

tipr: An R package for sensitivity analyses for unmeasured confounders

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DOI: 10.21105/joss.04495

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Submitted: 06 May 2022 Published: 05 September 2022

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Summary

The strength of evidence provided by epidemiological and observational studies is inherently limited by the potential for unmeasured confounding. We focus on three key quantities: the observed bound of the confidence interval closest to the null, the relationship between an unmeasured confounder and the outcome, for example a plausible residual effect size for an unmeasured continuous or binary confounder, and the relationship between an unmeasured confounder and the exposure, for example a realistic mean difference or prevalence difference for this hypothetical confounder between exposure groups. Building on the methods put forth by Cornfield et al. (1959), Bross (1966), Schlesselman (1978), Rosenbaum & Rubin (1983), Lin et al. (1998), Lash et al. (2009), Rosenbaum (1986), Cinelli & Hazlett (2020), VanderWeele & Ding (2017), and Ding & VanderWeele (2016), we can use these quantities to assess how an unmeasured confounder may tip our result to insignificance.

Statement of need

When assessing the relationship between an exposure and an outcome, the "no unmeasured confounders" assumption is crucial. (D'Agostino, 1998; Rubin, 1974) This assumption, however, is untestable, rendering sensitivity analyses necessary to quantify the potential impact of unmeasured confounding. There are several related methods for conducting sensitivity analyses for unmeasured confounders (Bross, 1966; Cinelli & Hazlett, 2020; Cornfield et al., 1959; Ding & VanderWeele, 2016; Lash et al., 2009; Lin et al., 1998; Rosenbaum, 1986; Rosenbaum & Rubin, 1983; Schlesselman, 1978; VanderWeele & Ding, 2017), some of which have their own R packages, for example methods in Cinelli & Hazlett (2020) can be implemented using the sensemakr R package, obsSens implements methods from Lin et al. (1998), and EValue implements methods in VanderWeele & Ding (2017). However, there is not currently a single R package that has a unified grammar allowing the user to conduct appropriate sensitivity analysis for their study. A unified grammar allows the user to easily transition between methods depending on the scenario under which they are conducting the sensitivity analysis as well as compare the results between methods. tipr seeks to fill this need.

Overview

The tipr R package (R Core Team, 2017) allows the user to conduct sensitivity analyses for unmeasured confounding. The user provides an "observed effect" (that is, some effect between an exposure of interest and an outcome of interest after adjusting for any observed confounders), and the sensitivity analyses will quantify how sensitive the effect is to a potential unmeasured confounder. The tipr functions fall into three categories:

(1) Functions that calculate how an observed effects would change with a specified unmeasured confounder (adjust functions)



- (2) Functions that calculate the magnitude of an unmeasured confounder needed to tip an observed effect to cross the null (tip functions)
- (3) Functions that calculate single number summaries of an observed effect's "sensitivity", such as the E-value (VanderWeele & Ding, 2017) or Robustness value (Cinelli & Hazlett, 2020).

tipr is available on CRAN and Github. Documentation can be found at https://lucymc-gowan.github.io/tipr/. It can be installed from CRAN using the following code:

install.packages("tipr")

The development version can be installed from Github using the following code:

devtools::install github("LucyMcGowan/tipr")

The library can be loaded by running:

library(tipr)

Unmeasured confounder parameterization

Informally, a confounder is a pre-exposure variable that is associated with both the exposure of interest and outcome of interest but is *not* on the causal pathway between the two.(Van Stralen et al., 2010; VanderWeele & Shpitser, 2013). There are two quantities that measure the strength of an unmeasured confounder – the relationship between the unmeasured confounder and the exposure and the strength of the unmeasured confounder and the outcome. The tipr package allows these quantities to be specified in a number of ways.

Unmeasured confounder - exposure relationship

If quantifying the impact of a standardized Normally distributed confounder, the impact of the unmeasured confounder on the exposure is parameterized as a difference in means between the unmeasured confounder in the exposed population and the unexposed population. By "standardized Normally distributed" we mean that the unmeasured confounder is Normally distributed with mean μ_1 for the exposed and μ_0 for the unexposed and unit variance. (Note a standardized Normally distributed confounder can be created from a Normally distributed confounder by dividing by the standard deviation). This is specified using the exposure_confounder_effect parameter. If quantifying the impact of a binary confounder, the impact is quantified using two parameters, exposed_confounder_prev: The estimated prevalence of the unmeasured confounder in the exposed population. Alternatively, the user can choose to quantify the magnitude of the unmeasured confounder-exposure relationship in terms of the percent of variation in the exposure explained by the unmeasured confounder. This is specified using the confounder_exposure_r2 parameter.

Unmeasured confounder - outcome relationship

If quantifying the impact of a standardized Normally distributed confounder or binary confounder, the impact of the unmeasured confounder on the outcome is parameterized as the anticipated effect size of the unmeasured confounder if it were included in the final outcome model. This effect will be on the same scale as the input observed effect. For example, if the observed effect is a *coefficient* in a linear regression model, this effect would also be a coefficient; if the observed effect is a *hazard ratio* from a Cox proportional hazards model, this effect would also be a hazard ratio. This is specified using the confounder_outcome_effect parameter. Alternatively, the user can choose to quantify the magnitude of the unmeasured confounder-outcome relationship in terms of the percent of variation in the outcome explained by the unmeasured confounder. This is specified using the confounder_outcome_r2 parameter.



Syntax

The functions in the tipr package follow a unified grammar. The function names follow this form: {action}_{effect}_with_{what}. For example, to adjust (action) a coefficient (effect) with a binary unmeasured confounder (what), we use the function adjust_coef_with_binary(). The full list of actions effects and whats is summarized in Table 1. The "default" function assumes the confounder type is Normally distributed, therefore adjust_coef() is equivalent to adjust_coef_with_continuous().

Table 1. Grammar of tipr functions.

category	Function term	Use
action	adjust	These functions adjust observed effects, requiring both the unmeasured confounder-exposure relationship and unmeasured confounder-outcome relationship to be specified.
	tip	These functions tip observed effects. Only one relationship, either the unmeasured confounder-exposure relationship or unmeasured confounder-outcome relationship needs to be specified.
effect	coef	These functions specify an observed coefficient from a linear, log-linear, logistic, or Cox proportional hazards model
	rr	These functions specify an observed relative risk
	or	These functions specify an observed odds ratio
	hr	These functions specify an observed hazard ratio
what	continuous	These functions specify an unmeasured standardized Normally distributed confounder. These functions will include the parameters exposure_confounder_effect and confounder_outcome_effect
	binary	These functions specify an unmeasured binary confounder. These functions will include the parameters exposed_confounder_prev, unexposed_confounder_prev, and confounder_outcome_effect
	r2	These functions specify an unmeasured confounder parameterized by specifying the percent of variation in the exposure / outcom explained by the unmeasured confounder. These functions will include the parameters confounder_exposure_r2 and outcome_exposure_r2

The first argument of each function is effect_observed quantifying the observed exposure-outcome relationship. The functions intended to adjust an effect with partial R^2 values specified are built on the sensemakr package (Cinelli et al., 2021) and have the following additional arguments: se: The standard error fo the observed exposure - outcome relationship and df: The residual degrees of freedom from the model used to fit the observed exposure - outcome relationship. This is the total number of observations minus the number of parameters estimated in your model. Often for models estimated with an intercept this is N-k-1 where k is the number of predictors in the model. The subsequent parameters in each function describe the unmeasured confounder's relationship with the exposure and outcome for the adjust functions, and one or the other for the tip functions.



The output for all functions is a data frame with the adjusted effect in the first column, the observed effect in the second column, and the specified sensitivity parameters in the subsequent columns. For example, if we want to know the impact of an unmeasured confounder with exposure_confounder_effect = 0.1 and confounder_outcome_effect = 1 on an observed effect = 1.5, we would run the following code.

```
library(tipr)
adjust_coef(effect_observed = 1.5,
            exposure_confounder_effect = 0.1,
            confounder outcome effect = 1)
## The observed effect (1.5) is updated to 1.4 by a confounder with the
## following specifications:
   * estimated difference in means: 0.1
##
   * estimated relationship between the unmeasured confounder and the
##
      outcome: 1
## # A tibble: 1 × 4
   effect_adjusted effect_observed exposure_confounder_effect confounder_outcom...
##
              <dbl>
                               <dbl>
                                                           <dbl>
                                                                              <dbl>
               1.4
                                1.5
                                                           0.1
##
                                                                                1
```

Additionally, there are two function, e_value, and r_value that calculate single number summaries of the sensitivity of a particular observed effect to unmeasured confounding, the E-value and Robustness value, respectively.

Tipping Point Example

The package includes a few example data sets. exdata_rr contains data simulated such that there is a binary exposure, a binary outcome, and two Normally distributed confounders, one "measured" (so included in the analysis) and one "unmeasured". There is no true causal effect between the exposure and outcome. The true causal diagram is shown below.

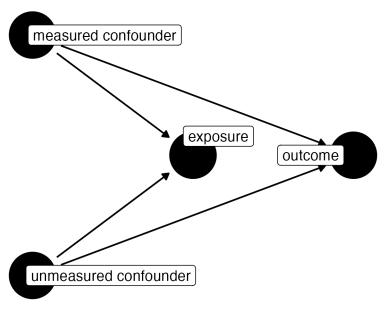


Figure 1: Causal Diagram illustrating two confounders, one measured and one unmeasured

We can fit the observable model as follows:



```
glm(outcome ~ exposure + measured_confounder,
    data = exdata_rr,
    family = poisson)
```

After fitting your model, you can determine the unmeasured confounder needed to tip your analysis. In this example, the exposure-outcome relationship is a relative risk of 1.5 (95% CI: 1.1, 2.1).

We are interested in a Normally distributed unmeasured confounder, so we can use the tip_rr_with_continuous() function. The function tip() is an alias for this function.

Let's assume the relationship between the unmeasured confounder and exposure is 0.5 (exposure_confounder_effect = 0.5, meaning the difference in means between the exposure groups is 0.5), let's solve for the relationship between unmeasured confounder and outcome needed to tip the analysis (in this case, we are solving for confounder_outcome_effect).

```
tip(1.1, exposure_confounder_effect = 0.5)
## The observed effect (1.1) WOULD be tipped by 1 unmeasured confounder
## with the following specifications:
##
    * estimated difference in means between the unmeasured
       confounder in the exposed population and unexposed population: 0.5
##
     * estimated association between the unmeasured confounder and
##
##
       the outcome: 1.21
##
##
## # A tibble: 1 × 5
     effect_adjusted effect_observed exposure_confound... confounder_outcome...
##
##
                                <dbl>
                                                   <dbl>
               <dbl>
                                                                         <dbl>
## 1
                                  1.1
                                                      0.5
                                                                          1.21
## # ... with 1 more variable: n unmeasured confounders <dbl>
```

A hypothetical unobserved continuous confounder that has a mean difference of 0.5 between the exposure groups would need a relationship with the outcome of 1.21 to tip this analysis at the 5% level.

Conclusion

The tipr package facilitates sensitivity analyses for unmeasured confounding, building on the methods put forth by Cornfield et al. (1959), Bross (1966), Schlesselman (1978), Rosenbaum & Rubin (1983), Lin et al. (1998), Lash et al. (2009), Rosenbaum (1986), Cinelli & Hazlett (2020), VanderWeele & Ding (2017), and Ding & VanderWeele (2016). The adjust functions allow an investigator to examine how a specific (or set of specific) confounders would change a result while the tip functions provide sensitivity analyses that allow an investigator to examine how extreme an unmeasured confounder would need to be in order to change the direction of the effect, and thus often the conclusions of the study.

References

Bross, I. (1966). Spurious effects from an extraneous variable. *Journal of Chronic Diseases*. https://doi.org/10.1016/0021-9681(66)90062-2

Cinelli, C., Ferwerda, J., & Hazlett, C. (2021). Sensemakr: Sensitivity analysis tools for regression models. https://CRAN.R-project.org/package=sensemakr

Cinelli, C., & Hazlett, C. (2020). Making sense of sensitivity: Extending omitted variable bias. Journal of the Royal Statistical Society: Series B (Statistical Methodology), 82(1), 39–67.



https://doi.org/10.1111/rssb.12348

- Cornfield, J., Haenszel, W., Hammond, E., Lilienfeld, A., Shimkin, M., & Wynder, E. (1959). Smoking and lung cancer: recent evidence and a discussion of some questions. *Journal of the National Cancer Institute*, 22(1), 173–203. https://doi.org/10.1093/jnci/22.1.173
- D'Agostino, R. (1998). Propensity score methods for bias reduction in the comparison of a treatment to a non-randomized control group. *Statistics in Medicine*, *17*(19), 2265–2281. https://doi.org/10.1002/(SICI)1097-0258(19981015)17:19%3C2265::AID-SIM918%3E3.0.CO;2-B
- Ding, P., & VanderWeele, T. J. (2016). Sensitivity Analysis Without Assumptions. *Epidemiology (Cambridge, Mass.)*, 27(3), 368–377. https://doi.org/10.1097/EDE. 0000000000000457
- Lash, T. L., Fox, M. P., Fink, A. K., & others. (2009). Applying quantitative bias analysis to epidemiologic data (Vol. 533). Springer. https://doi.org/10.1007/978-0-387-87959-8
- Lin, D., Psaty, B., & Kronmal, R. (1998). Assessing the sensitivity of regression results to unmeasured confounders in observational studies. *Biometrics*, *54*(3), 948–963. https://doi.org/10.2307/2533848
- R Core Team. (2017). R: A language and environment for statistical computing. R Foundation for Statistical Computing. https://www.R-project.org/
- Rosenbaum, P. (1986). Dropping out of high school in the united states: An observational study. *Journal of Educational Statistics*, 11(3), 207–224. https://doi.org/10.3102/10769986011003207
- Rosenbaum, P., & Rubin, D. (1983). Assessing sensitivity to an unobserved binary covariate in an observational study with binary outcome. *Journal of the Royal Statistical Society Series B.* https://doi.org/10.1111/j.2517-6161.1983.tb01242.x
- Rubin, D. (1974). Estimating causal effects of treatments in randomized and nonrandomized studies. *Journal of Educational Psychology*, 66(5), 688. https://doi.org/10.1037/h0037350
- Schlesselman, J. (1978). Assessing effects of confounding variables. *American Journal of Epidemiology*, 108(1), 3–8. https://doi.org/10.1093/oxfordjournals.aje.a112581
- Van Stralen, K., Dekker, F., Zoccali, C., & Jager, K. (2010). Confounding. *Nephron Clinical Practice*, 116(2), c143–c147. https://doi.org/10.1159/000315883
- VanderWeele, T. J., & Ding, P. (2017). Sensitivity Analysis in Observational Research: Introducing the E-Value. *Ann Intern Med.* https://doi.org/10.7326/M16-2607
- VanderWeele, T. J., & Shpitser, I. (2013). On the definition of a confounder. *Annals of Statistics*, 41(1), 196. https://doi.org/10.1214/12-aos1058