

Population and ODE-based models using Stan and Torsten

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StanCon 2019, Cambridge UK
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Outline

1. Course information
2. Introduction and modeling framework | Charles Margossian
3. Models in pharmacometrics | Charles Margossian
4. ODEs in Stan and Torsten | Charles Margossian
5. Numerical ODE integrators | Yi Zhang
6. Population models | Charles Margossian
7. ODE group integrators | Yi Zhang
8. PMX population solvers | Yi Zhang
9. Additional nonlinear ODE example | Charles Margossian

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Instructors

- ▶ Charles Margossian
 - ▶ Columbia University, Department of Statistics
- ▶ Yi Zhang
 - ▶ Metrum Research Group

TA

- ▶ Steve Bronder
 - ▶ Capital One

Outline

Day 1

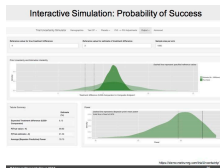
- ▶ Introduction and modeling framework
- ▶ Pharmacometrics models
- ▶ Ordinary differential equation(ODE) based models
- ▶ Numerical ODE integrators

Day 2

- ▶ Population models
- ▶ Group/Population ODE integrators and MPI parallelisation
- ▶ Group/Population solvers and MPI parallelisation

Logistics

METWORX™, cloud-based modeling & simulation platform by Metrum Research Group.



Logistics

Workshop package

- ▶ R scripts and Stan files to do the exercises
- ▶ These slides
- ▶ Outline of the course
- ▶ Additional documentation

We will be using:

- ▶ Torsten v0.87
- ▶ RStan v2.19.2
- ▶ ggplot, plyr, tidyr, dplyr

Outline

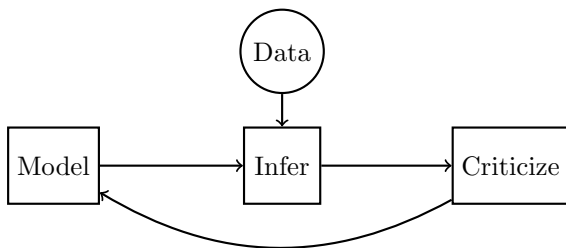
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Preliminary question

- ▶ Why Bayesian in a field such as pharmacometrics?
- ▶ Example - *Bayesian aggregation of average data: an application in drug development* [Weber et al., 2018]

Modeling framework

Box's loop



Inference

- ▶ find the set of parameters consistent with our model and our data
- ▶ approximate this set with draws from the posterior distribution

Sampling algorithm

- ▶ Use the NUTS to sample $\pi(\theta|y)$
- ▶ Requires users the specify $\log \pi(\theta, y) = \log \pi(y|\theta) + \log \pi(\theta)$

The "criticism" step

This step can be broken up in two parts:

1. did we sample from the correct distribution?
2. does our model capture the characteristics of the data we care about?

Diagnosing the inference algorithm

- ▶ look at the trace and the density plots
- ▶ look at \hat{R} and effective number of samples
- ▶ have any warning messages been issued, i.e. divergent transitions ?

Example: fitting a linear model

Likelihood:

$$Y \sim \text{Normal}(\mathbf{x}\beta, \sigma^2)$$

Prior:

$$\beta \sim \text{Normal}(2, 1)$$

$$\sigma^2 \sim \text{Normal}(1, 1)$$

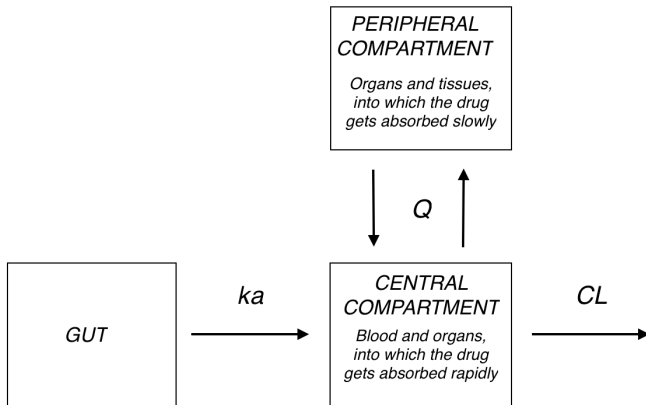
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What is the effect of a treatment on a patient?

- ▶ *pharmacokinetics (PK)*: how is the drug absorbed in the body?
- ▶ *pharmacodynamics (PD)*: once it is absorbed, what are its effects?

Example: Two compartment model



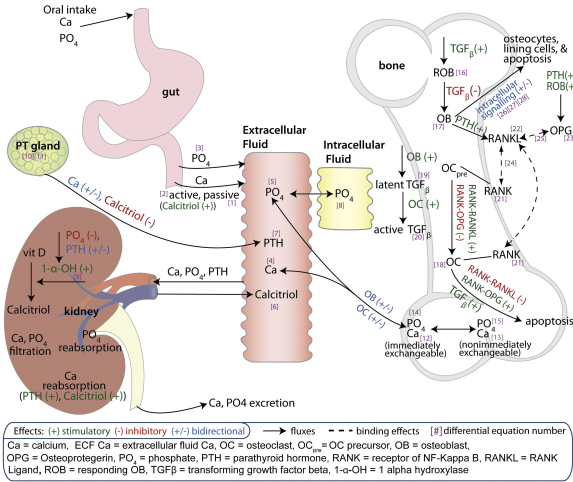
Two compartment model

$$y'_{\text{gut}} = -k_a y_{\text{gut}}$$

$$y'_{\text{cent}} = k_a y_{\text{gut}} - \left(\frac{CL}{V_{\text{cent}}} + \frac{Q}{V_{\text{cent}}} \right) y_{\text{cent}} + \frac{Q}{V_{\text{peri}}} y_{\text{peri}}$$

$$y'_{\text{peri}} = \frac{Q}{V_{\text{cent}}} y_{\text{cent}} - \frac{Q}{V_{\text{peri}}} y_{\text{peri}}$$

Example 2: Bone mineral density model from [Peterson and Riggs, 2012]



Two compartment model

Denote $\theta = \{CL, Q, VC, VP, K_a\}$, the ODE coefficients. Then

$$y' = f(y, t, \theta)$$

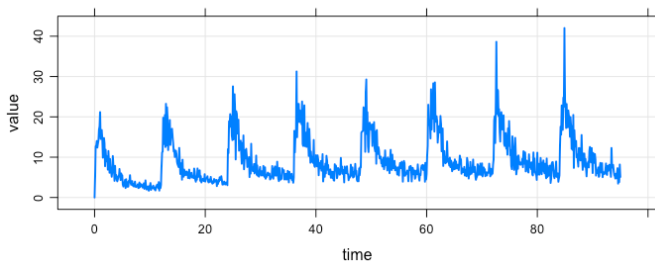
Given an initial condition $y_0 = y(t_0)$, solving the above ODE gives us the *{natural evolution}* of the system at any given time point.

The event schedule

An event can be:

- ▶ **State changer**: an (exterior) intervention that alters the state of the system; for example a bolus dosing or the beginning of an infusion.
- ▶ **Observation**: a measurement of a quantity of interest at a certain time.

Drug concentration in a patient's blood

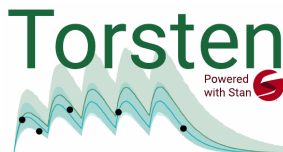


The event schedule

- ▶ There is no general theory for the event schedule :(
- ▶ The modeling software NONMEM® proposes a convention for pharmacometrics, which we adopt in Torsten.

Torsten functions

Torsten functions offers additional built-in functions to simulate data from a compartment model.



Each Torsten function requires users to specify:

- ▶ a system of ODEs and a method to solve it.
- ▶ An event schedule.

Torsten functions

```
matrix = pmx_solve_onecpt(real[] time, real[] amt, real[] rate,  
                           real[] ii, int[] evid, int[] cmt,  
                           real[] addl, int[] ss, real[]  
                               ↪ theta,  
                           real[] biovar, real[] tlag);
```

```
matrix = pmx_solve_twocpt(real[] time, real[] amt, real[] rate,  
                           real[] ii, int[] evid, int[] cmt,  
                           real[] addl, int[] ss, real[]  
                               ↪ theta,  
                           real[] biovar, real[] tlag);
```

- ▶ Analytically solutions for the one/two cpt models.
- ▶ Event schedule
- ▶ ODE coefficients, e.g. $\theta = \{CL, Q, VC, VP, ka\}$ for two-cpt model.
- ▶ bio-availability fraction and lag times.

Example

Clinical trial

- ▶ Single patient
- ▶ Bolus doses with 1200 mg, administered every 12 hours, for a total of 15 doses.
- ▶ Many observations for the first, second, and last doses.
- ▶ Additional observation every 12 hours.

Note: the observation are plasma drug concentration measurement.

See `data/twoCpt.data.r`.

Example

Model

- ▶ two compartment model with first-order absorption
- ▶ prior information based on clinical trial conducted on a large population
- ▶ normal error for the plasma drug concentration measurement.

Example

Prior

```
CL ~ lognormal(log(10), 0.25);  
Q ~ lognormal(log(15), 0.5);  
VC ~ lognormal(log(35), 0.25);  
VP ~ lognormal(log(105), 0.5);  
ka ~ lognormal(log(2.5), 1);  
sigma ~ cauchy(0, 1);
```

Likelihood

$$\log(cObs) \sim \text{Normal} \left(\log \left(\frac{y_2}{VC} \right), \sigma^2 \right)$$

Exercise 1: write and fit this model, using `twoCptModel.r` and `model/twoCptModel.stan`. *Exercise 2:* Write a generated quantities block and do posterior predictive checks.

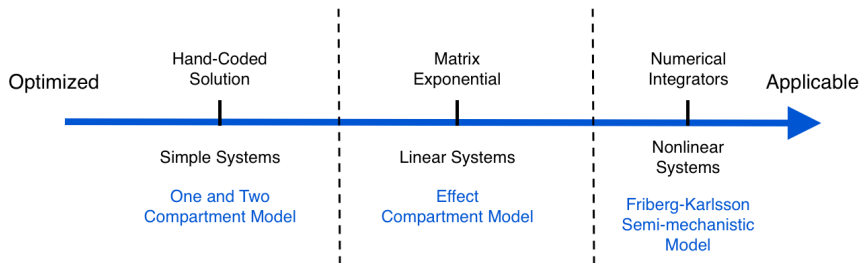
Resources

- ▶ Torsten repository:
`https://github.com/metrumresearchgroup/Torsten`
- ▶ Torsten User manual (on GitHub and in the workshop folder).

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Arsenal of tools



For some examples, see [Margossian and Gillespie, 2017a].

- ▶ the "optimized - applicable" spectrum is a heuristic; counter-examples can be built.
- ▶ coding effort may also be a criterion

Matrix exponential

Consider a system of linear ODEs:

$$y'(t) = Ky(t)$$

where K is a constant matrix.

Then

$$y(t) = e^{tK}y_0$$

Matrix Exponential

$$e^{tK} = \sum_{n=0}^{\infty} \frac{(tK)^n}{n!} = I + tK + \frac{(tK)^2}{2} + \frac{(tK)^3}{3!} + \dots$$

Matrix Exponential

For example, the two compartment model generates the following matrix:

$$K = \begin{bmatrix} -ka & 0 & 0 \\ ka & -(CL + Q)/V_c & Q/V_p \\ 0 & Q/V_c & -Q/V_p \end{bmatrix}$$

Linear ODE solver in Torsten

```
matrix = pmx_solve_linode(real[] time, real[] amt, real[] rate,  
                           real[] ii, int[] evid, int[] cmt,  
                           real[] addl, int[] ss,  
                           matrix K, real[] biovar, real[] tlag)
```

Numerical integrator

```
real[ , ] pmx_integrate_ode_rk45(ODE_RHS, real[] y0, real t0,  
  ↪ real[] ts, real[] theta, real[] x_r, int[] x_i, real rtol =  
  ↪ 1.e-6, real atol = 1.e-6, int max_step = 1e6);
```

- ▶ ODE_RHS: ODE right-hand-side f in $y' = f(y, t, \theta, x_r, x_i)$.
- ▶ y0: initial condition at time t0.
- ▶ t0: initial time.
- ▶ ts: times at which we require a solution.
- ▶ theta: parameters to be passed to f .
- ▶ x_r: real data to be passed to f .
- ▶ x_i: integer data to be passed to f .
- ▶ rtol, atol, and max_step are optional control parameters for *relative tolerance*, *absolute tolerance*, and *max number of time steps*, respectively. Their default values have no theoretical justification.

System function

```
functions {  
  real[] system(real time, real[] y,  
                real[] theta, real[] x_r, int[] x_i) {  
    real dydt[3];  
    real CL = theta[1];  
    real Q = theta[2];  
  
    /* .... */  
  
    return dydt;  
  }  
}
```

Torsten function

```
matrix pmx_solve_rk45(ODE_system, int nCmt, real[] time, real[]  
  ↪ amt, real[] rate, real[] ii, int[] evid, int[] cmt, real[]  
  ↪ addl, int[] ss, real[] theta, real[] biovar, real[] tlag,  
  ↪ real rel_tol, real abs_tol, int max_step);
```

Exercise 3: Write, fit, and diagnose the two compartment model using the `pmx_solve_rk45` function.

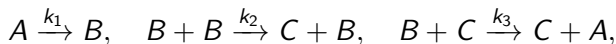
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Nonlinear ODEs without analytical solution

kinetics of an autocatalytic reaction [Robertson, 1966]

The structure of the reactions is



where k_1 , k_2 , k_3 are the rate constants and A , B and C are the chemical species involved. The corresponding ODEs are

$$x_1' = -k_1 x_1 + k_3 x_2 x_3$$

$$x_2' = k_1 x_1 - k_2 x_2^2 - k_3 x_2 x_3$$

$$x_3' = k_2 x_2^2$$

Given $k_1 = 0.04$, $k_2 = 3.0e7$, $k_3 = 1.0e4$, we make inference regarding the initial condition for $x_1(t = 0)$.

Nonlinear ODEs without analytical solution

$$x_1' = -k_1 x_1 + k_3 x_2 x_3$$

$$x_2' = k_1 x_1 - k_2 x_2^2 - k_3 x_2 x_3$$

$$x_3' = k_2 x_2^2$$

Given $k_1 = 0.04$, $k_2 = 3.0\text{e}7$, $k_3 = 1.0\text{e}4$, we make inference regarding the initial condition for $x_1(t = 0)$.

Exercise 4

Write Stan function for the above ODE's RHS.

Stan function for autocatalytic kinetics

$$x_1' = -k_1 x_1 + k_3 x_2 x_3$$

$$x_2' = k_1 x_1 - k_2 x_2^2 - k_3 x_2 x_3$$

$$x_3' = k_2 x_2^2$$

```
functions{  
  real[] reaction(real t, real[] x, real[] theta, real[] r,  
    ↪ int[] i){  
    real dxdt[3];  
    real k1 = theta[1];  
    real k2 = theta[2];  
    real k3 = theta[3];  
    dxdt[1] = -k1*x[1] + k3*x[2]*x[3];  
    dxdt[2] = k1*x[1] - k3*x[2]*x[3] - k2*(x[2])^2;  
    dxdt[3] = k2*(x[2])^2;  
    return dxdt;  
  }  
}
```

- What's the initial conditions for x_2 and x_3 ?

Numerical integrators

- ▶ Runge-Kutta 4th/5th (rk45)
 - ▶ non-stiff equations
 - ▶ Most popular, try this if you don't know the nature of the ODE, or what you're doing, or both.
- ▶ Backward differentiation formula (bdf)
 - ▶ stiff equations
 - ▶ More expensive to use
- ▶ Adams-Moulton
 - ▶ non-stiff equations
 - ▶ higher-order of accuracy(do you really need it?)
 - ▶ scales better with number of steps

Numerical integrators

Integrators	Stan	Torsten
rk45	integrate_ode_rk45	pmx_integrate_ode_rk45
BDF	integrate_ode_bdf	pmx_integrate_ode_bdf
Adams	integrate_ode_adams	pmx_integrate_ode_adams

```
real[ , ] pmx_integrate_ode_rk45(ODE_RHS, real[] y0, real t0,  
  ↪ real[] ts, real[] theta, real[] x_r, int[] x_i, real rtol =  
  ↪ 1.e-6, real atol = 1.e-6, int max_step = 1e6);
```

- ▶ ODE_RHS: ODE right-hand-side f in $y' = f(y, t, \theta, x_r, x_i)$.
- ▶ y0: initial condition at time t0.
- ▶ t0: initial time.
- ▶ ts: times at which we require a solution.
- ▶ theta: parameters to be passed to f .
- ▶ x_r: real data to be passed to f .
- ▶ x_i: integer data to be passed to f .

Model

- ▶ In each of 8 experiments performed x_3 is observed.
- ▶ Hierarchical model for $x_0[1]$

```
model {  
  y0_mu ~ lognormal(log(2.0), 0.5);  
  for (i in 1:nsub) {  
    y0_1[i] ~ lognormal(y0_mu, 0.5);  
  }  
  sigma ~ cauchy(0, 0.5);  
  obs ~ lognormal(log(x3), sigma);  
}
```

Data

Data available for the inference

```
data {  
  int<lower=1> nsub;      /* nb. of subjects */  
  int<lower=1> len[nsub]; /* nb. of results-extraction time  
    ↪ points for each subject */  
  int<lower=1> ntot;      /* total nb. of results-extraction  
    ↪ time points */  
  real ts[ntot];          /* concatenated array for  
    ↪ results-extraction time points */  
  real obs[ntot];         /* concatenated array for observed  
    ↪ x3 */  
}
```

Exercise 5

Given above data and model, write the rest of Stan code.

- ▶ Hint: use `chem.stan` as template, also see `chem.data.R` and `chem.init.R`.
- ▶ Reaction begins with A (on which is also what we'd like to make inference), the other two species are non-existent at the beginning of the reaction.
- ▶ Which numerical integrator are you using? Why?

Exercise 5

How to build & run?

Edit/Add cmdstan/make/local

```
TORSTEN_MPI = 1  # flag on torsten's MPI solvers  
CXXFLAGS += -isystem /usr/local/include    # path to MPI  
↪      library's headers
```

Build in cmdstan

```
make path_to_workshop/RScript/model/chemical_reactions/chem
```

Run

```
./chem sample adapt delta=0.95 random seed=1104508041 data  
↪   file=chem.data.R init=chem.init.R
```

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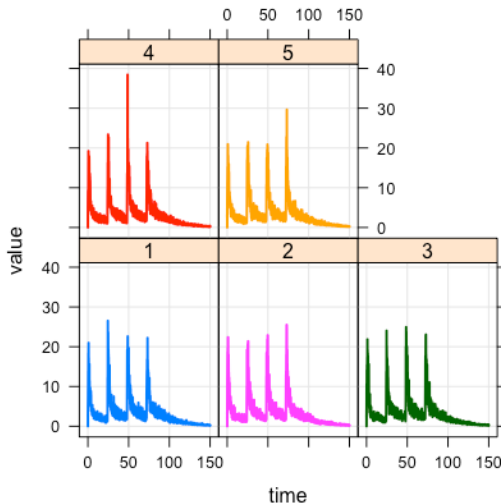
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Data pooled into groups

- ▶ sport measurements are grouped by players
- ▶ people's voting intention can be grouped by states, social status, etc.
- ▶ medical measurements are grouped by patients

Data pooled into groups

- ▶ medical measurements are grouped by patients
 - ▶ Simulated with mrgsolve <https://mrgsolve.github.io/>



Hierarchical model

With a hierarchical model, we can

- ▶ do partial pooling.
- ▶ estimate how similar the groups are to one another.
- ▶ estimate individual parameters.

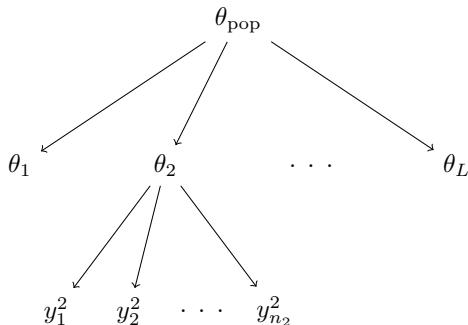
$$\theta = (\theta_1, \dots, \theta_L) \sim p(\theta | \theta_{\text{pop}})$$

$$y = (y_1, \dots, y_N) \sim p(y | \theta, x)$$

Hierarchical model

$$\theta = (\theta_1, \dots, \theta_L) \sim p(\theta | \theta_{\text{pop}})$$

$$y = (y_1, \dots, y_N) \sim p(y | \theta, x)$$



Example 3: Hierarchical two compartment model

Likelihood function:

$$\log \theta \sim \text{Normal}(\log \theta_{\text{pop}}, \Omega)$$

$$\Omega = \begin{pmatrix} \omega_1 & 0 & 0 & 0 & 0 \\ 0 & \omega_2 & 0 & 0 & 0 \\ 0 & 0 & \omega_3 & 0 & 0 \\ 0 & 0 & 0 & \omega_4 & 0 \\ 0 & 0 & 0 & 0 & \omega_5 \end{pmatrix}$$

$$\log(cObs) \sim \text{Normal} \left(\log \left(\frac{y_2}{VC} \right), \sigma^2 \right)$$

Exercise 6: Write, fit, and diagnose a hierarchical two

compartment model for a population of 10 patients. Use
`data/twoCptPop.data.r` and `twoCptPop.r` }

- ▶ *Start by running 3 chains with 30 iterations.*
- ▶ *Do you get any warning messages?*

Divergent transitions

- ▶ *Do you get any warning messages?*

There were 29 divergent transitions after warmup.

- ▶ A divergent transition occurs when we fail to accurately compute a Hamiltonian trajectory.
- ▶ This is because we *approximate* trajectories.
- ▶ Our sampler may not be refined enough to explore the entire typical set.

Divergent transitions

Consider the following hierarchical model:

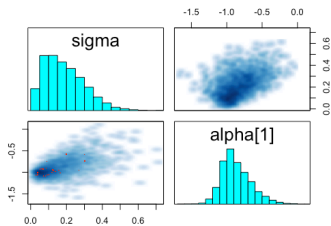
$$\alpha_i \sim \text{Normal}(\mu, \sigma)$$

$$y_i \sim p(y|\alpha_i)$$

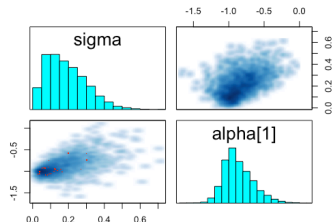
Divergent transitions

$$\alpha_i \sim \text{Normal}(\mu, \sigma)$$

Fitting this model yields the following pairs plot:



Divergent transitions



- ▶ This geometric shape is known as Neil's funnel [Neil, 2003].
- ▶ Its interactions with HMC is described in [Betancourt and Girolmi, 2015].
- ▶ It occurs in hierarchical models when we have sparse data and a centered prior.

Reparameterization

Proposition

Reparameterize the model to avoid the funnel shape. We will do so by standardizing α .

$$\alpha_{\text{std},i} := \frac{\alpha_i - \mu}{\sigma}$$

Then

$$\alpha_{\text{std}} \sim \text{Normal}(0, 1)$$

Reparameterization

Then

$$\alpha_i = \mu + \sigma \alpha_{\text{std},i}$$

Hence

$$y_i \sim p(\mu + \sigma \alpha_{\text{std},i})$$

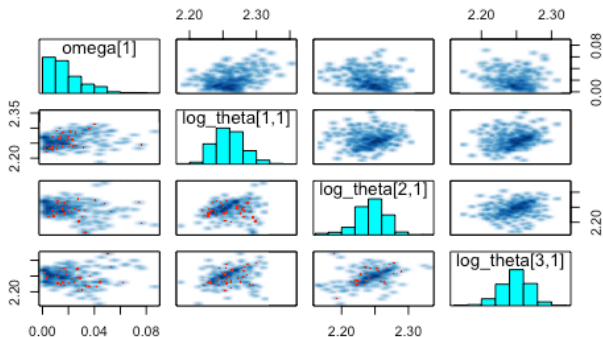
- ▶ Same data generating process; but how does this affect the geometry of the posterior?

Reparameterization

Our model is a little more complicated than the above example:

- ▶ a lot of parameters (100 +)!
- ▶ multiple population parameters and hierarchical structures.
- ▶ these parameters follow a log normal distribution (so we need a pairs plot with $\log \theta$).

Reparameterization



Reparameterization

Exercise 6: Reparametrize the two compartment population model and fit it.}

- ▶ First, work out the appropriate parametrization. You should start with $\log \theta_i \sim \text{Normal}(\theta_{\text{pop},i}, \omega)$
- ▶ Write, fit, and check the inference (run 100 chains).
- ▶ What kind of predictive checks can we do?

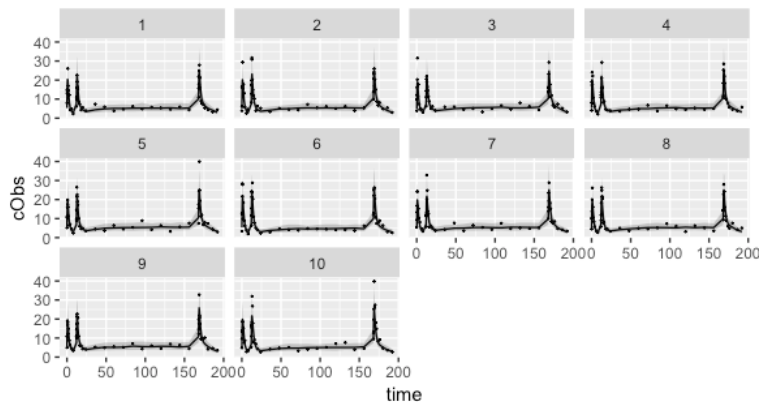
Reparameterization

Need:

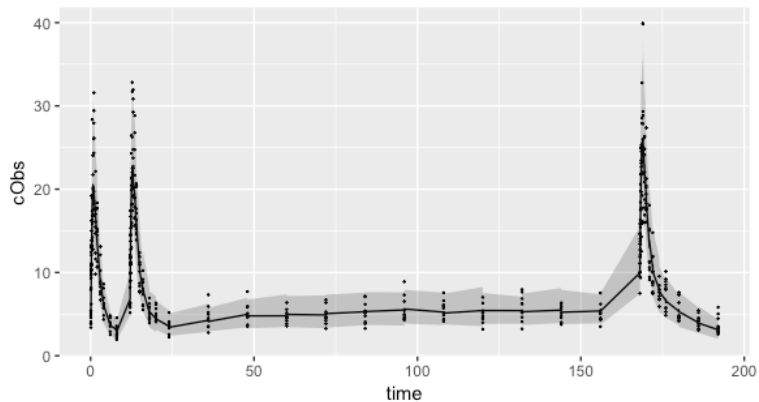
- ▶ predictions at an individual level
- ▶ predictions at a population level

As always, this comes down to properly writing the data generating process in the generated quantities block.

Individual predictions



Population predictions



Further reading

For a very good case study on hierarchical models, see, Bob Carpenter's *Pooling with Hierarchical Models for Repeated Binary Trials*

<https://mc-stan.org/users/documentation/case-studies/pool-binary-trials.html>

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ODE group integrators

Single ODE system

```
pmx_integrate_ode_rk45  
pmx_integrate_ode_bdf  
pmx_integrate_ode_adams
```

ODE group

```
pmx_integrate_ode_group_rk45  
pmx_integrate_ode_group_bdf  
pmx_integrate_ode_group_adams
```

Single ODE system

```
real[,]  
pmx_integrate_ode_xxx(  
  f,  
  real[] y0, real t0,  
  real[] ts,  
  real[] theta,  
  real[] x_r, int[] x_i,  
  ...);
```

ODE group

```
matrix  
pmx_integrate_ode_group_xxx(  
  f,  
  real[ , ] y0, real t0,  
  int[] len, real[] ts,  
  real[ , ] theta,  
  real[ , ] x_r, int[ , ] x_i,  
  ...);
```

ODE group integrators

Single ODE system

```
real[ , ]  
pmx_integrate_ode_xxx(  
    f,  
    real[] y0, real t0,  
    real[] ts,  
    real[] theta,  
    real[] x_r, int[] x_i,  
    ...);
```

ODE group

```
matrix  
pmx_integrate_ode_group_xxx(  
    f,  
    real[ , ] y0, real t0,  
    int[] len, real[] ts,  
    real[ , ] theta,  
    real[ , ] x_r, int[ , ] x_i,  
    ...);
```

- ▶ `len` specifies the length of data for each subject within the above ragged arrays, and the size of `len` is the size of the population.
- ▶ The group integrators return a single matrix ragged column-wise. The number of rows equals to the size of ODE system.

Exercise 7

autocatalytic reaction model: ODE group version

- ▶ Change the loop with the numerical integrator to use group integrator.
- ▶ Remember the return of the group integrator is a matrix
 - ▶ nb. of rows: nb. of states
 - ▶ nb. of cols: nb. of *total* results-extraction time points.

Exercise 7

Build and run

- ▶ Edit/Add cmdstan/make/local

```
TORSTEN_MPI = 1  # flag on torsten's MPI solvers
CXXFLAGS += -isystem /usr/local/include    # path to MPI
↳ library's headers
```

- ▶ Build in cmdstan

```
make ../example-models/chemical_reactions/chem_group
```

- ▶ Run

```
mpiexec -n 2 -l ./chem_group sample adapt delta=0.95 random
↳ seed=1104508041 data file=chem.data.R init=chem.init.R
```

Exercise 7

- ▶ What does output say?
- ▶ How many cores can you use until performance saturates? Why?
- ▶ (optional) Can you do it using Stan's `map_rect`? Is there a difference in style, output, and performance?

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PMX population solvers

Single ODE system	ODE group
<code>pmx_solve_rk45</code>	<code>pmx_solve_group_rk45</code>
<code>pmx_solve_bdf</code>	<code>pmx_solve_group_bdf</code>
<code>pmx_solve_adams</code>	<code>pmx_solve_group_adams</code>

Individual solvers

matrix

```
pmx_solve_bdf(f, int nCmt,  
  real[] time, real[] amt,  
  real[] rate, real[] ii,  
  int[] evid, int[] cmt,  
  real[] addl, int[] ss,  
  real[] theta, real[]  
    ↪ biovar,  
  real[] tlag, real rel_tol,  
  real abs_tol, int  
    ↪ max_step);
```

Population solvers

matrix

```
pmx_solve_group_bdf(f, int nCmt,  
  int[] len, real[] time,  
  real[] amt, real[] rate,  
  real[] ii, int[] evid,  
  int[] cmt, real[] addl,  
  int[] ss, real[ , ] theta,  
  real[ , ] biovar, real[ , ] tlag,  
  real rel_tol, real abs_tol,  
  int max_step);
```

PMX population solvers

matrix

```
pmx_solve_group_bdf(f, int nCmt, int[] len, real[] time, real[]  
  ↪ amt, real[] rate, real[] ii, int[] evid, int[] cmt, real[]  
  ↪ addl, int[] ss, real[,] theta, real[,] biovar, real[,]  
  ↪ tlag, real rel_tol, real abs_tol, int max_step);
```

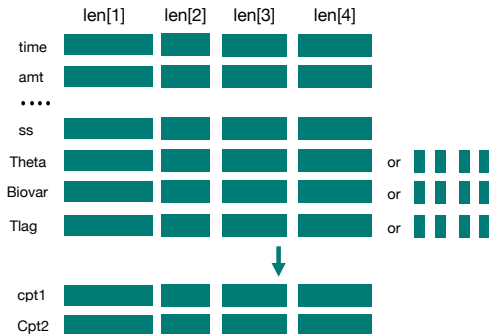


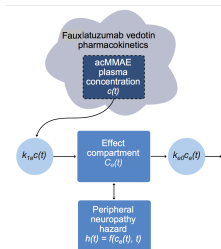
Figure: arguments and output of `pmx_solve_group_xxx`

Time-to-event model

We analyze the time to the first grade 2+ peripheral neuropathy (PN) event in patients treated with an antibody-drug conjugate (ADC) delivering monomethyl auristatin E (MMAE). We will simulate and analyze data using a simplified version of the model reported in [Lu et al., 2017].

- ▶ Fauxlatuzumab vedotin 1.2 mg/kg IV boluses q3w \times 6 does.
- ▶ 19 patients with 6 right-censored (simulated data).

Model scheme



Note

- ▶ To keep things simpler, we use the simulated individual CL and V values, and only model PD part of the problem.
- ▶ PN hazard is substantially delayed relative to PK exposure.
- ▶ Hazard increases over time to an extent not completely described by P

Likelihood

Likelihood for time to first $\text{PN} \geq 2$ event in the i^{th} patient:

$$L(\theta | t_{\text{PN},i}, \text{censor}_i, X_i) \\ = \begin{cases} h_i(t_{\text{PN},i} | \theta, X_i) e^{-\int_0^{t_{\text{PN},i}} h_i(u | \theta, X_i) du}, & \text{censor}_i = 0 \\ e^{-\int_0^{t_{\text{PN},i}} h_i(u | \theta, X_i) du}, & \text{censor}_i = 1 \end{cases}$$

where

$t_{\text{PN}} \equiv$ time to first $\text{PN} \geq 2$ or right censoring event

$\theta \equiv$ model parameters

$X \equiv$ independent variables / covariates

$\text{censor} \equiv \begin{cases} 1, & \text{PN} \geq 2 \text{ event is right censored} \\ 0, & \text{PN} \geq 2 \text{ event is observed} \end{cases}$

One can see the expression

$$e^{-\int_0^{t_{\text{PN},i}} h_i(u | \theta, X_i) du}$$

as the survival function at time t .

ODEs

Hazard of PN grade 2+ based on the Weibull distribution, with drug effect proportional to effect site concentration of MMAE:

$$h_j(t) = \beta E_{\text{drug}j}(t)^{\beta} t^{(\beta-1)}$$

$$E_{\text{drug}j}(t) = \alpha c_{ej}(t)$$

$$c'_{ej}(t) = k_{e0} (c_j(t) - c_{ej}(t)).$$

Overall ODE system including integration of the hazard function:

$$x'_1 = -\frac{CL}{V} x_1 \tag{1}$$

$$x'_2 = k_{e0} \left(\frac{x_1}{V} - x_2 \right) \tag{2}$$

$$x'_3 = h(t) \tag{3}$$

where $x_2(t) = c_e(t)$ and $x_3(t) = \int_0^t h(u) du$ aka cumulative hazard.

Exercise 8: write the ODE system

```
functions{
  real[] oneCptPNODE(real t, real[] x, real[] parms, real[]
    ↪ x_r, int[] x_i){
    real dxdt[3];
    real CL = parms[1];
    real V = parms[2];
    real ke0 = parms[3];
    real alpha = parms[4];
    real beta = parms[5];
    real Edrug;
    real hazard;
    /* ... */
    return dxdt;
  }
}
```

Exercise 8: the ODE system

```
real[] oneCptPNODE(real t, real[] x, real[] parms, real[] x_r,  
    ↪ int[] x_i){  
    real dxdt[3];  
    real CL = parms[1];  
    real V = parms[2];  
    real ke0 = parms[3];  
    real alpha = parms[4];  
    real beta = parms[5];  
    real Edrug;  
    real hazard;  
  
    dxdt[1] = -(CL / V) * x[1];  
    dxdt[2] = ke0 * (x[1] / V - x[2]);  
    Edrug = alpha * x[2];  
    if(t == 0){  
        hazard = 0;  
    }else{  
        hazard = beta * Edrug^beta * t^(beta - 1);  
    }  
    dxdt[3] = hazard;  
    return dxdt;  
}
```

Parameters

parameters

```
parameters{
  real<lower = 0> ke0;
  real<lower = 0> alpha;
  real<lower = 0> beta;
}

transformed parameters{
  vector<lower = 0>[nPNObs] survObs;
  row_vector<lower = 0>[nPNObs] EdrugObs;
  vector<lower = 0>[nPNObs] hazardObs;
  vector<lower = 0>[nPNCens] survCens;
  matrix<lower = 0>[3, nt] x;
  real<lower = 0> parms[nId, 5];

  for(j in 1:nId) {
    parms[j, ] = {CL[j], V[j], ke0, alpha, beta};
  }
  /* ... */
}
```

Parameters

```
transformed parameters{  
  vector<lower = 0>[nPNObs] survObs;  
  row_vector<lower = 0>[nPNObs] EdrugObs;  
  vector<lower = 0>[nPNObs] hazardObs;  
  vector<lower = 0>[nPNCens] survCens;  
  matrix<lower = 0>[3, nt] x;  
  real<lower = 0> parms[nId, 5];  
  
  for(j in 1:nId) {  
    parms[j, ] = {CL[j], V[j], ke0, alpha, beta};  
  }  
  /* ... */  
}
```

Exercise 9

- ▶ Use `pmx_solve_group_rk45` to solve for `x`.
- ▶ Write likelihood expressions for `survObs`, `EdrugObs`, `hazardObs`, and `survCens`.

Exercise 9

- Stan's target variable and user-defined likelihood.

```
model{  
  ke0 ~ normal(0, 0.0005);  
  alpha ~ normal(0, 0.000003);  
  beta ~ normal(0, 1.5);  
  
  target += log(hazardObs .* survObs); // observed PN event log  
    ↪ likelihood  
  target += log(survCens); // censored PN event log likelihood  
}
```

Exercise 9

Edit/Add cmdstan/make/local

```
TORSTEN_MPI = 1           # flag on torsten's MPI solvers  
CXXFLAGS += -isystem /usr/local/include # path to MPI library's  
↳ headers
```

Build in cmdstan

```
make ../example-models/ttpn2/ttpn2_group
```

Run

```
mpiexec -n 4 -l ttpn2_group sample num_warmup=500  
↳ num_samples=500 data file=ttpn2.data2.R init=ttpn2.init.R
```

Exercise 9

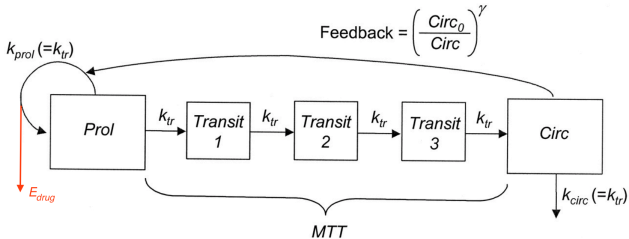
- ▶ The parallel performance is not optimal, why?
- ▶ Can you do it using Stan's `map_rect`?

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Friberg-Karlsson semi-mechanistic model

[Friberg et al., 2002]



ODE system of F-K model

$$y'_{\text{prol}} = k_{\text{tr}} y_{\text{prol}} (1 - E_{\text{drug}}) \left(\frac{Circ_0}{y_{\text{circ}}} \right)^{\gamma} - k_{\text{tr}} y_{\text{prol}}$$

$$y'_{\text{tr1}} = k_{\text{tr}} y_{\text{prol}} - k_{\text{tr}} y_{\text{tr1}}$$

$$y'_{\text{tr2}} = k_{\text{tr}} y_{\text{tr1}} - k_{\text{tr}} y_{\text{tr2}}$$

$$y'_{\text{tr3}} = k_{\text{tr}} y_{\text{tr2}} - k_{\text{tr}} y_{\text{tr3}}$$

$$y'_{\text{circ}} = k_{\text{tr}} y_{\text{tr3}} - k_{\text{tr}} y_{\text{circ}}$$

where $E_{\text{drug}} = \alpha \frac{y_{\text{cent}}}{V_{\text{cent}}}$, $k_{\text{tr}} = 4/MTT$, and $\alpha \approx 3e - 4$.

- ▶ y_{cent} is obtained from a two compartment model.
- ▶ Our PK/PD model therefore has a total of 8 equations.
- ▶ This problem can be solved using `pmx_solve_*`.

Coupled PK-PD system

Alternatively, we may elect to solve the PK ODEs **analytically** and the PD ODEs **numerically**.

- ▶ This can yield some speedup, in particular for problems that require ODE solutions and sensitivities (e.g [Margossian and Gillespie, 2017b]).

```
real[] pmx_solve_twocpt_rk45(reduced_ODE_system, int nOde,  
→ real[] time, real[] amt, real[] rate, real[] ii, int[]  
→ evid, int[] cmt, real[] addl, int[] ss, real[] theta,  
→ real[] biovar, real[] tlag, real rel_tol, real abs_tol,  
→ real max_step)
```

Coupled PK-PD system

```
real[] pmx_solve_twocpt_rk45(reduced_ODE_system, int nOde,  
  ↪ real[] time, real[] amt, real[] rate, real[] ii, int[]  
  ↪ evid, int[] cmt, real[] addl, int[] ss, real[] theta,  
  ↪ real[] biovar, real[] tlag, real rel_tol, real abs_tol,  
  ↪ real max_step)
```

- ▶ we now pass a "reduced system".
- ▶ we specify the number of ODEs to be solved numerically, not the number of compartments.
- ▶ theta now contains the parameters for the two cpt model, followed by the parameters that get passed to the numerical solver:

```
theta = {CL, Q, VC, VP, ka, /* ... */};
```

Reduced system

```
real[] reduced_system(real time, real[] y, real[] yPK, real[]  
  ↪ theta, real[] x_r, int[] x_i) {  
  real[3] dydt;  
  /* .... */  
  return dydt;  
}
```

Exercise 4 (optional): Write, fit, and diagnose a Friberg-Karlsson model with a two compartment with first order absorption PK. Use `FKModel.r` and `data/FKModel.data.r`.

Exercise 10

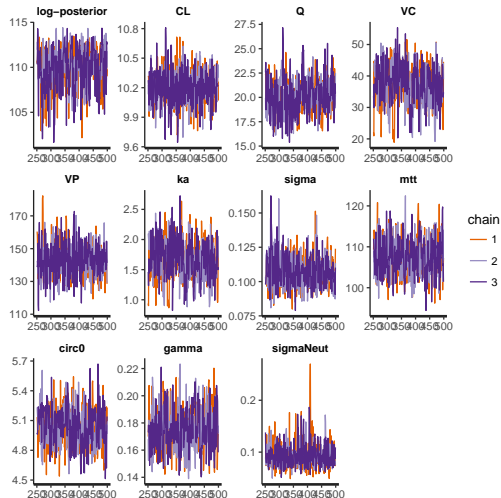
Write, fit, and diagnose a Friberg-Karlsson model with a two compartment with first order absorption PK. Use `FKModel.r` and `data/FKModel.data.r`.

- ▶ You may either use `pmx_solve_*` or `pmx_solve_twocpt_*`.
- ▶ Use $\alpha = 3e - 4$ and estimate all other 8 ODE coefficients, i.e. $\theta = \{CL, Q, VC, VP, ka, MTT, circ0, \gamma\}$.
- ▶ The initial state for the neutrophil count is $Circ_0$. Either edit the event schedule to reflect this at time 0, or write the solution to your ODEs as a deviation from the baseline.

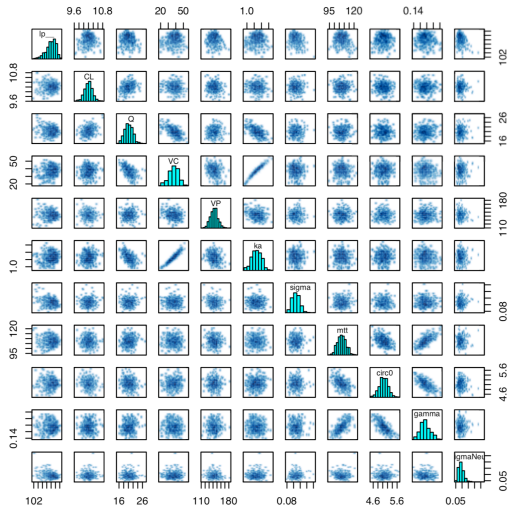
Exercise 10

- ▶ This exercise entails a few subtleties; in the interest of time we won't go through it in class.
- ▶ Here are however results I get from 3 chains with 500 iterations you can use as a benchmark.

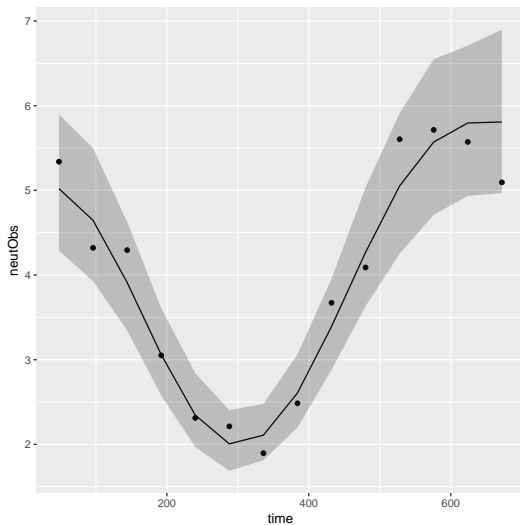
Exercise 10



Exercise 10



Exercise 10



Reference I



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