Practical 2: The brms package

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Exercise 1: Survival rate

We learn about the brms package and how to fit simple regression models.

Focus on: model output, convergence checks. Also some basic infos on priors and model predictions.

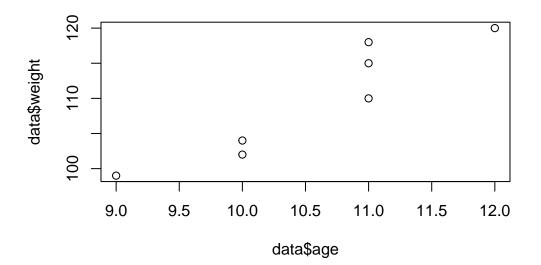
```
rm(list=ls())
library("brms")
library("ggplot2")
library("bayesplot")
library("loo")
library("cowplot")
try(dev.off())
```

Example 1: Linear regression

We start with our deer population and the simple weight~age example

Question: What's the average growth per year? (Slope in age)

Deterministic part: $\mu = a + b \cdot age$ Stochastic part: weight ~ Normal(μ, σ)



Basic brms functions

Instead of lm(), we use the brm() function. The formula notation is designed to be identical to lm, glm, lme4 (with few exceptions)

fit1 = brm(weight ~ age, data=data)

Looking at the summary table, we get a lot of infos:

If not specified otherwise, brms uses a normal distribution for the residuals: family=gaussian().

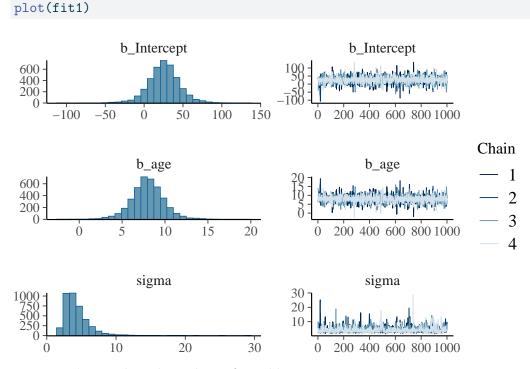
brms by default uses 4 chains, each with 1000 warmup and 1000 sampling iterations. The first thing you should look at are not parameter estimates, but Rhat and ESS. These indicate if the MCMC converged and the posterior distribution is properly sampled. Check if Rhat<1.01 and compare ESS to total number of draws.

```
summary(fit1)
```

```
Family: gaussian
Links: mu = identity; sigma = identity
Formula: weight ~ age
```

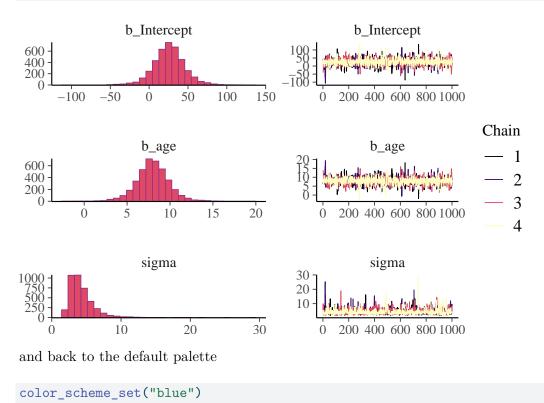
Data: data (Number of observations: 7) Draws: 4 chains, each with iter = 2000; warmup = 1000; thin = 1; total post-warmup draws = 4000 Regression Coefficients: Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS Intercept 25.56 21.23 -17.04 68.48 1.00 2328 1800 age 7.96 1.99 3.87 11.97 1.00 2336 1834 Further Distributional Parameters: Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS 4.31 1.90 2.18 9.12 1.00 1691 2047 sigma Draws were sampled 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).

Additionally, you should do a visual inspection of the MCMC. You get a histogram and a traceplot per parameter, which should look like a fuzzy caterpillar



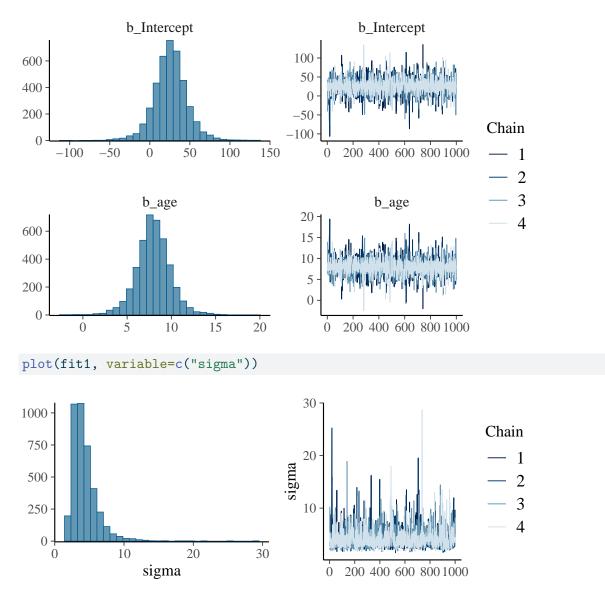
You can change the color palette if you like

```
color_scheme_set("viridisA")
plot(fit1)
```

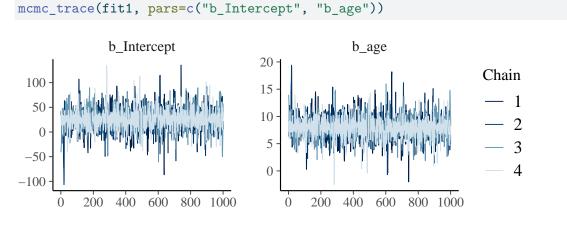


You can also specify to display just some selected parameters. Parameters of the deterministic model part begin with b_,the residual standard deviation is sigma

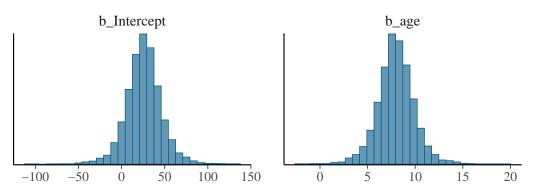
plot(fit1, variable=c("b_Intercept", "b_age"))



Histograms and traceplots can be plotted individually

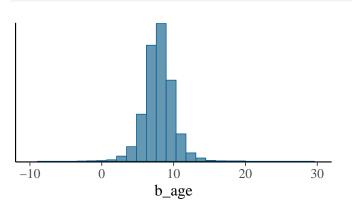


mcmc_hist(fit1, pars=c("b_Intercept", "b_age"))



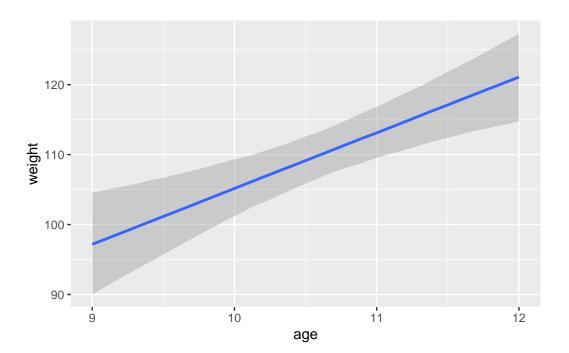
All the brms plots are done with ggplot2, so you can extract & modify them

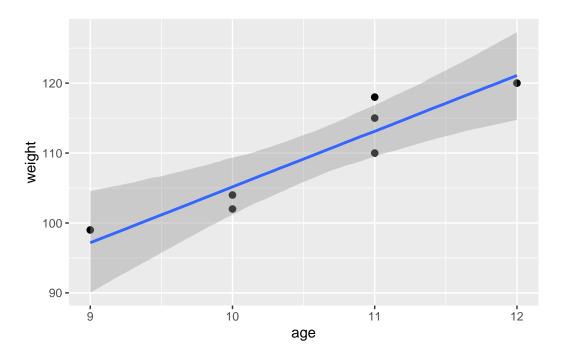
```
plot1 = mcmc_hist(fit1, pars=c("b_age"))
plot1 + xlim(-10,30)
```



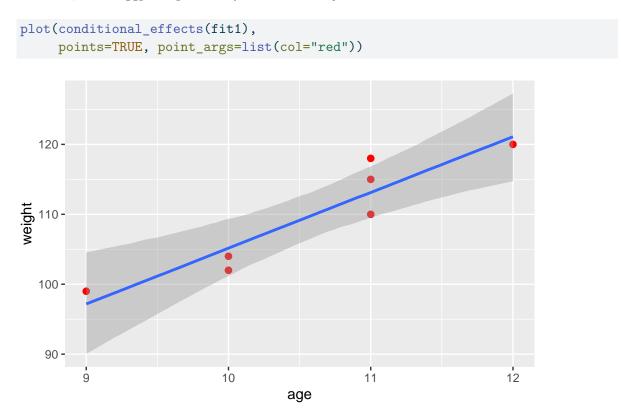
In this simple 1-predictor regression, model prediction are easily plotted vs data. conditional_effects() is a powerful function which we will use throughout the course.

plot(conditional_effects(fit1))





Again, this generates a ggplot object which can be modified, with some options in the plot function, or full ggplot options if you save the object



Note that we here only plot the uncertainty of the deterministic model part mu, more on that tomorrow. fitted() computes predictions of mu for each datapoint.

```
fitted(fit1) |> head()
```

Q2.5 Estimate Est.Error Q97.5 [1,] 105.18401 2.084831 101.18471 109.3446 [2,] 121.10918 3.186816 114.74524 127.2614 [3,] 113.14659 1.813679 109.52796 116.8500 [4,] 113.14659 1.813679 109.52796 116.8500 [5,] 97.22142 3.650626 90.06248 104.5488 [6,] 113.14659 1.813679 109.52796 116.8500

The brms package does not only offer model fitting via MCMC, it also has a lot of functions for model analysis and is compatible with a lot of other packages (e.g. emmeans). We will learn about some of these in the next days.

[1]	add_criterion	add_ic	as_draws_array	as_draws_df
[9]	as.array	as.data.frame	as.matrix	as.mcmc
[17]	coef	conditional_effects	conditional_smooths	control_params
[25]	fixef	formula	getCall	hypothesis
[33]	loo_compare	loo_epred	loo_linpred	loo_model_weigh
[41]	loo_subsample	100	L00	marginal_effect:
[49]	nchains	ndraws	neff_ratio	ngrps
[57]	nvariables	pairs	parnames	plot
[65]	posterior_linpred	posterior_predict	posterior_samples	posterior_smootl
[73]	predict	predictive_error	predictive_interval	prepare_predict:
[81]	ranef	reloo	residuals	restructure
[89]	summary	update	VarCorr	variables
see	'?methods' for accessing	help and source code		

brms specifications & priors

When we compare the results to frequentist lm-model, the slope is pretty close but there's ${\sim}0.5$ difference in intercepts.

fixef(fit1)

Estimate Est.ErrorQ2.5Q97.5Intercept 25.55816621.229873-17.04136268.47664age7.9625841.9904093.87254311.97261

lm(weight ~ age, data=data) |> coef()

(Intercept) age 26.2 7.9

So why are they different? What about priors, did we specify any?

The brm() function has A TON OF specifications, which we did not specify in the simple brm(weight~age, data=data) model, see the help function with ?brm. So brms uses default values.

?brm

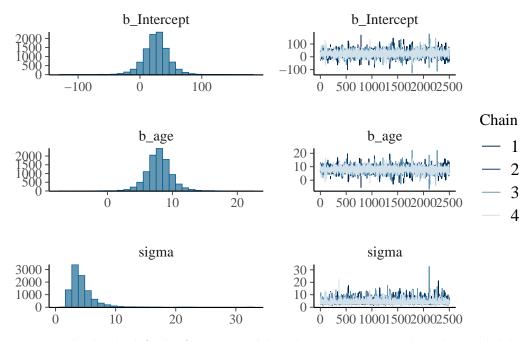
E.g., we can specify the number of chains & iterations manually. Per default, half of the iterations are used for warmup and are discared from the posterior sample.

With a larger number of samples (iter), we expect a more accurate approximation of the true posterior. Parameter means usually are quite correct even for low numbers, while outer quantiles (e.g. 90%, 95%) require larger numbers of samples.

summary(fit2)

```
Family: gaussian
  Links: mu = identity; sigma = identity
Formula: weight ~ age
   Data: data (Number of observations: 7)
  Draws: 4 chains, each with iter = 5000; warmup = 2500; thin = 1;
         total post-warmup draws = 10000
Regression Coefficients:
          Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
             26.25
                       21.61
                               -17.79
                                         68.66 1.00
                                                         5245
                                                                  4110
Intercept
              7.89
                        2.04
                                 3.94
                                          12.02 1.00
                                                         5257
                                                                  4131
age
Further Distributional Parameters:
      Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
sigma
          4.30
                    1.87
                             2.13
                                      9.10 1.00
                                                     3621
                                                              3481
Draws were sampled 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).
```

plot(fit2)



We can check the defaults for any model with default_prior(). The model does not have to be fitted, just model formula and data must be specified.

prior	class	coef	group	resp	dpar	nlpar	lb	ub	source
(flat)	b								default
(flat)	b	age							(vectorized)
student_t(3, 110, 11.9)	Intercept								default
student_t(3, 0, 11.9)	sigma						0		default

Alternatively, you can display the priors of any fitted model. Since we had not specified any priors, both outputs are the same here.

```
prior_summary(fit2)
```

prior	class	coef	group	resp	dpar	nlpar	lb u	b source
(flat)	b							default
(flat)	b	age						(vectorized)
student_t(3, 110, 11.9)	Intercept							default
student_t(3, 0, 11.9)	sigma						0	default

This table can be a bit confusing, but look at the column class: b is for effects / slopes. The first line tells you if there is a prior used for ALL effects, which is not the case (prior=flat). Second line is the prior for a specific coefficient (coef=age), there's also no prior specified.

But brms chooses a prior for Intercept and for the residual sdev sigma. These are automatically generated from the mean and the spread of the response. Note that internally, the brms machine uses mean-centered predictors. The Intercept parameter (and its prior) are based on mean-centered variables. What's displayed in the model summary is actually b_Intercept which is the intercept parameter transformed to the original, non mean-centered scale.

A short form is presented in the summary for prior=TRUE

summary(fit2, prior=TRUE)

```
Family: gaussian
  Links: mu = identity; sigma = identity
Formula: weight ~ age
   Data: data (Number of observations: 7)
  Draws: 4 chains, each with iter = 5000; warmup = 2500; thin = 1;
         total post-warmup draws = 10000
Priors:
Intercept ~ student t(3, 110, 11.9)
<lower=0> sigma ~ student_t(3, 0, 11.9)
Regression Coefficients:
          Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
                                -17.79
                                          68.66 1.00
                                                         5245
Intercept
             26.25
                       21.61
                                                                  4110
              7.89
                        2.04
                                  3.94
                                          12.02 1.00
                                                         5257
                                                                   4131
age
Further Distributional Parameters:
      Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk ESS Tail ESS
sigma
          4.30
                    1.87
                              2.13
                                       9.10 1.00
                                                     3621
                                                              3481
Draws were sampled 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).
```

Unless necessary, I would leave the brms defaults for Intercept & sigma. However, you should choose a prior for the slope, which currently has none.

This would set a prior for all slopes (if you have >1 predictors)

my_priors = prior(normal(5,1), class=b)

For setting a prior for a specific predictor, you specify it in **coef**. Since this model only has 1 predictor, both formulations are the same.

```
summary(fit3, prior=TRUE)
```

)

```
Family: gaussian
Links: mu = identity; sigma = identity
Formula: weight ~ age
Data: data (Number of observations: 7)
Draws: 4 chains, each with iter = 5000; warmup = 2500; thin = 1;
total post-warmup draws = 10000
Priors:
b_age ~ normal(5, 1)
```

```
Intercept ~ student_t(3, 110, 11.9)
<lower=0> sigma ~ student_t(3, 0, 11.9)
```

Regression Coefficients:

	Estimate	Est.Error	1-95% CI	u-95% C	I Rhat	Bulk_ESS	Tail_ESS
Intercept	49.08	10.00	30.68	69.5	4 1.00	6562	6790
age	5.74	0.93	3.84	7.4	8 1.00	6627	6763

Further Distributional Parameters:Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESSsigma 4.59 1.75 2.36 9.06 1.00 4814 5306

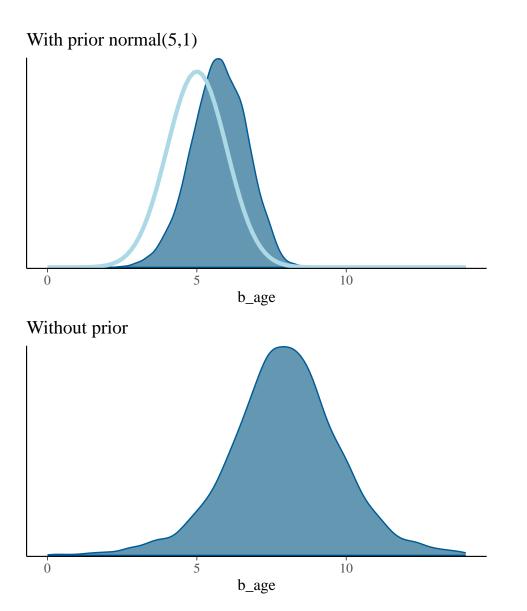
Draws were sampled using sampling(NUTS). For each parameter, Bulk_ESS and Tail_ESS are effective sample size measures, and Rhat is the potential scale reduction factor on split chains (at convergence, Rhat = 1).

We can draw prior & posterior in 1 plot by using mcmc_dens to plot the posterior distribution and adding the prior distribution (which we specified with normal(5,1) earlier). We also draw the old posterior without a slope-prior for comparison

```
plot3 = mcmc_dens(fit3, pars=c("b_age"))
plot3 = plot3 +
  geom_function(fun=dnorm, args=list(mean=5, sd=1), colour="lightblue", linewidth=1.5) +
  xlim(0,14) +
  ggtitle("With prior normal(5,1)")

plot1 = mcmc_dens(fit1, pars=c("b_age"))
plot1 = plot1 +
  xlim(0,14) +
  ggtitle("Without prior")

plot_grid(plot3, plot1, nrow=2)
```



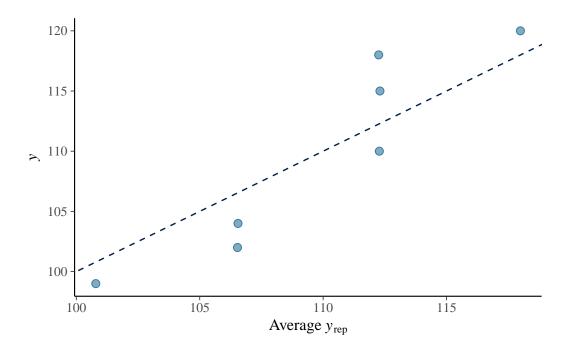
Here we only have a small dataset and an informative prior (mean=5) changes the posterior estimate of the slope.

Model analysis

Only after we checked MCMC convergence, we can go to the next step: Model evaluation / model checking. How well does our model describe the data?

A classical visual tool is observed vs predicted, which also works if you have multiple predictors.

pp_check(fit3, "scatter_avg")



Using all posterior draws for ppc type 'scatter_avg' by default.

bayes_R2 is the amount of explained variation. Its computation is a bit different from the classical frequentist R2, but conceptually it means the same.

bayes_R2(fit3)

Estimate Est.Error Q2.5 Q97.5 R2 0.680979 0.129089 0.3647646 0.8557636

More on that tomorrow, e.g. checking model assumptions.

Inference

OK, so we know that (a) MCMC converged and (b) model describes the data well. Only now can we make inference, i.e. quantitative statements about research questions. The summary already tells us mean and 95% confidence intervals for the slope (growth per year of age).

Different Credible intervals can be chosen in the summary, e.g. 90%-CI. 90% of posterior samples for slope were in this interval, we are 90% sure that the slope is in this interval.

summary(fit3, prob=0.90)

Family: gaussian Links: mu = identity; sigma = identity Formula: weight ~ age Data: data (Number of observations: 7) Draws: 4 chains, each with iter = 5000; warmup = 2500; thin = 1; total post-warmup draws = 10000 Regression Coefficients: Estimate Est.Error 1-90% CI u-90% CI Rhat Bulk_ESS Tail_ESS 33.04 6790 Intercept 49.08 10.00 65.99 1.00 6562 5.74 0.93 4.16 7.24 1.00 6627 6763 age Further Distributional Parameters: Estimate Est.Error 1-90% CI u-90% CI Rhat Bulk_ESS Tail_ESS 1.75 2.56 7.96 1.00 sigma 4.59 4814 5306 Draws were sampled 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).

Or you can extract specific quantiles of parameter estimates. fixef means "fixed effects" here.

fixef(fit3)

EstimateEst.ErrorQ2.5Q97.5Intercept49.0792229.996813030.67589769.541695age5.7413180.92776823.8420577.475878

fixef(fit3, probs=c(0.25, 0.5, 0.75))

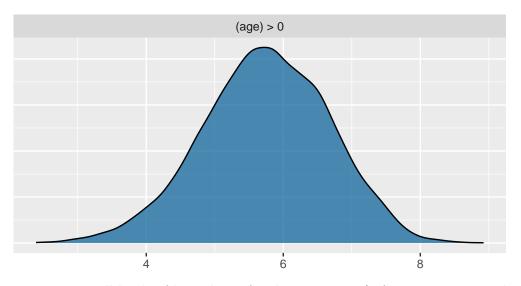
Estimate Est.ErrorQ25Q50Q75Intercept49.0792229.996813042.04393048.89011155.640038age5.7413180.92776825.1274925.7616616.395498

In a frequentist analysis you would want to know if the effect of age is "significant": p-values quantify the probability of observing such a pattern the data if the null hypothesis (b_age=0) was true (p small -> reject H0). Here, we can just calculate the probability that the slope is positive, $P(b_age > 0)$, with the hypothesis function. The column Post.Prob is the value of interest. It's =1 because all samples of slope were positive

hypothesis(fit3, "age>0")

```
Hypothesis Tests for class b:
    Hypothesis Estimate Est.Error CI.Lower CI.Upper Evid.Ratio Post.Prob Star
1 (age) > 0 5.74 0.93 4.16 7.24 Inf 1 *
---
'CI': 90%-CI for one-sided and 95%-CI for two-sided hypotheses.
'*': For one-sided hypotheses, the posterior probability exceeds 95%;
for two-sided hypotheses, the value tested against lies outside the 95%-CI.
Posterior probabilities of point hypotheses assume equal prior probabilities.
```

```
plot(hypothesis(fit3, "age>0"))
```



You can test all kinds of hypotheses for the parameters! If we were interested in the question if growth per year is bigger than 4, $P(b_age > 4)$, just put it in the hypothesis. The function is quite powerful and can handle all kinds of transformations of parameters.

hypothesis(fit3, "age>4")

```
Hypothesis Tests for class b:

Hypothesis Estimate Est.Error CI.Lower CI.Upper Evid.Ratio Post.Prob Star

1 (age)-(4) > 0 1.74 0.93 0.16 3.24 27.01 0.96 *
```

```
---

'CI': 90%-CI for one-sided and 95%-CI for two-sided hypotheses.

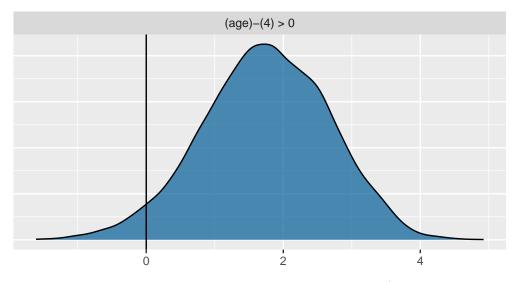
'*': For one-sided hypotheses, the posterior probability exceeds 95%;

for two-sided hypotheses, the value tested against lies outside the 95%-CI.

Posterior probabilities of point hypotheses assume equal prior probabilities.
```

It's transformed in the equivalent formulation age-4>0, and this is the probability distribution which is actually plotted.

```
plot1 = plot(hypothesis(fit3, "age>4"), plot=FALSE)
plot1[[1]] + geom_vline(xintercept=0)
```



The posterior probability is 0.96, which is also the integral (area under the curve) right of zero (age-4>0)

Exercise 1: Survival rate

Population counts from different habitats before and after winter.

Question: Is the average survival rate bigger than 2/3?

 $\begin{array}{ll} \text{Deterministic part:} & \mu = \theta, \quad \theta \in [0,1] \\ \text{Stochastic part:} & survived_i \sim \text{Binomial}(total_i,\mu) \end{array}$

Check the default prior!

Choose a meaningful prior for the Intercept parameter, use lb=0, ub=1. Fit the model & verify convergence. Re-run the analysis for different priors.

Intercept ~ student_t(3, 18, 3)

This brms default prior choice does not make any sense (mean=18), since it is chosen from the mean of the response survived (I guess??), while the parameter is actually a rate / probability $0 < \theta < 1$. Here brms messed up because we have overwritten the default link="logit" of the binomial distribution (again, I guess??). In most cases the default prior is fine, but better check it for generalized linear models (see also part 5 on GLMs).

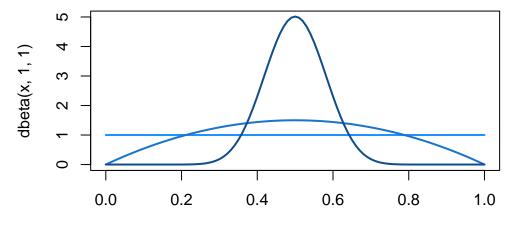
We use 3 different beta distribution priors (defined on interval [0,1]). Two shape parameters s_1, s_2 describe concentration to its mean $s_1/(s_1 + s_2)$

(1) beta(1,1) = uniform distribution

(2) beta(2,2) = weak prior, mean=0.5

(3) beta(20,20) = informative prior, mean=0.5

curve(dbeta(x,1,1), ylim=c(0,5), col="dodgerblue", lwd=2)
curve(dbeta(x,2,2), add=TRUE, col="dodgerblue3", lwd=2)
curve(dbeta(x,20,20), add=TRUE, , col="dodgerblue4", lwd=2)



Х

```
family = binomial(link="identity"),
prior = prior(beta(20,20), class=Intercept, lb=0, ub=1),
data = data)
```

summary(fit4)

```
Family: binomial
 Links: mu = identity
Formula: survived | trials(total) ~ 1
   Data: data (Number of observations: 10)
  Draws: 4 chains, each with iter = 2000; warmup = 1000; thin = 1;
         total post-warmup draws = 4000
Regression Coefficients:
         Estimate Est.Error 1-95% CI u-95% CI Rhat Bulk_ESS Tail_ESS
                    0.03
                                 0.65
                                         0.77 1.00
Intercept
              0.71
                                                        1518
                                                                 1824
Draws were sampled 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).

Here, the prior has only little effect on the outcome:

fixef(fit4)

Estimate Est.Error Q2.5 Q97.5 Intercept 0.708836 0.02882774 0.6515685 0.7655459 fixef(fit5)

Estimate Est.Error Q2.5 Q97.5 Intercept 0.7077716 0.02844394 0.6496433 0.7619462

fixef(fit6)

Estimate Est.Error Q2.5 Q97.5 Intercept 0.6818631 0.02743494 0.6242358 0.7356291

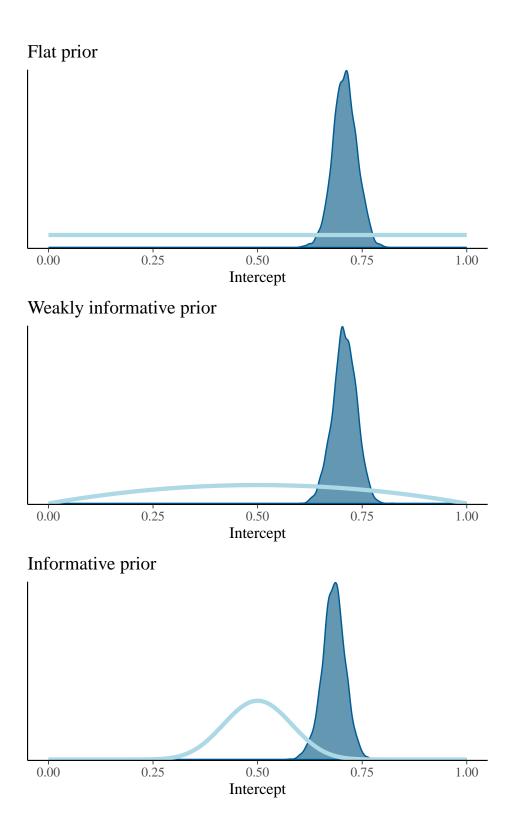
Plot prior & posterior for the different models:

```
plot1 = mcmc_dens(fit4, pars=c("Intercept"))
plot1 = plot1 +
   geom_function(fun=dbeta, args=list(1, 1), colour="lightblue", linewidth=1.5) +
   xlim(0,1) +
   ggtitle("Flat prior")

plot2 = mcmc_dens(fit5, pars=c("Intercept"))
plot2 = plot2 +
   geom_function(fun=dbeta, args=list(2, 2), colour="lightblue", linewidth=1.5) +
   xlim(0,1) +
   ggtitle("Weakly informative prior")

plot3 = mcmc_dens(fit6, pars=c("Intercept"))
plot3 = plot3 +
   geom_function(fun=dbeta, args=list(20, 20), colour="lightblue", linewidth=1.5) +
   xlim(0,1) +
   ggtitle("Informative prior")
```

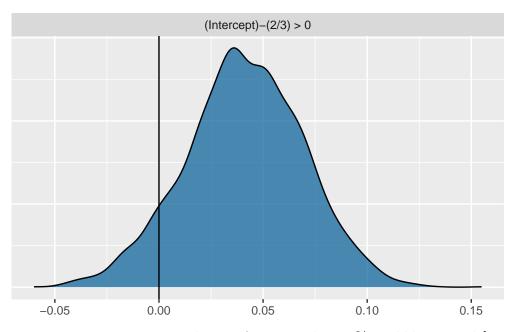
plot_grid(plot1, plot2, plot3, nrow=3)



We use the weakly informative prior model to test is survival rate is >2/3

```
hypothesis(fit5, "Intercept>2/3")
```

```
plot1 = plot(hypothesis(fit5, "Intercept>2/3"), plot=FALSE)
plot1[[1]] + geom_vline(xintercept=0)
```



We estimate a mean survival rate of 0.708 with a 95% credible interval [0.649,0.762]. Also, posterior probability $P(\theta > 2/3) = 0.92$, which means we are only 92% certain that average survival rate is larger than 2/3.