

ASDA3 Analysis Examples Replication Chapter 5

R Code

```
# data production prior to analysis examples
# need ncsr part1 part2 and population weight data
# nhanes subset of >=18 years of age
# hrs hh weight and financial respondents
# hrs respondent weight for individual level analyses
# can create subsets at the design data set level

setwd("P:/ASDA3/Replication R/Chapter 5")
getwd()

# data production
# load packages
library(survey)
library(haven)

#nhanes first
# example of how to read in from SAS, not used here
#nhanesdata <- read_sas("P:\\ASDA3\\Replication SAS\\nhanes1112_f.sas7bdat")

# load nhanes1112.rdata
load("P:/ASDA3/Data Sets for Analysis Examples and Stata R Code/nhanes1112.rdata")
nhanesdata <- nhanes1112
summary(nhanesdata)

#create factor variables
nhanesdata$racec <- factor(nhanesdata$ridreth1, levels = 1: 5 , labels =c("Mexican", "Other Hispanic", "White",
"Black", "Other"))
nhanesdata$marcatc <- factor(nhanesdata$marcat, levels = 1: 3, labels =c("Married", "Previously Married", "Never
Married"))
nhanesdata$edcatc <- factor(nhanesdata$edcat, levels = 1: 4, labels =c("0-11", "12", "13-15", "16+"))
nhanesdata$bp_catc <- factor(nhanesdata$bp_cat, levels = 1: 4, labels =c("Normal", "Pre-HBP", "Stage 1
HBP", "Stage 2 HBP"))
#nhanesdata$agescq <- (nhanesdata$age * nhanesdata$age)

names(nhanesdata)

nhanessvy2 <- svydesign(strata=~sdmvstra, id=~sdmvpsu, weights=~wtmec2yr, data=nhanesdata, nest=T)
subnhanes <- subset(nhanessvy2 , age >= 18)
names (nhanessvy2)

##
#ncs-r next

load("P:/ASDA3/Data Sets for Analysis Examples and Stata R Code/ncsr.rdata")
names(ncsr)
summary(ncsr)

#create factor versions with labels
ncsr$racec <- factor(ncsr$racecat, levels = 1: 4, labels =c("Other", "Hispanic", "Black", "White"))
ncsr$mar3catc <- factor(ncsr$mar3cat, levels = 1: 3, labels =c("Married", "Previously Married", "Never
Married"))
ncsr$ed4catc <- factor(ncsr$ed4cat, levels = 1: 4, labels =c("0-11", "12", "13-15", "16+"))
ncsr$sexc <- factor(ncsr$sex, levels = 1:2, labels=c("Male", "Female"))
ncsr$ag4catc <- factor(ncsr$ag4cat, levels = 1:4, labels=c("18-29", "30-44", "45-59", "60+"))
ncsr$mdec <- factor(ncsr$mde, level = 1:2, labels=c("No", "Yes"))
```

```

# part 1 data n=9282
ncsrsvyp1 <- svydesign(strata=~sestrat, id=~seclustr, weights=~ncsrwtsh, data=ncsr, nest=T)
names (ncsrsvyp1)

# part 2 data n=5692
ncsrp2 <- subset(ncsr, !is.na(ncsrwtlg))
ncsrsvyp2 <- svydesign(strata=~sestrat, id=~seclustr, weights=~ncsrwtlg, data=ncsrp2, nest=T)
names (ncsrsvyp2)

# rescale weight to sum to population in 2001
ncsr$popweight <- (ncsr$ncsrwtsh*(209128094/9282))

ncsrsvypop <- svydesign(strata=~sestrat, id=~seclustr, weights=~popweight, data=ncsr, nest=T)
summary(ncsrsvypop)

##
#hrs 2012 final data from SAS
#both hh and r weights are needed plus financial respondent for hh level analysis

#library(haven)
load ("P:/ASDA3/Data Sets for Analysis Examples and Stata R Code/hrs12.rdata")
names(hrs12)

hrssvyhh <- svydesign(strata=~stratum, id=~secu, weights=~nwgthh , data=hrs12, nest=T)
summary(hrssvyhh)

hrssvysub <-subset(hrssvyhh, nfinr==1)
summary(hrssvysub)

hrssvyr <- svydesign(strata=~stratum, id=~secu, weights=~nwgtr , data=hrs12, nest=T)
summary(hrssvyr)

# Figures 5.1 and 5.2
svyhist(~lbxtc, subset (nhanessvy2, age >=18), main="", col="grey80", xlab ="Histogram of Total Cholesterol")

# create a variable called gender for boxplot
nhanessvy2<-update(nhanessvy2, gender=cut(riagendr, c(1, 2, Inf), right=F))

svyboxplot(lbxtc~gender , subset (nhanessvy2, age >=18), col="grey80", ylab="Total Cholesterol", xlab ="1=Male
2=Female")

# Example 5.3
svytotal (~mde, ncsrsvypop, deff=T)
confint(svytotal(~mde, ncsrsvypop))

# Example 5.3 MDE over marital status
ex53 <- svyby (~mde, ~mar3catc, ncsrsvypop, svytotal)
ex53 <- svyby (~mde, ~mar3catc, ncsrsvypop, svytotal, deff=T)
ex53
confint(ex53)

# Example 5.4 HRS HH Level Wealth/Total Assets
svyby (~h11atota, ~I(nfinr==1), hrssvyhh, na.rm=T, svytotal)
confint(svyby (~h11atota, ~I(nfinr==1), hrssvyhh, na.rm=T, ci=T, svytotal))

# Example 5.5 HRS HH Income
svyby (~h11itot, ~I(nfinr==1), hrssvyhh, na.rm=T, svymean)

```

```

confint(svyby (~h11itot, ~I(nfinr==1), hrssvyhh, na.rm=T, ci=T, svymean))

# Example 5.5 using alternative Bayesian approach
# load all relevant libraries for csSampling

install.packages("devtools") # if you have not installed "devtools" package

# install_github("RyanHornby/csSampling") # lots of packages needed, will take a second, already loaded here

library(csSampling)
library(rstan)
library(brms)
library(survey)
rstan_options(auto_write = TRUE)

# read in the full HRS data set for the third edition
load("P:\\ASDA3\\Data Sets for Analysis Examples and Stata R Code\\hrs12.rdata")

# create complete data set with variables of interest
hrs.red <- hrs12[hrs12$nfinr == 1, c("h11itot", "secu", "stratum", "nwgthh")]
hrs.red <- hrs.red[complete.cases(hrs.red),]

# need to normalize HRS weights to match what is done for Stan modeling
hrs.red$wtsc <- hrs.red$nwgthh / mean(hrs.red$nwgthh)

# survey design object
hrs.des <- svydesign(id = ~secu, strata = ~stratum, weights = ~wtsc, nest = T, data = hrs.red)

# Bayesian approach, flat prior (non-informative), which is default
set.seed(41279)
model_formula <- formula("h11itot|weights(wtsc) ~ 1")
mod.brms <- cs_sampling_brms(svydes = hrs.des, brmsmod = brmsformula(model_formula), data = hrs.red, family =
gaussian())

# Print results
mod.brms$stan_fit

# With Normal Prior, mean = 70000, SD (SE) = 1000
mod.brms2 <- cs_sampling_brms(svydes = hrs.des, brmsmod = brmsformula(model_formula), prior =
c(set_prior("normal(70000,1000)", class = "Intercept")), data = hrs.red, family = gaussian())

# Example 5.6 Mean Systolic Blood Pressure, NHANES data
a <- svymean(~bpxsy1, subset (nhanessvy2, age >=18), na.rm=TRUE)
coef(a)
SE(a)
confint(a)

# Example 5.6 alternative Bayesian approach
# create complete data set with variables of interest
nhanes.red <- nhanes1112[nhanes1112$age >= 18 & !is.na(nhanes1112$age), c("bpxsy1", "sdmvpsu", "sdmvstra",
"wtmec2yr")]
nhanes.red <- nhanes.red[complete.cases(nhanes.red),]

# need to normalize NHANES weights to match what is done for Stan modeling
nhanes.red$wtsc <- nhanes.red$wtmec2yr / mean(nhanes.red$wtmec2yr)

# survey design object
nhanes.des <- svydesign(id = ~sdmvpsu, strata = ~sdmvstra, weights = ~wtsc, nest = T, data = nhanes.red)

```

```

# Bayesian approach, flat prior
set.seed(41279)
model_formula <- formula("bpxsy1|weights(wtsc) ~ 1")
mod.brms <- cs_sampling_brms(svydes = nhanes.des, brmsmod = brmsformula(model_formula), data = nhanes.red,
family = gaussian())

# Print results
mod.brms$stan_fit

# Example 5.7
svyby (~h11atota, ~I(nfinr==1), hrssvyhh, na.rm=T, ci=T, svymean)
confint(svyby(~h11atota, ~I(nfinr==1), hrssvyhh, na.rm=T, ci=T, svymean))

# Example 5.7 Alternative Bayesian Approach
# create complete data set with variables of interest
hrs.red <- hrs12[hrs12$nfinr == 1, c("h11atota", "secu", "stratum", "nwgthh")]
hrs.red <- hrs.red[complete.cases(hrs.red),]

# need to normalize HRS weights to match what is done for Stan modeling
hrs.red$wtsc <- hrs.red$nwgthh / mean(hrs.red$nwgthh)

# survey design object
hrs.des <- svydesign(id = ~secu, strata = ~stratum, weights = ~wtsc, nest = T, data = hrs.red)

# With Normal Prior, mean = 400000, SD (SE) = 20000
set.seed(41279)
model_formula <- formula("h11atota|weights(wtsc) ~ 1")
mod.brms <- cs_sampling_brms(svydes = hrs.des, brmsmod = brmsformula(model_formula), prior =
c(set_prior("normal(400000,20000)", class = "Intercept")), data = hrs.red, family = gaussian())

# Print results
mod.brms$stan_fit

# Plot posteriors, figure 5.4
plot(mod.brms, varnames = "Intercept")

# Example 5.8 Standard Deviation of Total HH Wealth HRS data
# create a data object with weights only but no design variables
hrswgt <- svydesign(id=~1, weights=~nwgthh, data=hrs12)
summary(hrswgt)

# obtain mean
a <- svymean(~h11atota, design=hrswgt, na.rm=T, deff="replace")
a

# use sqrt of variance to obtain standard deviation
sd <- sqrt(svyvar(~h11atota, design = hrswgt, na.rm=T))
sd

# refer back to Example 5.7 for Bayesian Results from intercept only regression, not shown again here

# Example 5.9 Population Percentiles for total HH Wealth HRS data, in subset of nfinr=1
q <- svyquantile(~h11atota, hrssvysub, c(.25,.5,.75), na.rm=T, ci=T)
q
# Obtain SE from confidence intervals, see R documentation for details
SE(q)

```

```

# Example 5.10 Lorenz Curve and GINI coefficient not available in R Survey Package

# Example 5.10 Available in "convey" package, requires vardpoor too

library(vardpoor)
library(convey)

# linearized design, use hrssvsub created previously
hrssvyhh_c <- convey_prep(hrssvyhh)

# now can subset to financial respondents (after convey_prep)
sub_hrssvyhh_c <- subset( hrssvyhh_c , nfinr==1)

# run svygini and svylorenz using subset, note that R does not require negative set to 0 as Stata does
svygini( ~h11atota, design = sub_hrssvyhh_c)

# note that the results will differ from Stata due to negative values set to 0 in Stata but not R
svylorenz( ~h11atota, sub_hrssvyhh_c, seq(0,1,.1), alpha = .01 )

# Fig 5.5 example 5.11 Relationship between 2 continuous variables, note this is weighted and design based
svyplot(lbdhdd ~ lbxtc, subset(subnhanes, age >= 18), style="bubble", ylab="HDL", xlab="Total Cholesterol")

# Fig 5.5 with survey-weighted LOWESS smoother
smoother <- svsmooth(lbdhdd ~ lbxtc, subset(subnhanes, age >= 18), bandwidth=10)
# add smooth curve to plot
lines(smoother, col="black", lwd=4, lty=1)

# Example 5.11 Correlation between Total and High Cholesterol, NHANES DATA
# create standardized versions of variables first, then use in regression
nhanesdata$stdlbxtc <- (nhanesdata$lbxtc - 194.4355)/41.05184
summary(nhanesdata$lbxtc + nhanesdata$stdlbxtc)

nhanesdata$stdlbdhdd <- (nhanesdata$lbdhdd - 52.83826)/14.93157
summary(nhanesdata$stdlbxtc)

#reset survey design and subset
nhanessvy2 <- svydesign(strata=~sdmvstra, id=~sdmvpsu, weights=~wtmec2yr, data=nhanesdata, nest=T)
subnhanes <- subset(nhanessvy2 , age >= 18)

#Design based linear regression to obtain correlation and correct SE
summary(Ex5_11_svyglm <- svyglm(stdlbxtc ~ stdlbdhdd, design=subnhanes))
confint(Ex5_11_svyglm)

# Example 5.12 Ratio Estimator for HDD to Total Cholesterol
ex5_12 <- svyby (~lbdhdd, denominator=~lbxtc, by=~I(age >= 18), nhanessvy2, na.rm=T, ci=T, svyratio)
ex5_12
confint(ex5_12)

# Example 5.13 Proportions of Diabetes by Gender in Subpopulation of Age >=70
subhrs70 <- subset(hrssvyr, nage >= 70)
ex5_13 <- svyby(~diabetes, ~gender, subhrs70, svymean, keep.names=T, na.rm=T)
print(ex5_13)
confint(ex5_13)

```

```

# Example 5.14 Mean Systolic Blood Pressure by Gender, Age 46+ NHANES
subnhanes46 <- subset(nhanessvy2, age >= 46)
# riagendr 1=Male 2=Female
ex5_14 <- svyby(~bpxsy1, ~riagendr, subnhanes46, svymean, keep.names=T, DEFF=T, na.rm=T)
print(ex5_14)
confint(ex5_14)

# Example 5.15 Differences in Mean HH Wealth by Educational Attainment, HRS data
# Codes for edcat : 1=0-11 2=12 3=13-15 4=16+ years of education
summary(hrssvysub)

ex5_15 <- svyby(~h11atota, ~edcat, hrssvysub, svymean, na.rm=T)
print(ex5_15)
confint(ex5_15)
svycontrast(ex5_15, list(avg=c(.5,0,0,.5), diff=c(1,0,0,-1)))

# Example 5.16 Differences in Total Wealth over Time 2010 to 2012, HRS data
# Use 2010 and 2012 data set prepared in SAS

library(haven)
# use sas data input
hrs_2010_2012 <- read_sas("P:/ASDA3/Replication SAS/hrs_2010_2012_both_f.sas7bdat")
summary(hrs_2010_2012)
names(hrs_2010_2012)

hrs2010_2012 <- svydesign(strata=~stratum, id=~secu, weights=~hhweight, data=hrs_2010_2012, nest=T)
subhrs2010 <- subset(hrs2010_2012, finr2010_2012==1)

ex5_16 <- svyby (~totwealth, ~year, design=subhrs2010, keep.vars=T, svymean)
# get coefficients
coef(ex5_16)
# get standard errors
se(ex5_16)

# use svycontrast
contrast <- svycontrast(ex5_16, list(avg=c(.5,.5), diff=c(1,-1)))
print(contrast)

```

R Results

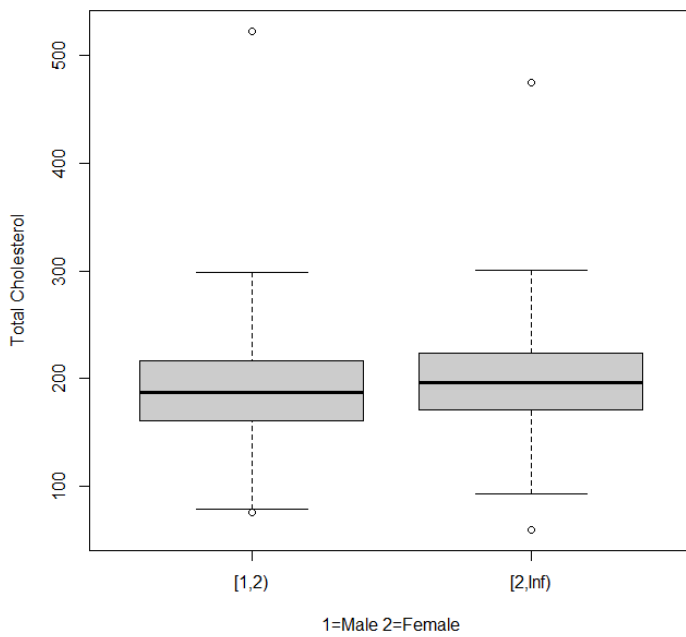
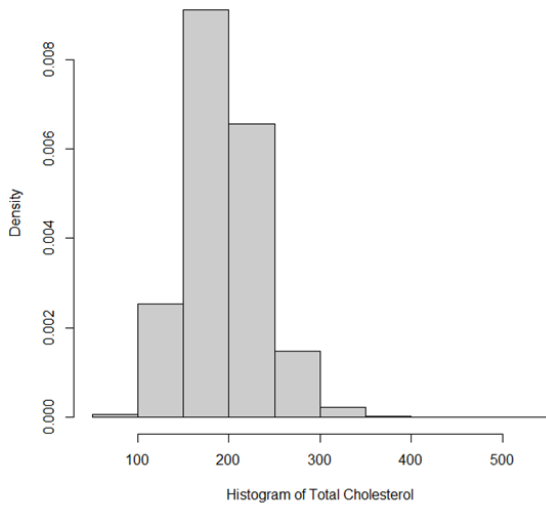
```
# figures 5.1 and 5.2
```

```
svyhist(~lbxtc, subset (nhanessvy2, age >=18), main="", col="grey80", xlab ="Histogram of Total Cholesterol")
```

```
#Create a variables called gender for boxplot
```

```
nhanessvy2<-update(nhanessvy2, gender=cut(riagendr, c(1, 2, Inf), right=F))
```

```
svyboxplot(lbxtc~gender , subset (nhanessvy2, age >=18), col="grey80", ylab="Total Cholesterol", xlab ="1=Male  
2=Female")
```



```

> #Example 5.3
> svytotal (~mde, ncsrsvypop, deff=T)
      total      SE  DEff
mde 40092207 2567488 9.028
> confint(svytotal(~mde, ncsrsvypop))
      2.5 %   97.5 %
mde 35060023 45124390
>
> #Example 5.3 MDE over marital status
> ex53 <- svyby (~mde, ~mar3catc, ncsrsvypop, svytotal)
> ex53 <- svyby (~mde, ~mar3catc, ncsrsvypop, svytotal, deff=T)
> ex53
      mar3catc      mde      se DEff.mde
Married      Married 20304191 1584108.6 6.817920
Previously Married Previously Married 10360671 702621.5 2.966192
Never Married      Never Married 9427345 773137.6 3.063915
> confint(ex53)
      2.5 %   97.5 %
Married      17199395 23408986
Previously Married 8983558 11737783
Never Married      7912024 10942667

> #Example 5.4 HRS HH Level Wealth/Total Assets
> svyby (~H11ATOTA, ~I(nfinr==1), hrssvyhh, na.rm=T, svytotal)
      I(nfinr == 1)  H11ATOTA      se
FALSE      FALSE 1.701334e+13 1.031603e+12
TRUE      TRUE 2.526686e+13 1.353710e+12
> confint(svyby (~H11ATOTA, ~I(nfinr==1), hrssvyhh, na.rm=T, ci=T, svytotal))
      2.5 %   97.5 %
FALSE 1.499144e+13 1.903525e+13
TRUE 2.261364e+13 2.792009e+13

> #Example 5.5 HRS HH Income
> svyby (~H11ITOT, ~I(nfinr==1), hrssvyhh, na.rm=T, svymean)
      I(nfinr == 1)  H11ITOT      se
FALSE      FALSE 98737.91 3007.883
TRUE      TRUE 71382.40 1937.229
> confint(svyby (~H11ITOT, ~I(nfinr==1), hrssvyhh, na.rm=T, ci=T, svymean))
      2.5 %   97.5 %
FALSE 92842.57 104633.3
TRUE 67585.50 75179.3

```

```
> Example 5.5 Bayesian Approach
# Print results
> mod.brms$stan_fit
Inference for Stan model: anon_model.
1 chains, each with iter=2000; warmup=1000; thin=1;
post-warmup draws per chain=1000, total post-warmup draws=1000.
```

	mean	se_mean	sd	2.5%	25%	50%
Intercept	71338.32	50.82	1071.99	69280.69	70581.08	71356.31
sigma	122158.21	23.03	733.41	120675.49	121675.94	122159.07
lprior	-27.03	0.00	0.05	-27.13	-27.07	-27.03
b_Intercept	71338.32	50.82	1071.99	69280.69	70581.08	71356.31
lp__	-186370.73	0.06	1.06	-186373.56	-186371.16	-186370.40

	75%	97.5%	n_eff	Rhat
Intercept	72051.2	73460.27	445	1
sigma	122674.3	123567.19	1014	1
lprior	-27.0	-26.93	499	1
b_Intercept	72051.2	73460.27	445	1
lp__	-186370.0	-186369.70	365	1

Samples were drawn using NUTS(diag_e) at Tue Mar 11 09:02:41 2025.
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).

```
> # With Normal Prior, mean = 70000, SD (SE) = 1000
> mod.brms2 <- cs_sampling_brms(svydes = hrs.des, brmsmod = brmsformula(model_formula), prior =
c(set_prior("normal(70000,1000)", class = "Intercept")), data = hrs.red, family = gaussian())
[1] "compiling stan model"
recompiling to avoid crashing R session
[1] "stan fitting"
```

```
SAMPLING FOR MODEL 'anon_model' NOW (CHAIN 1).
Chain 1:
Chain 1: Gradient evaluation took 0.00427 seconds
Chain 1: 1000 transitions using 10 leapfrog steps per transition would take 42.7 seconds.
Chain 1: Adjust your expectations accordingly!
Chain 1:
Chain 1:
Chain 1: Iteration: 1 / 2000 [ 0%] (Warmup)
Chain 1: Iteration: 200 / 2000 [ 10%] (Warmup)
Chain 1: Iteration: 400 / 2000 [ 20%] (Warmup)
Chain 1: Iteration: 600 / 2000 [ 30%] (Warmup)
Chain 1: Iteration: 800 / 2000 [ 40%] (Warmup)
Chain 1: Iteration: 1000 / 2000 [ 50%] (Warmup)
Chain 1: Iteration: 1001 / 2000 [ 50%] (Sampling)
Chain 1: Iteration: 1200 / 2000 [ 60%] (Sampling)
Chain 1: Iteration: 1400 / 2000 [ 70%] (Sampling)
Chain 1: Iteration: 1600 / 2000 [ 80%] (Sampling)
Chain 1: Iteration: 1800 / 2000 [ 90%] (Sampling)
Chain 1: Iteration: 2000 / 2000 [100%] (Sampling)
Chain 1:
Chain 1: Elapsed Time: 105.602 seconds (Warm-up)
Chain 1: 14.096 seconds (Sampling)
Chain 1: 119.698 seconds (Total)
Chain 1:
[1] "gradient evaluation"
Warning message:
```

```
In mrbweights(design$cluster, design$strata, design$fpc, ...) :
```

```
Design is sampled with replacement: only first stage used
```

```
> mod.brms2$stan_fit
```

```
Inference for Stan model: anon_model.
```

```
1 chains, each with iter=2000; warmup=1000; thin=1;
```

```
post-warmup draws per chain=1000, total post-warmup draws=1000.
```

	mean	se_mean	sd	2.5%	25%	50%
Intercept	70639.61	27.83	714.29	69248.30	70175.67	70623.06
sigma	122149.39	27.22	691.39	120831.13	121682.20	122145.02
lprior	-22.93	0.02	0.58	-24.55	-23.12	-22.71
b_Intercept	70639.61	27.83	714.29	69248.30	70175.67	70623.06
lp__	-186366.52	0.05	0.97	-186369.12	-186366.87	-186366.22

	75%	97.5%	n_eff	Rhat
Intercept	71118.65	72034.69	659	1.00
sigma	122574.12	123551.22	645	1.00
lprior	-22.52	-22.46	605	1.00
b_Intercept	71118.65	72034.69	659	1.00
lp__	-186365.83	-186365.60	394	1.01

```
Samples were drawn using NUTS(diag_e) at Tue Mar 11 09:06:19 2025.
```

```
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).
```

```
> #Example 5.6 Mean Systolic Blood Pressure, NHANES data
```

```
> a <- svymean(~BPXSY1 , subset (nhanessvy2, age >=18), na.rm=TRUE)
```

```
> coef(a)
```

```
BPXSY1
```

```
122.0292
```

```
> SE(a)
```

```
BPXSY1
```

```
BPXSY1 0.6163389
```

```
> confint(a)
```

```
2.5 % 97.5 %
```

```
BPXSY1 120.8212 123.2372
```

```

> Example 5.6
> # Bayesian approach, flat prior
> set.seed(41279)
> model_formula <- formula("bpxsy1|weights(wtsc) ~ 1")
> mod.brms <- cs_sampling_brms(svydes = nhanes.des, brmsmod = brmsformula(model_formula), data = nhanes.red,
family = gaussian())
[1] "compiling stan model"
recompiling to avoid crashing R session
[1] "stan fitting"

SAMPLING FOR MODEL 'anon_model' NOW (CHAIN 1).
Chain 1:
Chain 1: Gradient evaluation took 0.001451 seconds
Chain 1: 1000 transitions using 10 leapfrog steps per transition would take 14.51 seconds.
Chain 1: Adjust your expectations accordingly!
Chain 1:
Chain 1:
Chain 1: Iteration:    1 / 2000 [ 0%] (Warmup)
Chain 1: Iteration:   200 / 2000 [ 10%] (Warmup)
Chain 1: Iteration:   400 / 2000 [ 20%] (Warmup)
Chain 1: Iteration:   600 / 2000 [ 30%] (Warmup)
Chain 1: Iteration:   800 / 2000 [ 40%] (Warmup)
Chain 1: Iteration:  1000 / 2000 [ 50%] (Warmup)
Chain 1: Iteration:  1001 / 2000 [ 50%] (Sampling)
Chain 1: Iteration:  1200 / 2000 [ 60%] (Sampling)
Chain 1: Iteration:  1400 / 2000 [ 70%] (Sampling)
Chain 1: Iteration:  1600 / 2000 [ 80%] (Sampling)
Chain 1: Iteration:  1800 / 2000 [ 90%] (Sampling)
Chain 1: Iteration:  2000 / 2000 [100%] (Sampling)
Chain 1:
Chain 1: Elapsed Time: 5.301 seconds (Warm-up)
Chain 1:                4.441 seconds (Sampling)
Chain 1:                9.742 seconds (Total)
Chain 1:
[1] "gradient evaluation"
Warning message:
In mrbweights(design$cluster, design$strata, design$fpc, ...) :
  Design is sampled with replacement: only first stage used
>
> # Print results
> mod.brms$stan_fit
Inference for Stan model: anon_model.
1 chains, each with iter=2000; warmup=1000; thin=1;
post-warmup draws per chain=1000, total post-warmup draws=1000.

      mean se_mean  sd      2.5%      25%      50%
Intercept  122.02   0.01 0.24    121.54    121.86    122.02
sigma      17.37   0.01 0.17     17.05    17.26    17.38
lprior     -7.47   0.00 0.01     -7.49    -7.48    -7.47
b_Intercept 122.02   0.01 0.24    121.54    121.86    122.02
lp__      -21937.91  0.05 1.02   -21940.46 -21938.30 -21937.61
      75%      97.5% n_eff Rhat
Intercept  122.18  122.46 1001    1
sigma      17.49   17.72  957    1
lprior     -7.46   -7.44  974    1
b_Intercept 122.18  122.46 1001    1
lp__      -21937.19 -21936.95 412    1
> #Example 5.7

```

```

> svyby (~H11ATOTA, ~I(nfinr==1), hrssvyhh, na.rm=T, ci=T, svymean)
      I(nfinr == 1) H11ATOTA      se
FALSE          FALSE 563269.1 26670.34
TRUE           TRUE  428470.8 17353.77
> confint(svyby(~H11ATOTA, ~I(nfinr==1), hrssvyhh, na.rm=T, ci=T, svymean))
      2.5 %   97.5 %
FALSE 510996.2 615542.0
TRUE  394458.0 462483.5

```

#Example 5.7 Using Alternative Bayesian Approach

```

> # With Normal Prior, mean = 400000, SD (SE) = 20000
> # Print results
> mod.brms$stan_fit
Inference for Stan model: anon_model.
1 chains, each with iter=2000; warmup=1000; thin=1;
post-warmup draws per chain=1000, total post-warmup draws=1000.

```

	mean	se_mean	sd	2.5%	25%	50%
Intercept	423606.38	353.97	7955.77	407199.96	418380.45	423627.66
sigma	1113687.40	234.70	6466.77	1101343.40	1108952.54	1113625.82
lprior	-29.93	0.02	0.48	-31.14	-30.19	-29.85
b_Intercept	423606.38	353.97	7955.77	407199.96	418380.45	423627.66
lp__	-217737.15	0.04	0.86	-217739.44	-217737.53	-217736.89

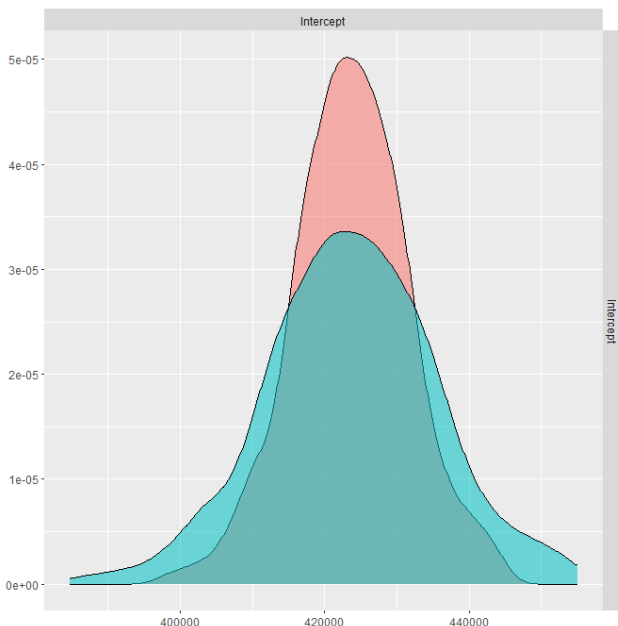
	75%	97.5%	n_eff	Rhat
Intercept	428870.98	439998.28	505	1.00
sigma	1118325.22	1125996.74	759	1.00
lprior	-29.58	-29.22	514	1.00
b_Intercept	428870.98	439998.28	505	1.00
lp__	-217736.53	-217736.26	482	1.01

Samples were drawn using NUTS(diag_e) at Tue Mar 11 09:25:04 2025.
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).

```

# Plot posteriors from previous model, intercept only
> plot(mod.brms, varnames = "Intercept")

```



```

#Example 5.8 Standard Deviation of Total HH Wealth HRS data
#Create a data object with weights only but no design variables
> hrswgt <- svydesign(id=~1, weights=~nwgthh, data=hrs)
> #obtain mean
> a <- svymean(~H11ATOTA, design=hrswgt, na.rm=T, deff="replace")
> a
      mean      SE  DEff
H11ATOTA 474129 11592 2.0536
> # use sqrt of variance to obtain standard deviation
> sd <- sqrt(svyvar(~H11ATOTA, design = hrswgt, na.rm=T))
> sd
      variance      SE
H11ATOTA 1143716 1.657e+11

```

```

#Example 5.9 Population Percentiles for total HH Wealth HRS data, in subset of nfinr=1
> q <- svyquantile(~H11ATOTA, hrssvysub, c(.25,.5,.75), na.rm=T, ci=T)
> q
$H11ATOTA
  quantile ci.2.5 ci.97.5      se
0.25    22000  18000   26900 2221.401
0.5     142000 127000  158000 7737.463
0.75    440000 404500  480000 18844.465

attr("hasci")
[1] TRUE
attr("class")
[1] "newsvyquantile"
> # Obtain SE from confidence intervals, see R documentation for details
> SE(q)
H11ATOTA.0.25 H11ATOTA.0.5 H11ATOTA.0.75
      2221.401      7737.463      18844.465

```

```

> #Example 5.10 Lorenz Curve and GINI coefficient not available in R Survey Package

```

```

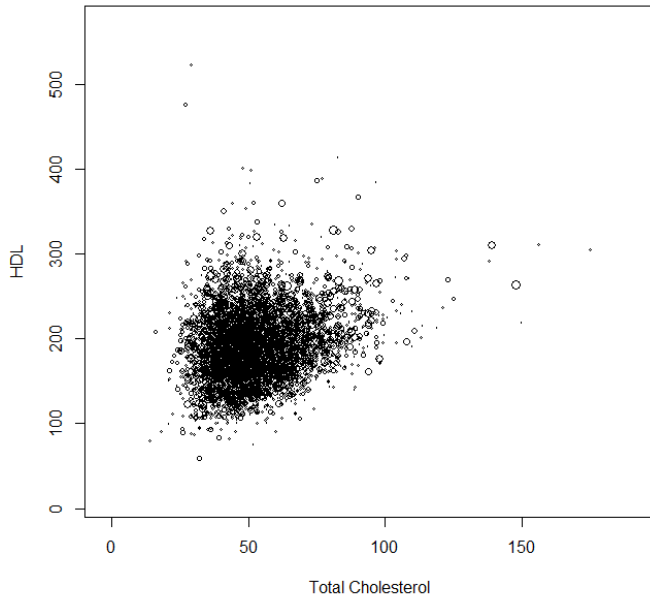
> #Example 5.10 Available in "convey" package
> library(vardpoor)
> library(convey)
> # linearized design, use hrssvysub created previously
> hrssvyhh_c <- convey_prep(hrssvyhh)
> # now can subset to financial respondents (after convey_prep)
> sub_hrssvyhh_c <- subset( hrssvyhh_c , nfinr==1)
> # run svygini and svylorenz using subset, note that R does not require negative set to 0 as Stata
> svygini( ~H11ATOTA, design = sub_hrssvyhh_c)
      gini      SE
H11ATOTA 0.73897 0.0094
> svylorenz( ~H11ATOTA, sub_hrssvyhh_c, seq(0,1,.1), alpha = .01 )
      lorenz      SE
L(0)    0.0000000 0.0000
L(0.1) -0.0071723 0.0008
L(0.2) -0.0065605 0.0009
L(0.3) -0.0012259 0.0011
L(0.4)  0.0131167 0.0018

```

```
L(0.5) 0.0388491 0.0029
L(0.6) 0.0811900 0.0046
L(0.7) 0.1464493 0.0068
L(0.8) 0.2498023 0.0099
L(0.9) 0.4234323 0.0140
```

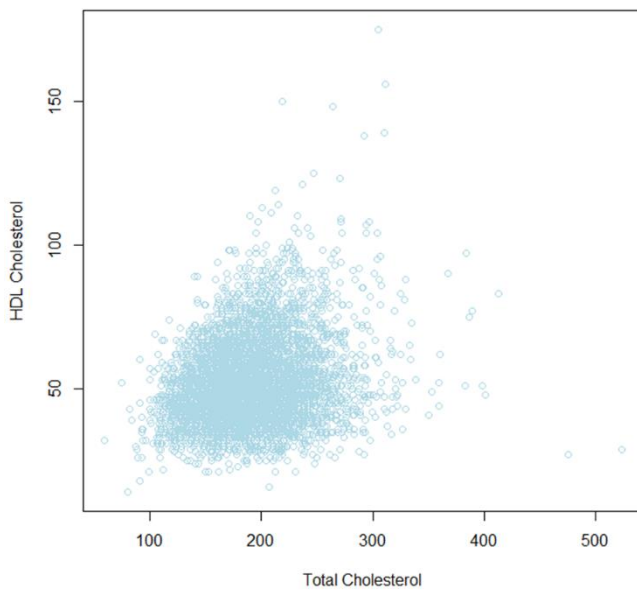
#Example 5.11, Figure 5.5 Plot of Relationship between 2 continuous variables, note this is weighted and design based

```
> svyplot(lbxtc~lbdhdd, subset(subnhanes, age>=18), style="bubble", ylab="HDL", xlab="Total Cholesterol")
```



```
> # unweighted scatter plot
```

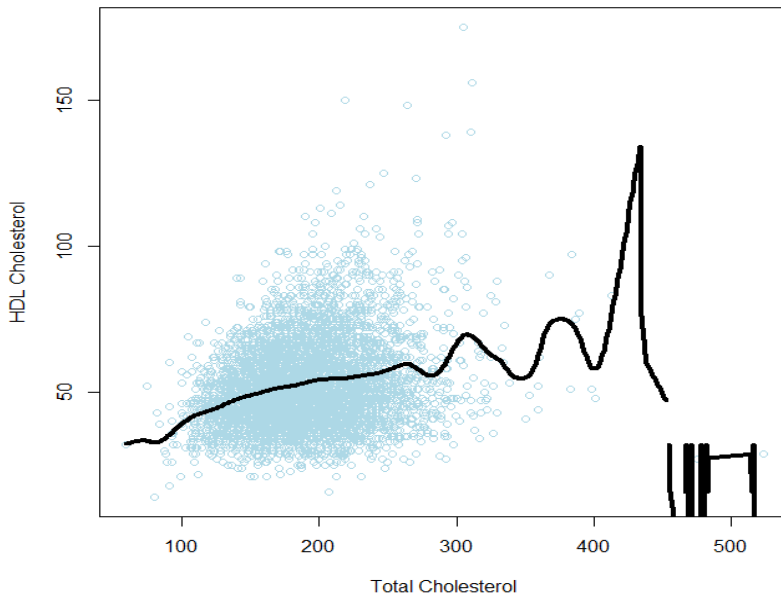
```
> plot(lbdhdd ~ lbxtc, subset(nhanes1112, age >= 18), xlab="Total Cholesterol", ylab="HDL Cholesterol", col="lightblue")
```



```

> # survey-weighted LOWESS smoother
> smoother <- svsmooth(lbdhdd ~ lbxtc, subset(nhanes.des, age >= 18), bandwidth=10)
> # add smooth curve to plot
> lines(smoother, col="black", lwd=4, lty=1)

```



#EXAMPLE 5.11 Correlation between Total and High Cholesterol, NHANES Data

```

> summary(nhanesdata$lbxtc + nhanesdata$stdlbxtc)
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   NA's
  55.7  153.0  178.6  182.9  209.4  531.0  2768

> #nhanesdata$stdlbdhdd <- (nhanesdata$LBDHDD-52.83826)/14.93157
> summary(nhanesdata$stdlbxtc)
  Min. 1st Qu.  Median    Mean 3rd Qu.    Max.   NA's
-3.2991 -0.9850 -0.3760 -0.2737  0.3548  8.0036  2768

> #reset survey design and subset
> nhanessvy2 <- svydesign(strata=~sdmvsra, id=~sdmvpsu, weights=~WTMEC2YR, data=nhanesdata, nest=T)
> subnhanes <- subset(nhanessvy2 , age >= 18)

> #Design based linear regression to obtain correlation and correct SE
> summary(Ex5_11_svyglm <- svyglm(stdlbxtc ~ stdlbdhdd, design=subnhanes))
Call:
svyglm(formula = stdlbxtc ~ stdlbdhdd, design = subnhanes)

```

Survey design:

```
subset(nhanessvy2, age >= 18)
```

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	-8.916e-07	2.568e-02	0.00	1
stdlbdhdd	2.414e-01	1.104e-02	21.87	2.4e-13 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
(Dispersion parameter for gaussian family taken to be 1.003306)
Number of Fisher Scoring iterations: 2

```

#Example 5.12 Ratio Estimator for HDD to Total Cholesterol
> ex5_12 <- svyby(~lbdhdd, denominator=~ltxtc, by=~I(age >= 18), nhanessvy2, na.rm=T, ci=T, svyratio)
> confint(ex5_12)
      2.5 %    97.5 %
FALSE 0.3265713 0.3374500
TRUE  0.2660798 0.2774246

```

```

#Example 5.13 Proportions of Diabetes by Gender in Subpopulation of Age >=70
> subhrs70 <- subset(hrssvyr, nage >= 70)
> ex5_13 <- svyby(~diabetes, ~gender, subhrs70, svymean, keep.names=T, na.rm=T)
> print(ex5_13)
  gender diabetes      se
1      1 0.2736113 0.007468441
2      2 0.2269743 0.008554564
> confint(ex5_13)
      2.5 %    97.5 %
1 0.2589734 0.2882491
2 0.2102077 0.2437409
>

```

```

> #Example 5.14 Mean Systolic Blood Pressure by Gender, Age 46+ NHANES
> subnhanes46 <-subset(nhanessvy2, age >= 46)
>
> #riagendr 1=M 2=F
> ex5_14 <- svyby(~BPXSY1, ~riagendr, subnhanes46, svymean, keep.names=T, na.rm=T)
> print(ex5_14)
  riagendr BPXSY1      se
1      1 128.3005 0.8687054
2      2 128.1820 0.9460163
> confint(ex5_14)
      2.5 %    97.5 %
1 126.5979 130.0032
2 126.3278 130.0361

```

```

#Example 5.14 Mean Systolic Blood Pressure by Gender, Age 46+ NHANES
> subnhanes46 <-subset(nhanessvy2, age >= 46)
>
> #riagendr 1=M 2=F
> ex5_14 <- svyby(~BPXSY1, ~riagendr, subnhanes46, svymean, keep.names=T, na.rm=T)
> print(ex5_14)
  riagendr BPXSY1      se
1      1 128.3005 0.8687054
2      2 128.1820 0.9460163
> confint(ex5_14)
      2.5 %    97.5 %
1 126.5979 130.0032
2 126.3278 130.0361

```

```

> #Example 5.15 Differences in Mean HH Wealth by Educational Attainment, HRS data
> #CODES FOR EDCAT: 1=0-11 2=12 3=13-15 4=16+ YEARS OF EDUCATION
> summary(hrssvsub)
Stratified 1 - level Cluster Sampling design (with replacement)
With (112) clusters.
subset(hrssvyhh, nfinr == 1)
Probabilities:
      Min.   1st Qu.   Median     Mean   3rd Qu.     Max.
5.594e-05 1.959e-04 3.110e-04      Inf 5.961e-04      Inf
Stratum Sizes:
      1  2  3  4  5  6  7  8  9 10 11 12 13 14 15 16
obs      257 409 247 228 194 245 281 275 253 228 185 253 134 105 125 96
design.PSU 2  2  2  2  2  2  2  2  2  2  2  2  2  2  2  2
actual.PSU 2  2  2  2  2  2  2  2  2  2  2  2  2  2  2  2
      17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32
obs      192 123 204 273 233 129 77 52 61 217 424 308 350 364 364 314
design.PSU 2  2  2  2  2  2  2  2  2  2  2  2  2  2  2  2
actual.PSU 2  2  2  2  2  2  2  2  2  2  2  2  2  2  2  2
      33 34 35 36 37 38 39 40 41 42 43 44 45 46 47
obs      405 339 217 358 136 258 242 454 345 349 498 445 406 467 308
design.PSU 2  2  2  2  2  2  2  2  2  2  2  2  2  2  2
actual.PSU 2  2  2  2  2  2  2  2  2  2  2  2  2  2  2
      48 49 50 51 52 53 54 55 56
obs      203 244 463 387 111 96 40 124 96
design.PSU 2  2  2  2  2  2  2  2  2
actual.PSU 2  2  2  2  2  2  2  2
Data variables:
 [1] "hhid"      "pn"          "R11BMI"      "nage"        "nfinr"
 [6] "gender"    "secu"        "stratum"     "nwgthh"      "nwgtr"
[11] "H11ATOTA"  "H11ITOT"    "marcat"      "edcat"       "racecat"
[16] "diabetes"  "numfalls24" "age65p"      "arthritis"   "finr"
[21] "female"    "age70"
> ex5_15 <- svyby(~H11ATOTA, ~edcat, hrssvsub, svymean, na.rm=T)
> print(ex5_15)
  edcat H11ATOTA      se
1     1 122088.6 10595.60
2     2 259027.2  9802.47
3     3 336308.6 17201.79
4     4 834141.0 46477.79
> confint(ex5_15)
  2.5 %  97.5 %
1 101321.6 142855.6
2 239814.7 278239.6
3 302593.7 370023.5
4 743046.2 925235.8
> svycontrast(ex5_15, list(avg=c(.5,0,0,.5), diff=c(1,0,0,-1)))
  contrast      SE
avg    478115 23835
diff   -712052 47670

```

```
# Example 5.15 Alternative Bayesian approach
> # Bayesian approach, flat prior
> set.seed(41279)
> model_formula <- formula("h11atota|weights(wtsc) ~ edcat1")
> mod.brms <- cs_sampling_brms(svydes = hrs.des, brmsmod = brmsformula(model_formula), data = hrs.red, family =
gaussian())
[1] "compiling stan model"
[1] "stan fitting"
```

SAMPLING FOR MODEL 'anon_model' NOW (CHAIN 1).

Chain 1:

Chain 1: Gradient evaluation took 0.002207 seconds

Chain 1: 1000 transitions using 10 leapfrog steps per transition would take 22.07 seconds.

Chain 1: Adjust your expectations accordingly!

Chain 1:

Chain 1:

Chain 1: Iteration: 1 / 2000 [0%] (Warmup)

Chain 1: Iteration: 200 / 2000 [10%] (Warmup)

Chain 1: Iteration: 400 / 2000 [20%] (Warmup)

Chain 1: Iteration: 600 / 2000 [30%] (Warmup)

Chain 1: Iteration: 800 / 2000 [40%] (Warmup)

Chain 1: Iteration: 1000 / 2000 [50%] (Warmup)

Chain 1: Iteration: 1001 / 2000 [50%] (Sampling)

Chain 1: Iteration: 1200 / 2000 [60%] (Sampling)

Chain 1: Iteration: 1400 / 2000 [70%] (Sampling)

Chain 1: Iteration: 1600 / 2000 [80%] (Sampling)

Chain 1: Iteration: 1800 / 2000 [90%] (Sampling)

Chain 1: Iteration: 2000 / 2000 [100%] (Sampling)

Chain 1:

Chain 1: Elapsed Time: 112.688 seconds (Warm-up)

Chain 1: 7.492 seconds (Sampling)

Chain 1: 120.18 seconds (Total)

Chain 1:

[1] "gradient evaluation"

Warning message:

In mrbweights(design\$cluster, design\$strata, design\$fpc, ...) :

Design is sampled with replacement: only first stage used

>

> # Print results

> mod.brms\$stan_fit

Inference for Stan model: anon_model.

1 chains, each with iter=2000; warmup=1000; thin=1;

post-warmup draws per chain=1000, total post-warmup draws=1000.

	mean	se_mean	sd	2.5%	25%	50%
b[1]	-714328.45	1382.52	39318.31	-787398.12	-742169.73	-713721.08
Intercept	489282.21	643.77	18720.77	455206.21	476119.83	488701.43
sigma	1475791.70	385.55	12459.57	1451900.58	1467600.99	1475073.37
lprior	-34.73	0.00	0.14	-35.00	-34.82	-34.72
b_Intercept	831857.96	605.38	22082.74	788225.76	817499.64	831468.02
lp__	-99090.33	0.05	1.16	-99093.46	-99090.83	-99090.01
	75%	97.5%	n_eff	Rhat		
b[1]	-687158.32	-638514.87	809	1		
Intercept	501496.04	527016.88	846	1		
sigma	1484257.31	1499886.78	1044	1		
lprior	-34.63	-34.46	815	1		
b_Intercept	847382.64	874227.85	1331	1		
lp__	-99089.50	-99089.02	506	1		

Samples were drawn using NUTS(diag_e) at Tue Mar 11 09:55:42 2025.
 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).

```
> # With informative prior for coefficient
> mod.brms <- cs_sampling_brms(svydes = hrs.des, brmsmod = brmsformula(model_formula), prior =
c(set_prior("normal(-700000,40000)", class = "b", coef="edcat1")), data = hrs.red, family = gaussian())
[1] "compiling stan model"
[1] "stan fitting"
```

SAMPLING FOR MODEL 'anon_model' NOW (CHAIN 1).

Chain 1:

Chain 1: Gradient evaluation took 0.002221 seconds

Chain 1: 1000 transitions using 10 leapfrog steps per transition would take 22.21 seconds.

Chain 1: Adjust your expectations accordingly!

Chain 1:

Chain 1:

```
Chain 1: Iteration: 1 / 2000 [ 0%] (Warmup)
Chain 1: Iteration: 200 / 2000 [ 10%] (Warmup)
Chain 1: Iteration: 400 / 2000 [ 20%] (Warmup)
Chain 1: Iteration: 600 / 2000 [ 30%] (Warmup)
Chain 1: Iteration: 800 / 2000 [ 40%] (Warmup)
Chain 1: Iteration: 1000 / 2000 [ 50%] (Warmup)
Chain 1: Iteration: 1001 / 2000 [ 50%] (Sampling)
Chain 1: Iteration: 1200 / 2000 [ 60%] (Sampling)
Chain 1: Iteration: 1400 / 2000 [ 70%] (Sampling)
Chain 1: Iteration: 1600 / 2000 [ 80%] (Sampling)
Chain 1: Iteration: 1800 / 2000 [ 90%] (Sampling)
Chain 1: Iteration: 2000 / 2000 [100%] (Sampling)
```

Chain 1:

Chain 1: Elapsed Time: 107.866 seconds (Warm-up)

Chain 1: 6.471 seconds (Sampling)

Chain 1: 114.337 seconds (Total)

Chain 1:

[1] "gradient evaluation"

Warning message:

In mrbweights(design\$cluster, design\$strata, design\$fpc, ...) :

Design is sampled with replacement: only first stage used

> mod.brms\$stan_fit

Inference for Stan model: anon_model.

1 chains, each with iter=2000; warmup=1000; thin=1;

post-warmup draws per chain=1000, total post-warmup draws=1000.

	mean	se_mean	sd	2.5%	25%	50%
b[1]	-706037.28	938.40	28602.24	-767783.07	-725412.31	-706001.45
Intercept	491640.72	547.15	17964.01	455484.21	479337.62	492298.13
sigma	1476180.16	355.25	12712.81	1451441.85	1467936.16	1475660.19
lprior	-46.53	0.02	0.40	-47.73	-46.65	-46.43
b_Intercept	830240.21	541.82	20544.48	793277.62	814591.42	829873.99
lp__	-99101.91	0.05	1.18	-99104.88	-99102.42	-99101.63
	75%	97.5%	n_eff	Rhat		
b[1]	-686439.35	-651279.80	929	1		
Intercept	503305.84	525090.92	1078	1		
sigma	1484201.16	1503216.29	1281	1		
lprior	-46.30	-46.06	451	1		

```
b_Intercept 845749.59 866541.10 1438 1
lp__        -99101.05 -99100.58 511 1
```

Samples were drawn using NUTS(diag_e) at Tue Mar 11 10:00:35 2025.

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`).

```
#Example 5.16 Differences in Total Wealth over Time 2010 to 2012, HRS data
> ex5_16 <- svyby(~totwealth, ~year, design=subhrs2010, keep.vars=T, svymean)
> coef(ex5_16)
      2010      2012
432829.6 437807.6
> SE(ex5_16)
[1] 16010.53 17016.29
> contrast <- svycontrast(ex5_16, list(avg=c(.5,.5), diff=c(1,-1)))
Warning message:
In vcov.svyby(stat) : Only diagonal elements of vcov() available
> print(contrast)
      contrast      SE
avg 435318.6 11682
diff -4978.1 23364
> confint(contrast)
      2.5 %      97.5 %
avg 412421.98 458215.21
diff -50771.29 40815.16
```