

3_STAN

Stan code

Next we encode similar models in Stan. We start with the presidents, whom we model by student's t likelihood. This is a robust version of normal likelihood, because the heavier tails of the t distribution account better for outliers. Moreover, we let the data determine the heaviness of the tails by fitting the shape parameter nu on the data (in classical stats this parameter is called df - degrees of freedom) model:

$$\begin{aligned}y_i &\sim \text{student}(\nu, \mu, \sigma) \\ \nu &\sim \text{student}(2, 1, 50) \\ \mu &\sim \text{normal}(180, 10) \\ \sigma &\sim \text{student}(4, 0, 5)\end{aligned}$$

ülesanne - kirjutage see mudel ümber nii, et see kasutaks normaaljaotuse tõepära ja normaaljaotuse prioreid, kusjuures tehke sigma prior vähem informatiivseks.

```
modelString <- "

data {
  int<lower=0> N; //the nr of data points
  vector[N] heights; //a vector of N datapoints called heights.
}
parameters {
  real<lower=1> nu; //shape parameter cannot be <1
  real mu; //mean
  real<lower=0> sigma; //standard deviation
}
model {
  nu ~ student_t(2, 1, 50); //prior for shape parameter
  mu ~ normal(180, 10); //prior for the mean
  sigma ~ student_t(4, 0, 5); //prior for the SD
  heights ~ student_t(nu, mu, sigma); //regression model
}

"

stanDso <- stan_model(model_code = modelString)

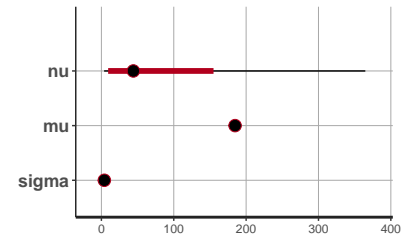
heights <- tibble(value = c(183, 192, 182, 183, 177, 185, 188, 188, 182, 185, 188))

dataList <- list(heights = heights$value, N = length(heights$value))
```

```
fit1 <- sampling(object = stanDso, data = dataList, chains = 3, cores = 3, iter = 1000,
  warmup = 200, thin = 1)
```

```
write_rds(fit1, "models/fit1.rds")
```

```
fit1 <- read_rds("models/fit1.rds")
plot(fit1)
```



```
fit1 %>% tidy()
```

```
## # A tibble: 3 x 3
##   term estimate std.error
##   <chr>      <dbl>      <dbl>
## 1 nu         83.2        223.
## 2 mu        185.         1.36
## 3 sigma      4.26         1.15
```

```
fit1
```

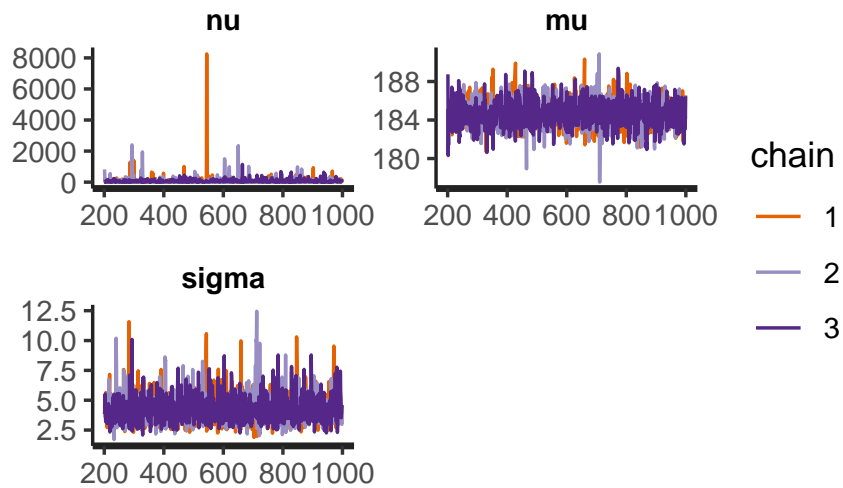
```
## Inference for Stan model: df64bdb04a46456ae5e57c45f44b989e.
## 3 chains, each with iter=1000; warmup=200; thin=1;
## post-warmup draws per chain=800, total post-warmup draws=2400.
##
```

	mean	se_mean	sd	2.5%	25%	50%	75%	97.5%	n_eff	Rhat
nu	83.22	5.51	223.16	3.75	20.77	44.23	85.62	366.07	1640	1
mu	184.71	0.04	1.36	181.97	183.87	184.72	185.51	187.42	1383	1
sigma	4.26	0.03	1.15	2.58	3.51	4.06	4.82	6.94	1873	1
lp__	-21.83	0.05	1.47	-25.72	-22.50	-21.44	-20.80	-20.21	921	1

```
##
## Samples were drawn using NUTS(diag_e) at Tue Aug 4 17:58:15 2020.
## For each parameter, n_eff is a crude measure of effective sample size,
## and Rhat is the potential scale reduction factor on split chains (at
## convergence, Rhat=1).
```

```
markov chains
```

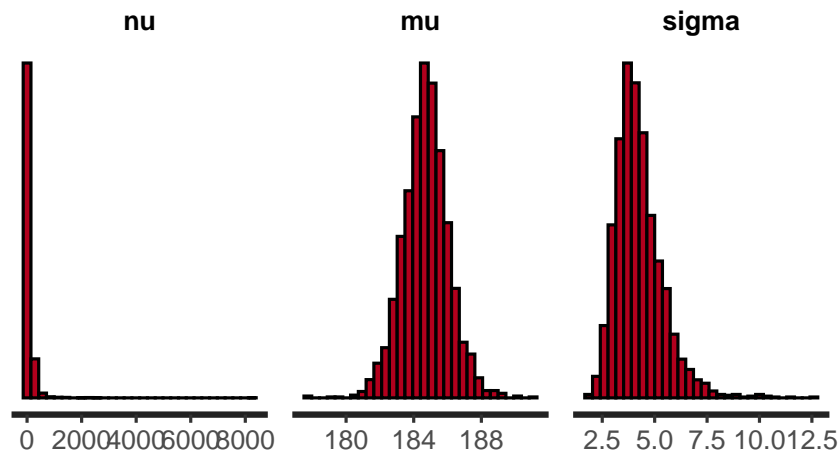
```
traceplot(fit1, nrow = 2)
```



posteriors:

```
# plot(fit1, plotfun = 'hist')
stan_hist(fit1)
```

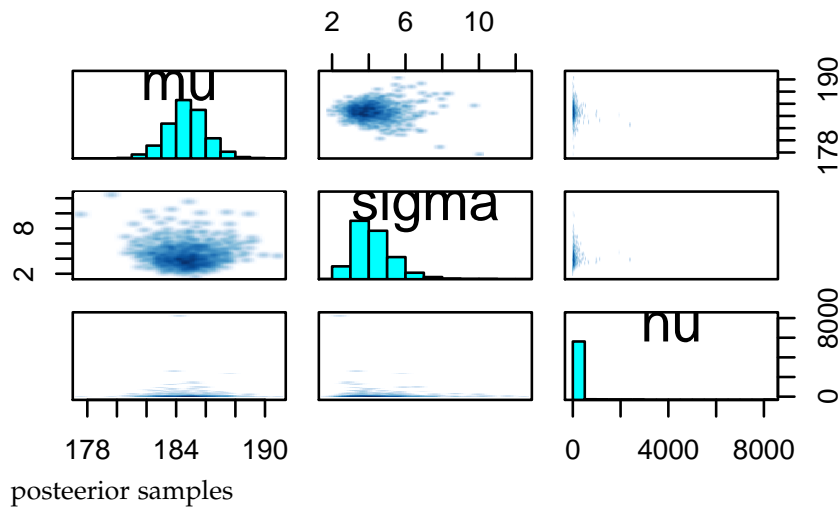
```
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
```



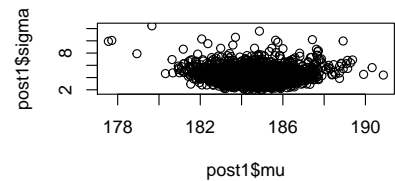
```
pairs(fit1, pars = c("mu", "sigma", "nu"))
```

```
## Warning in KernSmooth::bkde2D(x, bandwidth = bandwidth, gridsize = nbin, :
## Binning grid too coarse for current (small) bandwidth: consider increasing
## 'gridsize'
```

```
## Warning in KernSmooth::bkde2D(x, bandwidth = bandwidth, gridsize = nbin, :
## Binning grid too coarse for current (small) bandwidth: consider increasing
## 'gridsize'
```



```
post1 <- as.data.frame(fit1)
plot(post1$mu, post1$sigma)
```



```
pres_brms <- brm(value ~ 1, data = heights, family = student())
write_rds(pres_brms, "models/pres_brms")
```

```
pres_brms <- read_rds("models/pres_brms")
tidy(pres_brms)
```

##	term	estimate	std.error	lower	upper
## 1	b_Intercept	184.811496	1.447901	182.461040	187.140324
## 2	sigma	4.350663	1.214018	2.777434	6.537025
## 3	nu	20.378918	13.723315	4.539076	47.855902
## 4	lp__	-37.590361	1.377186	-40.288812	-36.106543

exerciser – run the same for normal likelihood

expand this into regression

We simulate data:

```
set.seed(12)
data <- tibble(value = c(rnorm(n = 8, mean = 100, sd = 10), rnorm(8, 150, 15)), indeks = rep(c(0,
1), each = 8))

t.test(data = data, value ~ indeks)

##
## Welch Two Sample t-test
##
## data: value by indeks
```

```

## t = -10.449, df = 13.276, p-value = 8.89e-08
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -60.21584 -39.61759
## sample estimates:
## mean in group 0 mean in group 1
##      93.75789      143.67460

lm(data = data, value ~ indeks)

##
## Call:
## lm(formula = value ~ indeks, data = data)
##
## Coefficients:
## (Intercept)      indeks
##      93.76      49.92

modelString <- "

data {
  int<lower=0> N;
  vector[N] y;
  vector[N] x;
}
parameters {
  real<lower=1> nu;
  real<lower=0> alpha;
  real beta;
  real<lower=0> sigma;
}
model {
  nu ~ student_t(4, 2, 30); //prior for shape parameter
  alpha ~ normal(100, 50); //prior for the mean
  beta ~ normal(50, 50);
  sigma ~ student_t(4, 0, 20); //prior for the SD
  \ty ~ student_t(nu, alpha + beta * x, sigma);
}

"

stanDso <- stan_model(model_code = modelString)

dataList <- list(y = data$value, x = data$indeks, N = nrow(data))

fit2 <- sampling(object = stanDso, data = dataList, chains = 3, cores = 3, iter = 1000,

```

```

warmup = 400, thin = 1)

write_rds(fit2, "models/fit2.rds")

fit2 <- read_rds("models/fit2.rds")
fit2

## Inference for Stan model: da9c5cf4c8c4dcd7b2ad28f7c999ddec.
## 3 chains, each with iter=1000; warmup=400; thin=1;
## post-warmup draws per chain=600, total post-warmup draws=1800.
##
##          mean se_mean    sd   2.5%   25%   50%   75%  97.5% n_eff Rhat
## nu       34.12    0.99 33.42   3.06  12.91  25.55  44.09 116.01 1146   1
## alpha    93.46    0.13  3.79  86.17  90.92  93.43  95.91 100.87  831   1
## beta     50.33    0.18  5.37  39.33  47.07  50.29  53.72  61.26  865   1
## sigma    9.72    0.06  2.11   6.31   8.20   9.49  10.89  14.49 1126   1
## lp__    -41.35    0.06  1.62 -45.57 -42.12 -41.01 -40.15 -39.36  718   1
##
## Samples were drawn using NUTS(diag_e) at Tue Aug  4 18:50:47 2020.
## For each parameter, n_eff is a crude measure of effective sample size,
## and Rhat is the potential scale reduction factor on split chains (at
## convergence, Rhat=1).

post2 <- as.data.frame(fit2)
gr2 <- post2$alpha + post2$beta
hist(gr2)

plot(fit2, pars = c("alpha", "beta"))

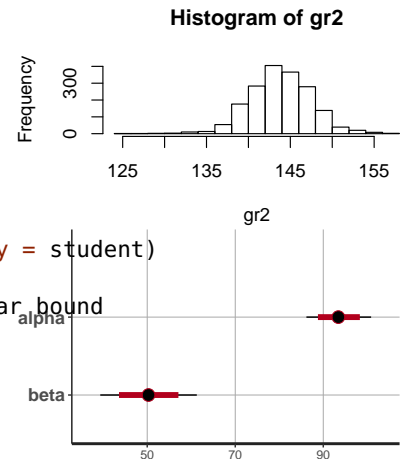
get_prior(data = data, bf(value ~ indeks, sigma ~ indeks), family = student)

##          prior      class coef group resp dpar nlpar bound
## 1                b
## 2                b indeks
## 3 student_t(3, 123, 39) Intercept
## 4      gamma(2, 0.1)      nu
## 5                b          sigma
## 6                b indeks          sigma
## 7 student_t(3, 0, 10) Intercept          sigma

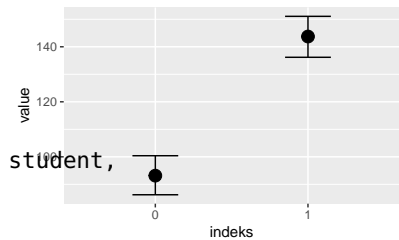
data1 <- data
data1$indeks <- as.character(data1$indeks)

fit3 <- brm(data = data1, value ~ indeks, family = student, prior = c(prior(student_t(4,
  2, 30), class = nu), prior(normal(100, 50), class = Intercept), prior(normal(50,
  50), class = b), prior(student_t(4, 0, 20), class = sigma)), cores = 3, chains = 3)
write_rds(fit3, "models/fit3.rds")

```



```
fit3 <- read_rds("models/fit3.rds")
conditional_effects(fit3)
```



```
fit4 <- brm(data=data1, bf(value~indeks, sigma~indeks), family = student,
            cores=3, chains = 3)
write_rds(fit4, "models/fit4.rds")
```

```
(fit4 <- read_rds("models/fit4.rds"))
```

```
## Family: student
## Links: mu = identity; sigma = log; nu = identity
## Formula: value ~ indeks
##          sigma ~ indeks
## Data: data1 (Number of observations: 16)
## Samples: 3 chains, each with iter = 2000; warmup = 1000; thin = 1;
##          total post-warmup samples = 3000
##
## Population-Level Effects:
##           Estimate Est.Error l-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
## Intercept          93.20      4.20   84.91   101.66 1.00    2071    1376
## sigma_Intercept      2.30      0.34    1.63     3.02 1.00    2065    1317
## indeks1             50.47      5.67   39.04    61.89 1.00    1998    1595
## sigma_indeks1       -0.15      0.45   -1.02     0.79 1.00    2129    2043
##
## Family Specific Parameters:
##           Estimate Est.Error l-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
## nu      19.51     13.70      2.91    55.80 1.00     2094     1206
##
## Samples were drawn using sampling(NUTS). For each parameter, Bulk_ESS
## and Tail_ESS are effective sample size measures, and Rhat is the potential
## scale reduction factor on split chains (at convergence, Rhat = 1).
```

```
exp(2.31)
```

```
## [1] 10.07442
```

```
exp(-0.17 + 2.31)
```

```
## [1] 8.499438
```

```
data1 %>% group_by(indeks) %>% summarise(SD = sd(value))
```

```
## 'summarise()' ungrouping output (override with '.groups' argument)
```

```
## # A tibble: 2 x 2
```

```
##   indeks    SD
```

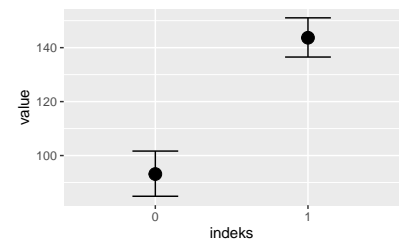


```
##   <chr>  <dbl>
## 1 0      10.6
## 2 1       8.36

t.test(data = data1, value ~ indeks)

##
## Welch Two Sample t-test
##
## data:  value by indeks
## t = -10.449, df = 13.276, p-value = 8.89e-08
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
##  -60.21584 -39.61759
## sample estimates:
## mean in group 0 mean in group 1
##      93.75789      143.67460

conditional_effects(fit4)
```



model the proportion via binomial distribution

Here we have a covid-19 antibody study of 3330 americans, of whom 50 tested positive. Remember, p is the single parameter in this model.
raw ratio:

50/3330

```
## [1] 0.01501502
```

or 1.5%

```
modelString <- "
```

```
data {
  int<lower = 0> y;
  int<lower = 0> n;
}
parameters {
  real<lower=0, upper = 1> p;
}
model {
  p ~ beta(1,1);
  y ~ binomial(n, p);
}
```

```

"
stanDso2 <- stan_model(model_code = modelString)

y <- 50
n <- 3330

dataList <- list(y = y, n = n)

stanFit3 <- sampling(object = stanDso2, data = dataList, chains = 3, cores = 3, iter = 1000,
  warmup = 200, thin = 1)

write_rds(stanFit3, "models/stanFit3.rds")

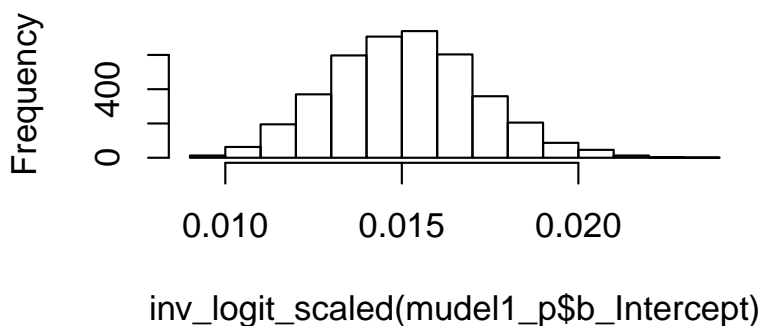
## # A tibble: 1 x 3
##   term estimate std.error
##   <chr>      <dbl>      <dbl>
## 1 p          0.0122    0.00264

mudel1 <- brm(success | trials(3330) ~ 1, data = list(success = 50), family = binomial)
write_rds(mudel1, "models/mudel1.rds")

mudel1 <- read_rds("models/mudel1.rds")
mudel1_p <- posterior_samples(mudel1)
hist(inv_logit_scaled(mudel1_p$b_Intercept))

```

istogram of inv_logit_scaled(mudel1_p\$b_In1



võtame arvesse testi sensitiivsuse ja spetsiifilisuse

Seda määrati uuringu käigus eraldi, kasutades inimesi, kes konkreetseid proove võtsid ja analüüsisid

```
50/3330 * 103/122 + (1 - 50/3330) * (1 - 399/401)
```

```
## [1] 0.01758925
```

ehk 1.8%

siin jätan priori defineerimata - Stan teeb selle tasaseks prioriks.

```
modelString <- "data {
  int<lower = 0> y_sample;
  int<lower = 0> n_sample;
  real<lower = 0, upper = 1> spec;
  real<lower = 0, upper = 1> sens;
}
parameters {
  real<lower=0, upper = 1> p;
}
model {
  real p_sample = p * sens + (1 - p) * (1 - spec);
  y_sample ~ binomial(n_sample, p_sample);
}"
```

```
stanDso3 <- stan_model(model_code = modelString)
```

```
y_sample <- 50
n_sample <- 3330
spec <- 399/401
sens <- 103/122
```

```
dataList <- list(y_sample = y_sample, n_sample = n_sample, spec = spec, sens = sens)
```

```
stanFit3 <- sampling(object = stanDso3, data = dataList, chains = 3, cores = 3, iter = 1000,
  warmup = 200, thin = 1)
write_rds(stanFit3, "models/stanFit3.rds")
```

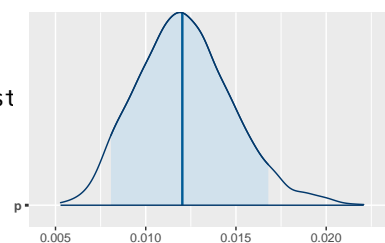
```
stanFit3 <- read_rds("models/stanFit3.rds")
tidy(stanFit3)
```

```
## # A tibble: 1 x 3
##   term estimate std.error
##   <chr>      <dbl>      <dbl>
## 1 p          0.0122    0.00264
```

```
mcmc_areas(stanFit3, pars = c("p"), prob = 0.9)
```

```
## Warning: 'expand_scale()' is deprecated; use 'expansion()' inst
```

```
modelString <- "data {
  int<lower = 0> y_sample;
  int<lower = 0> n_sample;
```



```

  int<lower = 0> y_spec;
  int<lower = 0> n_spec;
  int<lower = 0> y_sens;
  int<lower = 0> n_sens;
}
parameters {
  real<lower=0, upper = 1> p;
  real<lower=0, upper = 1> spec;
  real<lower=0, upper = 1> sens;
}
model {
  real p_sample = p * sens + (1 - p) * (1 - spec);
  y_sample ~ binomial(n_sample, p_sample);
  y_spec ~ binomial(n_spec, spec);
  y_sens ~ binomial(n_sens, sens);
}

```

```
stanDso4 <- stan_model(model_code = modelString)
```

```

y_sample <- 50
n_sample <- 3330
y_spec <- 399
n_spec <- 401
y_sens <- 103
n_sens <- 122

```

```
dataList <- list(y_sample = y_sample, n_sample = n_sample, y_spec = y_spec, n_spec = n_spec,
  y_sens = y_sens, n_sens = n_sens)
```

```
stanFit4 <- sampling(object = stanDso4, data = dataList, chains = 3, cores = 3, iter = 1000,
  warmup = 200, thin = 1)
```

```
write_rds(stanFit4, "models/stanFit4.rds")
```

```
stanFit4 <- read_rds("models/stanFit4.rds")
tidy(stanFit4)
```

```

## # A tibble: 3 x 3
##   term estimate std.error
##   <chr>      <dbl>      <dbl>
## 1 p          0.0103    0.00472
## 2 spec       0.993     0.00350
## 3 sens       0.838     0.0328

```

lets slap some priors on p, sens and spec. Especially for p I'm putting a strong prior that says that most likely incidence is 1% and

incidences over 3% are very unlikely. I am vaguer for sensitivity and specificity where the prior has heavy tails allowing for a wide range of possibilities.

```

modelString <- "data {
  int<lower = 0> y_sample;
  int<lower = 0> n_sample;
  int<lower = 0> y_spec;
  int<lower = 0> n_spec;
  int<lower = 0> y_sens;
  int<lower = 0> n_sens;
}
parameters {
  real<lower=0, upper = 1> p;
  real<lower=0, upper = 1> spec;
  real<lower=0, upper = 1> sens;
}
model {
  real p_sample = p * sens + (1 - p) * (1 - spec);
  p ~ normal(0.01, 0.01);
  spec ~ student_t(6, 0.9, 0.1);
  sens ~ student_t(3, 0.8, 0.1);
  y_sample ~ binomial(n_sample, p_sample);
  y_spec ~ binomial(n_spec, spec);
  y_sens ~ binomial(n_sens, sens);
}"

stanDso5 <- stan_model(model_code = modelString)

y_sample <- 50
n_sample <- 3330
y_spec <- 399
n_spec <- 401
y_sens <- 103
n_sens <- 122

dataList <- list(y_sample = y_sample, n_sample = n_sample, y_spec = y_spec, n_spec = n_spec,
  y_sens = y_sens, n_sens = n_sens)

stanFit5 <- sampling(object = stanDso5, data = dataList, chains = 3, cores = 3, iter = 1000,
  warmup = 300, thin = 1)
write_rds(stanFit5, "models/stanFit5.rds")

stanFit5 <- read_rds("models/stanFit5.rds")
tidy(stanFit5)

```

```
## # A tibble: 3 x 3
##   term estimate std.error
##   <chr>     <dbl>     <dbl>
## 1 p         0.0106    0.00420
## 2 spec      0.993     0.00321
## 3 sens      0.833     0.0338

0.01059813 - 1.96 * 0.004201813

## [1] 0.002362577
```

lower bound for the p is about 0.2% incidence
higher bound is about 1.9%

```
0.01059813 + 1.96 * 0.004201813
```

```
## [1] 0.01883368
```

```
mcmc_dens(stanFit3, pars = c("p"))
```

```
mcmc_dens(stanFit4, pars = c("p"))
```

```
mcmc_dens(stanFit5, pars = c("p"))
```

