

MR example for Supplementary Material

Example R and Stata code to perform a multivariable Mendelian randomization (MVMR) analysis

R code

- Read in the data

```
suppressPackageStartupMessages({  
  library(tidyverse)  
  library(haven)  
})  
dat <- read_dta("https://raw.githubusercontent.com/remlapmot/mrrobust/master/dodata.dta")  
dat <- dat %>% filter(ldlcp2 < 1e-8)
```

Example code using the MendelianRandomization package

- Install the package and load it into memory

```
library(MendelianRandomization)
```

- Convert our data frame to the required class

```
datfmt <- mr_mvinput(  
  bx = as.matrix(cbind(dat$ldlcbeta, dat$hdlcbeta, dat$tgbeta)),  
  bxse = as.matrix(cbind(dat$ldlcse, dat$hdlcse, dat$tgse)),  
  by = dat$chdbeta,  
  byse = dat$chdse,  
  exposure = "exposure",  
  outcome = "outcome",  
  snps = "snps",  
  effect_allele = dat$a1,  
  other_allele = dat$a2,  
  eaf = NA  
)
```

- Fit an MVMR/MVIVW model

```
mvivwfit <- mr_mvivw(datfmt)  
mvivwfit
```

```
##
## Multivariable inverse-variance weighted method
## (variants uncorrelated, random-effect model)
##
## Number of Variants : 73
##
## -----
##      Exposure Estimate Std Error  95% CI      p-value
## exposure_1      0.429      0.061  0.309, 0.548    0.000
## exposure_2     -0.194      0.131 -0.451, 0.062    0.138
## exposure_3      0.226      0.123 -0.016, 0.468    0.067
## -----
## Residual standard error =  1.490
## Heterogeneity test statistic = 155.3766 on 70 degrees of freedom, (p-value = 0.0000)
```

- Fit an MVMR-Egger model

```
mvmreggerfit <- mr_mvegger(datfmt)
mvmreggerfit
```

```
##
## Multivariable MR-Egger method
## (variants uncorrelated, random-effect model)
##
## Orientated to exposure : 1
## Number of Variants : 73
##
## -----
##      Exposure Estimate Std Error  95% CI      p-value
## exposure_1      0.567      0.100  0.371, 0.764    0.000
## exposure_2     -0.136      0.133 -0.398, 0.125    0.306
## exposure_3      0.274      0.125  0.030, 0.518    0.028
## (intercept)    -0.009      0.005 -0.020, 0.001    0.084
## -----
## Residual standard error =  1.469
## Heterogeneity test statistic = 148.9290 on 69 degrees of freedom, (p-value = 0.0000)
```

Example code using the MVMR package

- Install the package and load it into memory

```
library(MVMR)
```

- Create a data object of the required structure

```
r_input <- format_mvnr(
  BXGs = dat[,c("ldlcbeta", "hdlcbeta", "tgbeta")],
  BYG = dat$chdbeta,
  seBXGs = dat[,c("ldlcse", "hdlcse", "tgse")],
  seBYG = dat$chdse,
  RSID = dat$rsid
)
```

- Fit an MVMR model

```
mvmrfit <- ivw_mvvr(r_input)
```

```
##
## Multivariable MR
##
##           Estimate Std. Error   t value    Pr(>|t|)
## exposure1  0.4286200  0.0609661   7.030464 1.099077e-09
## exposure2 -0.1941989  0.1308289  -1.484372 1.421994e-01
## exposure3  0.2260456  0.1232828   1.833554 7.097168e-02
##
## Residual standard error: 1.49 on 70 degrees of freedom
```

- Heterogeneity statistic

```
strength_mvvr(r_input)
```

```
## Warning in strength_mvvr(r_input): Covariance between effect of genetic variants on each exposure
## not specified. Fixing covariance at 0.
```

```
##
## Conditional F-statistics for instrument strength
##
##           exposure1 exposure2 exposure3
## F-statistic  126.7447  35.29937  39.32731

##           exposure1 exposure2 exposure3
## F-statistic  126.7447  35.29937  39.32731
```

Stata code

- Load the Statamarkdown package to enable Stata code chunks in an R Markdown file

```
library(Statamarkdown)
```

- Read in the data and create an indicator variable to select observations with p -value between the genotype and LDL-C $< 10^{-8}$

```
use https://raw.githubusercontent.com/remlapmot/mrrobust/master/dodata, clear
gen byte sel1 = (ldlcp2 < 1e-8)
```

Example code using the mrrobust package

- Install the mrrobust package using the github package

```
// Note: output suppressed
net install mrrobust, from("https://raw.githubusercontent.com/remlapmot/mrrobust/master/")
mrdeps
```

- Fit and MVMR model with phenotypes LDL-c and HDL-c (Burgess, Dudbridge, and Thompson 2015).

```
mvmr chdbeta ldldbeta hdlcbeta [aw=1/(chdse^2)] if sel1==1
```

```

                                     Number of genotypes = 73
                                     Number of phenotypes = 2
                                     Standard errors: Random effect
                                     Residual standard error = 1.514
-----
|          Coef.   Std. Err.      z    P>|z|    [95% Conf. Interval]
-----+-----
chdbeta |
  ldldbeta |   .4670719   .0581901    8.03   0.000    .3530214   .5811224
  hdlcbeta |  -.2930048   .1211822   -2.42   0.016   -.5305175  -.0554921
-----
```

- Additionally include a third phenotype – triglycerides.

```
mvmr chdbeta ldldbeta hdlcbeta tgbeta [aw=1/(chdse^2)] if sel1==1
```

```

                                     Number of genotypes = 73
                                     Number of phenotypes = 3
                                     Standard errors: Random effect
                                     Residual standard error = 1.490
-----
|          Coef.   Std. Err.      z    P>|z|    [95% Conf. Interval]
-----+-----
chdbeta |
  ldldbeta |   .42862   .0609661    7.03   0.000    .3091286   .5481113
  hdlcbeta |  -.1941989   .1308289   -1.48   0.138   -.4506189   .0622211
  tgbeta |   .2260456   .1232828    1.83   0.067   -.0155842   .4676755
-----
```

- Report the QA statistic for instrument validity and the conditional F-statistics for instrument strength for each phenotype (Sanderson et al. 2019; Sanderson, Spiller, and Bowden 2020).

```
mvmr chdbeta ldldbeta hdlcbeta tgbeta [aw=1/(chdse^2)] if sel1==1, gxse(ldlcse hdlcse tgse)
```

```
> e hdlcse tgse)
```

```

                                     Number of genotypes = 73
                                     Number of phenotypes = 3
                                     Standard errors: Random effect
                                     Residual standard error = 1.490
-----
|          Coef.   Std. Err.      z    P>|z|    [95% Conf. Interval]
-----+-----
```

chdbeta							
ldlcbeta		.42862	.0609661	7.03	0.000	.3091286	.5481113
hdlcbeta		-.1941989	.1308289	-1.48	0.138	-.4506189	.0622211
tgbeta		.2260456	.1232828	1.83	0.067	-.0155842	.4676755

Q_A statistic for instrument validity; chi2(70) = 152.88 (p = 0.0000)

Conditional F-statistics for instrument strength:

F_x1 = 130.31 (ldlcbeta)

F_x2 = 36.29 (hdlcbeta)

F_x3 = 40.44 (tgbeta)

- Fit an MVMR-Egger regression (Rees, Wood, and Burgess 2017), orienting the model with respect to the first phenotype in the main *varlist*.

```
mrmvegger chdbeta ldlcbeta hdlcbeta tgbeta [aw=1/(chdse^2)] if sel1==1
```

MVMR-Egger model oriented wrt: ldlcbeta

Number of genotypes = 73

Number of phenotypes = 3

Residual standard error = 1.469

		Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
chdbeta							
ldlcbeta		.5672993	.1002611	5.66	0.000	.370791	.7638075
hdlcbeta		-.1364113	.1332727	-1.02	0.306	-.3976209	.1247983
tgbeta		.2739803	.1246927	2.20	0.028	.0295871	.5183735
_cons		-.0093655	.0054187	-1.73	0.084	-.019986	.001255

We can also orient the model with respect to HDL-C.

```
mrmvegger chdbeta ldlcbeta hdlcbeta tgbeta [aw=1/(chdse^2)] if sel1==1, orient(2)
```

```
> t(2)
```

MVMR-Egger model oriented wrt: hdlcbeta

Number of genotypes = 73

Number of phenotypes = 3

Residual standard error = 1.501

		Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
chdbeta							
ldlcbeta		.4286398	.0614056	6.98	0.000	.308287	.5489926
hdlcbeta		-.1989637	.1541909	-1.29	0.197	-.5011723	.1032449
tgbeta		.2256794	.1243221	1.82	0.069	-.0179875	.4693463
_cons		.0002155	.0036218	0.06	0.953	-.006883	.0073141

Or we can orient the model with respect to triglycerides.

```
mrmmvegger chdbeta ldldbeta hdlcbeta tgbeta [aw=1/(chdse^2)] if sel1==1, orient(3)
```

```
> t(3)
```

MVMR-Egger model oriented wrt: tgbeta

Number of genotypes = 73

Number of phenotypes = 3

Residual standard error = 1.499

	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
chdbeta						
ldldbeta	.4203073	.0660026	6.37	0.000	.2909447	.54967
hdlcbeta	-.1903089	.1321536	-1.44	0.150	-.4493252	.0687075
tgbeta	.2065651	.1365427	1.51	0.130	-.0610537	.474184
_cons	.0013499	.003951	0.34	0.733	-.0063939	.0090936

R session information for reproducibility

```
library(sessioninfo)
session_info()
```

```
- Session info -----
setting value
version R version 4.1.1 (2021-08-10)
os Windows 10 x64
system x86_64, mingw32
ui RTerm
language (EN)
collate English_United Kingdom.1252
ctype English_United Kingdom.1252
tz Europe/London
date 2021-09-20
```

```
- Packages -----
package * version date lib source
arrangements 1.1.9 2020-09-13 [1] CRAN (R 4.1.0)
assertthat 0.2.1 2019-03-21 [1] CRAN (R 4.1.0)
backports 1.2.1 2020-12-09 [1] CRAN (R 4.1.0)
broom 0.7.9 2021-07-27 [1] CRAN (R 4.1.0)
cellranger 1.1.0 2016-07-27 [1] CRAN (R 4.1.0)
cli 3.0.1 2021-07-17 [1] CRAN (R 4.1.0)
codetools 0.2-18 2020-11-04 [2] CRAN (R 4.1.1)
colorspace 2.0-2 2021-06-24 [1] CRAN (R 4.1.0)
conquer 1.0.2 2020-08-27 [1] CRAN (R 4.1.0)
crayon 1.4.1 2021-02-08 [1] CRAN (R 4.1.0)
curl 4.3.2 2021-06-23 [1] CRAN (R 4.1.0)
data.table 1.14.0 2021-02-21 [1] CRAN (R 4.1.0)
DBI 1.1.1 2021-01-15 [1] CRAN (R 4.1.0)
dbplyr 2.1.1 2021-04-06 [1] CRAN (R 4.1.0)
```

DEoptimR	1.0-9	2021-05-24	[1]	CRAN	(R 4.1.0)
digest	0.6.27	2020-10-24	[1]	CRAN	(R 4.1.0)
dplyr	* 1.0.7	2021-06-18	[1]	CRAN	(R 4.1.0)
ellipsis	0.3.2	2021-04-29	[1]	CRAN	(R 4.1.0)
evaluate	0.14	2019-05-28	[1]	CRAN	(R 4.1.0)
fansi	0.5.0	2021-05-25	[1]	CRAN	(R 4.1.0)
fastmap	1.1.0	2021-01-25	[1]	CRAN	(R 4.1.0)
forcats	* 0.5.1	2021-01-27	[1]	CRAN	(R 4.1.0)
foreach	1.5.1	2020-10-15	[1]	CRAN	(R 4.1.0)
fs	1.5.0	2020-07-31	[1]	CRAN	(R 4.1.0)
generics	0.1.0	2020-10-31	[1]	CRAN	(R 4.1.0)
ggplot2	* 3.3.5	2021-06-25	[1]	CRAN	(R 4.1.0)
glmnet	4.1-2	2021-06-24	[1]	CRAN	(R 4.1.0)
glue	1.4.2	2020-08-27	[1]	CRAN	(R 4.1.0)
gmp	0.6-2	2021-01-07	[1]	CRAN	(R 4.1.0)
gtable	0.3.0	2019-03-25	[1]	CRAN	(R 4.1.0)
haven	* 2.4.3	2021-08-04	[1]	CRAN	(R 4.1.0)
hms	1.1.0	2021-05-17	[1]	CRAN	(R 4.1.0)
htmltools	0.5.2	2021-08-25	[1]	CRAN	(R 4.1.1)
htmlwidgets	1.5.4	2021-09-08	[1]	CRAN	(R 4.1.1)
httr	1.4.2	2020-07-20	[1]	CRAN	(R 4.1.0)
iterators	1.0.13	2020-10-15	[1]	CRAN	(R 4.1.0)
iterpc	0.4.2	2020-01-10	[1]	CRAN	(R 4.1.0)
jsonlite	1.7.2	2020-12-09	[1]	CRAN	(R 4.1.0)
knitr	1.34	2021-09-09	[1]	CRAN	(R 4.1.1)
lattice	0.20-44	2021-05-02	[2]	CRAN	(R 4.1.1)
lazyeval	0.2.2	2019-03-15	[1]	CRAN	(R 4.1.0)
lifecycle	1.0.0	2021-02-15	[1]	CRAN	(R 4.1.0)
lubridate	1.7.10	2021-02-26	[1]	CRAN	(R 4.1.0)
magrittr	2.0.1	2020-11-17	[1]	CRAN	(R 4.1.0)
Matrix	1.3-4	2021-06-01	[2]	CRAN	(R 4.1.1)
MatrixModels	0.5-0	2021-03-02	[1]	CRAN	(R 4.1.0)
matrixStats	0.61.0	2021-09-17	[1]	CRAN	(R 4.1.1)
MendelianRandomization	* 0.5.1	2021-04-16	[1]	CRAN	(R 4.1.0)
modelr	0.1.8	2020-05-19	[1]	CRAN	(R 4.1.0)
munsell	0.5.0	2018-06-12	[1]	CRAN	(R 4.1.0)
MVMR	* 0.3	2021-08-11	[1]	Github	(wspiller/mvmr@a6388a8)
pillar	1.6.2	2021-07-29	[1]	CRAN	(R 4.1.0)
pkgconfig	2.0.3	2019-09-22	[1]	CRAN	(R 4.1.0)
plotly	4.9.4.1	2021-06-18	[1]	CRAN	(R 4.1.0)
purrr	* 0.3.4	2020-04-17	[1]	CRAN	(R 4.1.0)
quantreg	5.86	2021-06-06	[1]	CRAN	(R 4.1.0)
R6	2.5.1	2021-08-19	[1]	CRAN	(R 4.1.1)
RadialMR	1.0	2021-07-12	[1]	Github	(WSpiller/RadialMR@d63d3fc)
Rcpp	1.0.7	2021-07-07	[1]	CRAN	(R 4.1.0)
readr	* 2.0.1	2021-08-10	[1]	CRAN	(R 4.1.1)
readxl	1.3.1	2019-03-13	[1]	CRAN	(R 4.1.0)
remotes	2.4.0	2021-06-02	[1]	CRAN	(R 4.1.0)
reprex	2.0.1	2021-08-05	[1]	CRAN	(R 4.1.0)
rjson	0.2.20	2018-06-08	[1]	CRAN	(R 4.1.0)
rlang	0.4.11	2021-04-30	[1]	CRAN	(R 4.1.0)
rmarkdown	2.11	2021-09-14	[1]	CRAN	(R 4.1.1)
robustbase	0.93-8	2021-06-02	[1]	CRAN	(R 4.1.0)
rstudioapi	0.13	2020-11-12	[1]	CRAN	(R 4.1.0)

rvest	1.0.1	2021-07-26	[1]	CRAN (R 4.1.0)
scales	1.1.1	2020-05-11	[1]	CRAN (R 4.1.0)
sessioninfo	* 1.1.1	2018-11-05	[1]	CRAN (R 4.1.0)
shape	1.4.6	2021-05-19	[1]	CRAN (R 4.1.0)
SparseM	1.81	2021-02-18	[1]	CRAN (R 4.1.0)
Statamarkdown	* 0.7.0	2021-09-15	[1]	Github (Hemken/Statamarkdown@a68a8b9)
stringi	1.7.4	2021-08-25	[1]	CRAN (R 4.1.1)
stringr	* 1.4.0	2019-02-10	[1]	CRAN (R 4.1.0)
survival	3.2-13	2021-08-24	[2]	CRAN (R 4.1.1)
tibble	* 3.1.4	2021-08-25	[1]	CRAN (R 4.1.1)
tidyr	* 1.1.3	2021-03-03	[1]	CRAN (R 4.1.0)
tidyselect	1.1.1	2021-04-30	[1]	CRAN (R 4.1.0)
tidyverse	* 1.3.1	2021-04-15	[1]	CRAN (R 4.1.0)
tzdb	0.1.2	2021-07-20	[1]	CRAN (R 4.1.0)
utf8	1.2.2	2021-07-24	[1]	CRAN (R 4.1.0)
vctrs	0.3.8	2021-04-29	[1]	CRAN (R 4.1.0)
viridisLite	0.4.0	2021-04-13	[1]	CRAN (R 4.1.0)
withr	2.4.2	2021-04-18	[1]	CRAN (R 4.1.0)
xfun	0.26	2021-09-14	[1]	CRAN (R 4.1.1)
xml2	1.3.2	2020-04-23	[1]	CRAN (R 4.1.0)
yaml	2.2.1	2020-02-01	[1]	CRAN (R 4.1.0)

[1] C:/Users/eptmp/Documents/R/win-library/4.1

[2] C:/Program Files/R/R-4.1.1/library

Stata session information for reproducibility

```
di c(version)
ado describe mrrobust
```

16.1

```
-----
[89] package mrrobust from https://raw.github.com/remlapmot/mrrobust/master
-----
```

TITLE

'mrrobust': Stata package for two-sample Mendelian randomization analyses

DESCRIPTION/AUTHOR(S)

Author: Tom Palmer

Distribution-Date: 20210917

INSTALLATION FILES

```
m\mrmedian.ado
m\mrmedian.sthlp
m\mrmedianobs.ado
m\mrmedianobs_work.ado
m\mrmedianobs.sthlp
m\mregger.ado
m\mregger.sthlp
```



```

m\mrrobust.sthlp
m\mreggerplot.ado
m\mreggerplot.sthlp
m\mrmodal.ado
m\mrmodal.sthlp
m\mrratio.ado
m\mrratio.sthlp
m\mrivests.ado
m\mrivests.sthlp
m\mrforest.ado
m\mrforest.sthlp
m\mreggersimex.ado
m\mreggersimex.sthlp
m\mreggersimexonce.ado
m\mrmodalplot.ado
m\mrmodalplot.sthlp
m\mrfunnel.ado
m\mrfunnel.sthlp
m\mrdeps.ado
m\mrdeps.sthlp
m\mr.ado
m\mr.sthlp
m\mrmvivw.ado
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m\mvivw.ado
m\mvivw.sthlp
m\mvmr.ado
m\mvmr.sthlp
m\mrmovegger.ado
m\mrmovegger.sthlp
m\mrleaveoneout.ado
m\mrleaveoneout.sthlp
m\mrrobust-author.ihlp

```

INSTALLED ON
 20 Sep 2021

References

- Burgess, S, F Dudbridge, and SG Thompson. 2015. "Multivariable Mendelian randomization: the use of pleiotropic genetic variants to estimate causal effects." *American Journal of Epidemiology* 181: 251--260. <https://doi.org/10.1093/aje/kwu283>.
- Rees, J, A Wood, and S Burgess. 2017. "Extending the MR-Egger method for multivariable Mendelian randomization to correct for both measured and unmeasured pleiotropy." *Statistics in Medicine* 36: 4705--18. <https://doi.org/10.1002/sim.7492>.
- Sanderson, E, G Davey Smith, F Windmeijer, and J Bowden. 2019. "An examination of multivariable Mendelian randomization in the single-sample and two-sample summary data settings." *International Journal of Epidemiology* 48: 713--27. <https://doi.org/10.1093/ije/dyy262>.
- Sanderson, E, W Spiller, and J Bowden. 2020. "Testing and Correcting for Weak and Pleiotropic Instruments in Two-Sample Multivariable Mendelian Randomisation." *bioRxiv*. <https://doi.org/10.1101/2020.04.02.021980>.