

What range could your causal effect lie between if the IV assumptions held?

Find out with our bpbounds R package and Shiny app!

# bpbounds: R package and web app

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## Introduction

- We present our bpbounds R package and Shiny web app for the nonparametric bounds for the average causal effect (ACE) due to Balke and Pearl (Palmer et al. 2018).
- This is an R implementation of our Stata programs (Palmer et al. 2011).
- The package can be installed from CRAN:

```
install.packages("bpbounds")
```

- Code development is on the GitHub repository: <https://github.com/remlapmot/bpbounds>

## Methods

- Balke and Pearl (1997) showed it is possible to derive bounds for the ACE.
- Bounds have the interpretation of:

There is some distribution of the unobserved confounders (of the exposure-outcome association) that yields a true ACE as small as the lower bound, while another choice produces an ACE as large as the upper bound.

- There are at least two ways to implement the Balke-Pearl bounds:

- i. conditional probabilities from contingency tables;
  - ii. the polytope method due to Dawid (2003).
- We implemented the polytope method since it is generalisable for identified IV models with exposures, outcomes, and instruments with more categories. Currently, we allow for a binary or 3 category instrument, and binary exposure and outcome.

## Example Mendelian randomization analysis

- We extract an example from Meleady et al. (2003).
- We have a 3 category instrument and binary phenotype and outcomes.
- We use the 677CT polymorphism (rs1801133) in the MTHFR gene, involved in folate metabolism, as an instrumental variable to investigate the causal effect of homocysteine on the risk of cardiovascular disease.
- The code is shown on the right.
- The ACE lies between a risk difference of -9% to 74% increase in absolute risk.
- Additionally, we see that the monotonicity inequality is not satisfied.

## Conclusion

- Use of bounds in instrumental variable analyses is regaining interest (Swanson et al. 2018; Labrecque and Swanson 2018).
- Our R package and app provide a convenient interface to the bounds.

## References

Balke, A., and J. Pearl. 1997. "Bounds on treatment effects from studies with imperfect compliance." *Journal of the American Statistical Association* 92 (439): 1172–6.

Dawid, A. P. 2003. "Causal Inference Using Influence Diagrams: The Problem of Partial Compliance (with Discussion)." In *Highly Structured Stochastic Systems*, edited by P. J. Green, N. L. Hjort, and S. Richardson, 45–65. New York: Oxford University Press.

Labrecque, Jeremy, and Sonja A Swanson. 2018. "Understanding the Assumptions Underlying Instrumental Variable Analyses: A Brief Review of Falsification Strategies and Related Tools." *Current Epidemiology Reports* 5 (3): 214–20.

Meleady, Raymond, Per M Ueland, Henk Blom, Alexander S Whitehead, Helga Refsum, Leslie E Daly, Stein Emil Vollset, et al. 2003. "Thermolabile Methyltetrahydrofolate Reductase, Homocysteine, and Cardiovascular Disease Risk: The European Concerted Action Project." *The American Journal of Clinical Nutrition* 77 (3): 43–50.

Palmer, T. M., R. Ramsahai, V. Didelez, and N. A. Sheehan. 2018. *bpbounds: R package implementing Balke-Pearl bounds for the average causal effect*. <https://CRAN.R-project.org/package=bpbounds>.

Palmer, T. M., R. Ramsahai, V. Didelez, and N. A. Sheehan. 2011. "Nonparametric Bounds for the Causal Effect in a Binary Instrumental-Variable Model." *Stata Journal* 11 (3): 345–67. <http://www.stata-journal.com/article.html?article=st0232>.

Swanson, Sonja A., Miguel A. Hernán, Matthew Miller, James M. Robins, and Thomas S. Richardson. 2018. "Partial Identification of the Average Treatment Effect Using Instrumental Variables: Review of Methods for Binary Instruments, Treatments, and Outcomes." *Journal of the American Statistical Association* 113 (523): 931–47. <https://doi.org/10.1080/01621459.2018.1434530>.

## Extra Figures & Tables

```
library(bpbounds)

mt3 <- c(.83, .05, .11, .01,
        .88, .06, .05, .01,
        .72, .05, .20, .03)

p3 <- array(mt3, dim = c(2, 2, 3),
           dimnames = list(x = c(0, 1),
                          y = c(0, 1),
                          z = c(0, 1, 2)))

bpres3 <- bpbounds(as.table(p3))
summary(bpres3)

##
## Data:                               trivariate
## Instrument categories:              3
##
## Instrumental inequality: TRUE
## Causal parameter Lower bound Upper bound
## ACE -0.09 0.74000
## P(Y|do(X=0)) 0.06 0.12000
## P(Y|do(X=1)) 0.03 0.80000
## CRR 0.25 13.33333
##
## Monotonicity inequality: FALSE
```



Figure 1: Shiny app <https://remlapmot.shinyapps.io/bpbounds>

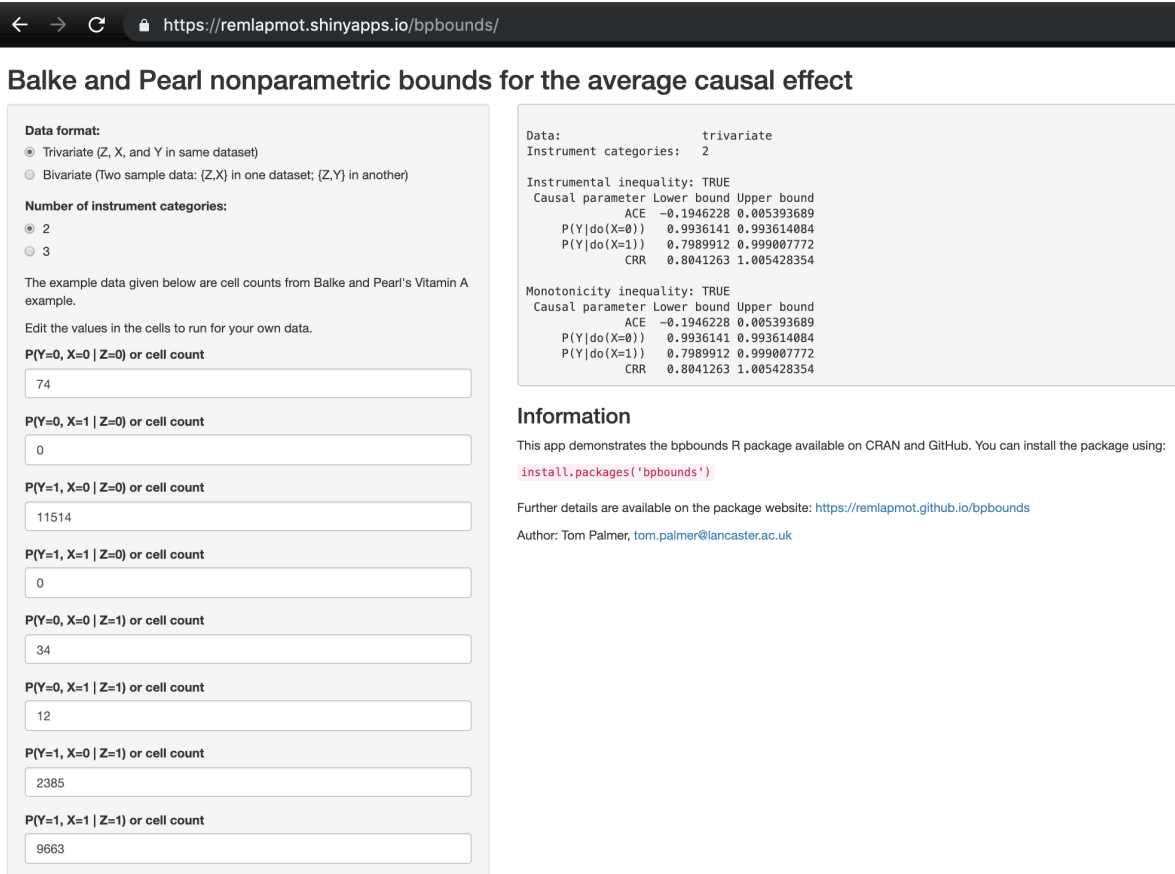


Figure 2: Screenshot of our Shiny app.



Figure 3: Package website <https://remlapmot.github.io/bpbounds/>

