
                        0.2850000
                                    0.3018206
                                                0.3201153
                                                           0.3162773
51
       332
            0.2348606
                        0.2428118
                                    0.2729613
                                                0.3097688
                                                           0.3174629
61
       333
            0.2838424
                        0.2895550
                                    0.2767693
                                                0.2998097
                                                           0.2971710
       334
            0.2654731
                        0.2762012
                                    0.2433647
                                                0.2546723
                                                           0.2790244
81
       335
            0.2416083
                        0.2739472
                                    0.2544907
                                                0.2708021
                                                           0.2514519
                                                0.3461222
91
       337
            0.3123666
                        0.3138058
                                    0.2916112
                                                           0.3657324
```

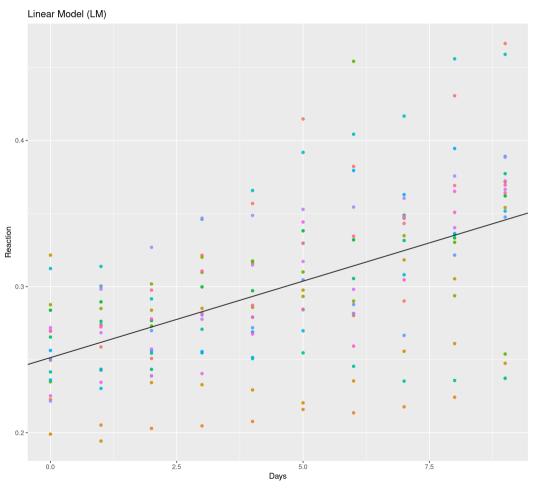
Note that with this format should a subject not have measurement for the entire set of days, R would pad these
out with NAs

#### Sleep example: R-INLA code for LM

Code Plot

## Sleep example: R-INLA code for LM

Code Plot



• all the subjects share the same intercept and slope

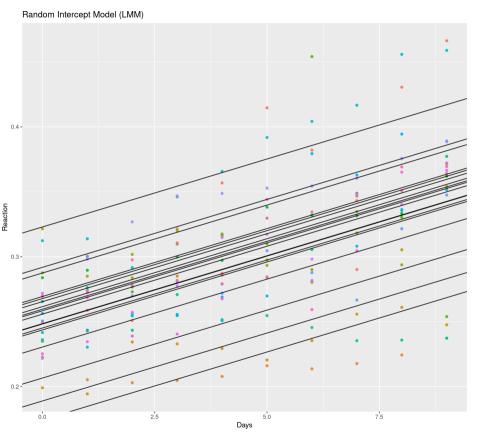
#### Sleep example: R-INLA code for LMM

Code Plot

```
> library(INLA)
> formula_LMM <- Reaction ~ Days + f(Subject,model="iid")
> lmm <- inla(formula_LMM, data=sleepstudy, family="gaussian", control.compute = list(waic=TRUE))
> #Plot
> p <- ggplot(data = sleepstudy, aes(x=Days, y=Reaction, group=Subject))
> p + geom_point(aes(colour = Subject), alpha = .9) + geom_abline(slope=lmm$summary.fixed[2,1], intercept + theme(legend.position = "none")
```

# Sleep example: R-INLA code for LMM

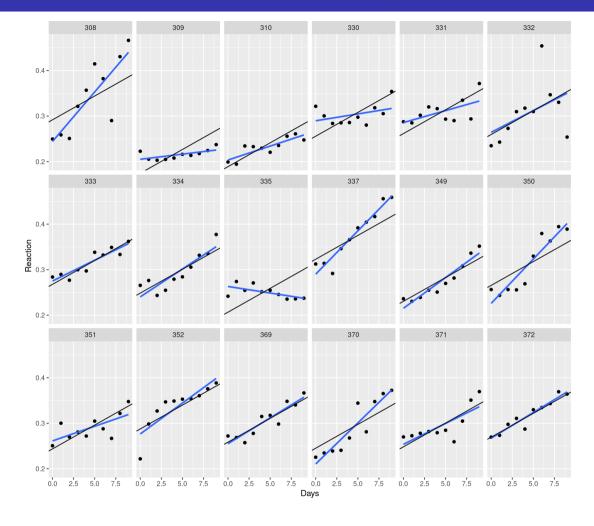
Code Plot



- each individual has a different regression line
- but all individuals have the same slope (parallel lines)

# Interpretation

## Sleep Example: revisiting the data



- blue: regression line for each subject;
- black: regression line from the LMM
- Note that some of the subjects do not fit well in the *parallel lines* scenario
- So we add random slopes to the hierarchical model

## Sleep Example: adding random slopes

• Modify LMM to allow a separate slope for each individual:

$$egin{aligned} y_{it} &\sim ext{Normal}(\mu_{it}, \sigma^2) \ \mu_{it} &= lpha_i + oldsymbol{eta_i t} \end{aligned}$$

• As for the  $\{\alpha_i\}$ , assume that the  $\{\beta_i\}$  follow *common* prior distributions with vague priors on their hyperparameters

#### Sleep Example: adding random slopes

• Modify LMM to allow a separate slope for each individual:

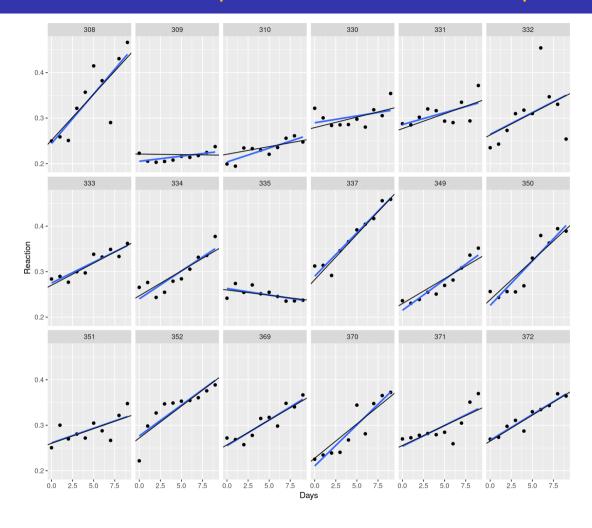
$$egin{aligned} y_{it} &\sim ext{Normal}(\mu_{it}, \sigma^2) \ \mu_{it} &= lpha_i + oldsymbol{eta_i t} \end{aligned}$$

- As for the  $\{\alpha_i\}$ , assume that the  $\{\beta_i\}$  follow *common* prior distributions with vague priors on their hyperparameters
- In R-INLA

```
> Subject2<-sleepstudy$Subject
> formula_LMM2 <- Reaction ~ f(Subject, model="iid") + f(Subject2, Days, model="iid")
> lmm2 <- inla(formula_LMM2, data=sleepstudy, family="gaussian", control.compute = list(waic=TRUE))
> #To access the summaries of the beta_i
> lmm2$summary.random$Subject2[1:5,]
```

```
0.025quant
  ID
                                                 0.5quant 0.975quant
                                                                               mode
                                                                                             kld
              mean
1 308
                                0.015404885
                                             0.0202640016 0.025107546
      0.0202626344 0.002468318
                                                                       0.0202669017 1.423632e-13
2 309
      -0.0002377739 0.002572373 -0.005251515 -0.0002505348 0.004841343 -0.0002767014 4.747123e-13
3 310
      0.0033595434 0.002578983 -0.001667998
                                             0.0033471788 0.008450352 0.0033218474 5.995693e-13
4 330
      0.0045935386 0.002512368 -0.000380402
                                             0.0046067329 0.009491057
                                                                       0.0046344311 7.386043e-13
5 331
      0.0065542981 0.002500534 0.001603901
                                             0.0065671570 0.011430067
                                                                       0.0065941373 7.456124e-13
```

#### HAMD Example: random intercepts and slopes



- LMM with random intercepts only:
  - each individual has a different regression line
  - but for each treatment, only intercept varies by individual
- LMM with random intercepts and random slopes:
  - now intercepts and slopes both vary
  - better fit for each individual

#### References

Belenky, G., N. J. Wesensten, D. R. Thorne, et al. (2003). "Patterns of performance degradation and restoration during sleep restriction and subsequent recovery: A sleep dose-response study". In: *Journal of sleep research* 12.1, pp. 1-12. Gómez-Rubio, V. (2020). *Bayesian inference with INLA*. CRC Press.