

Meta-analysis

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Overview

- What is meta-analysis?
- When can/should I use meta-analysis?
- What are the steps in meta-analysis?
- Meta-analysis in R
- Pitfalls and limitations of meta-analysis

Suggested reading:

Viechtbauer, W. (2010). Conducting meta-analyses in R with the metafor package. *Journal of Statistical Software*, 36(3), 1–48.
Available at: <https://www.jstatsoft.org/article/view/v036i03>

What is meta-analysis?

- A meta-analysis = systematic review with quantitative analysis of the extracted data:
 - *Single summary of the effect*
 - *Heterogeneity of effects*
 - *How effects differ depending on features of primary studies*
 - *Whether studies with more favourable findings are more likely to be published ('publication bias')*

When can/should I use meta-analysis?

- Best for well-defined and relatively narrow research questions e.g., a group difference, intervention effect, or association between two variables
- Primary studies must be quantitative
- Technically requires only two studies but greatest value when there are enough studies to conduct moderator analysis

Steps in meta-analysis

- Specify research question, search, and eligibility criteria
- Systematic search
- Assess against inclusion/exclusion criteria
- Quality assessment and **data extraction**
- **Data analysis**
 - *Calculate/convert effect sizes*
 - *Meta-analytic model(s)*
 - Pooled effect size estimate
 - Heterogeneity
 - Meta-regression
 - *Assess publication bias*

Data extraction

- Need to extract **effect size** measure and associated sampling variance
- Effect sizes = standardised metric to facilitate comparisons across studies
- Appropriate effect size measure depends on study designs
- Examples:
 - **Odds ratio, relative risk ratio, risk difference** for 2x2 data (e.g., two groups with binary outcome)
 - **Standardised mean difference** for comparing two groups on continuous outcome (e.g., sex differences in neuroticism)
 - Raw and transformed **correlation coefficient** for assessing strength of association
 - **Proportion** (e.g., for prevalence studies)

Data extraction

- Also extract/derive other study information for descriptive purposes and/or moderator analysis
 - *Study year*
 - *Quality*
 - *Lab of origin*
 - *Type of intervention*
 - *Study design features*

Fitting the meta-analysis model

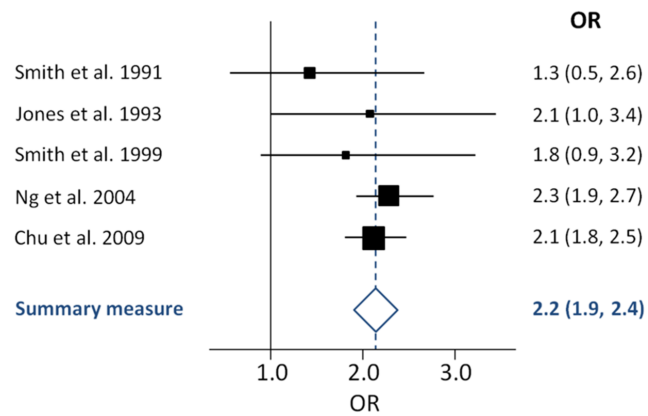
- Calculated using either a fixed-effect or random-effects model
 - *Fixed-effect model assumes all studies estimate common effect, with variation due only to sampling variation*
 - *Random-effects model assumes that the sampled studies come from a broader population of studies varying in true effect*
 - Provides an estimate of variation in true effect τ^2
 - *Choice between fixed or random should be made a priori, not based on estimated heterogeneity*
- Weighted estimation can be used to up-weight studies with greater precision (lower SEs/bigger sample size)
- Effect sizes can be regressed on predictors to estimate moderator effects

Results from meta-analytic models

- **Pooled effect size estimate** provides estimate of true effect (or average true effect for random-effects meta-analysis)
- I^2 = % of variation between studies
 - *Bigger I^2 means greater heterogeneity*
- β **coefficients** and associated p-values estimate effect and significance of moderators

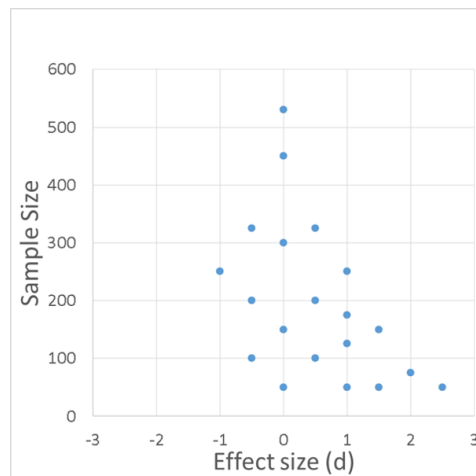
Plotting meta-analytic results

- Forest plots used to display meta-analytic results



Publication bias

- Funnel plots can be used to help identify publication bias
- x-axis is effect size, y-axis is N, SE, or sampling variance



Statistical methods to assess publication bias

- Statistical tests of the relation between effect size and precision also possible
 - *Egger test (regression-based method)*
 - *Rank correlation test*
- Trim and fill method
 - *Estimates number of unpublished studies needed to bring pooled effect size estimate to non-significant*
 - *Con: not possible for meta-analytic models with moderators*

Meta-analysis in R

- Several software options but best option in R is metafor package
- Functions for:
 - *calculating effect sizes*
 - *Fitting meta-analytic models (including moderators)*
 - *Plotting results*
 - *Statistically assessing publication bias*

Example: the effectiveness of BCG vaccination against tuberculosis

```
library(metafor)
data("dat.bcg", package="metafor")
print(dat.bcg)
```

##	trial	author	year	tpos	tneg	cpos	cneg	ablat	alloc
## 1	1	Aronson	1948	4	119	11	128	44	random
## 2	2	Ferguson & Simes	1949	6	300	29	274	55	random
## 3	3	Rosenthal et al	1960	3	228	11	209	42	random
## 4	4	Hart & Sutherland	1977	62	13536	248	12619	52	random
## 5	5	Frimodt-Moller et al	1973	33	5036	47	5761	13	alternate
## 6	6	Stein & Aronson	1953	180	1361	372	1079	44	alternate
## 7	7	Vandiviere et al	1973	8	2537	10	619	19	random
## 8	8	TPT Madras	1980	505	87886	499	87892	13	random
## 9	9	Coetzee & Berjak	1968	29	7470	45	7232	27	random
## 10	10	Rosenthal et al	1961	17	1699	65	1600	42	systematic
## 11	11	Comstock et al	1974	186	50448	141	27197	18	systematic
## 12	12	Comstock & Webster	1969	5	2493	3	2338	33	systematic
## 13	13	Comstock et al	1976	27	16886	29	17825	33	systematic

- 13 primary studies
- In each study two groups (treated vs control) with a binary outcome (tested positive vs negative for TB)
 - *We can use OR, RR, RD as our effect size measure*
- Additional information about study design and latitude of study

Calculate effect sizes

- We can calculate the effect sizes and associated sampling variation using `escalc()` function
- We supply the Ns in each of the four cells of the 2-by-2 table:

TB+ TB-

Treated ai bi

Control ci di

```
dat.bcg<-escalc(measure="OR", ai=tpos, bi=tneg, ci=cpos, di=cneg, data=dat.bcg, append=T)
head(dat.bcg)
```

```
## trial author year tpos tneg cpos cneg ablat alloc yi
## 1 1 Aronson 1948 4 119 11 128 44 random -0.9387
## 2 2 Ferguson & Simes 1949 6 300 29 274 55 random -1.6662
## 3 3 Rosenthal et al 1960 3 228 11 209 42 random -1.3863
## 4 4 Hart & Sutherland 1977 62 13536 248 12619 52 random -1.4564
## 5 5 Frimodt-Moller et al 1973 33 5036 47 5761 13 alternate -0.2191
## 6 6 Stein & Aronson 1953 180 1361 372 1079 44 alternate -0.9581
## vi
## 1 0.3571
## 2 0.2081
## 3 0.4334
## 4 0.0203
## 5 0.0520
## 6 0.0099
```

Fit random-effects model

- We can fit a random-effects model using the `rma()` function, supplying the newly calculated ORs and sampling variance:

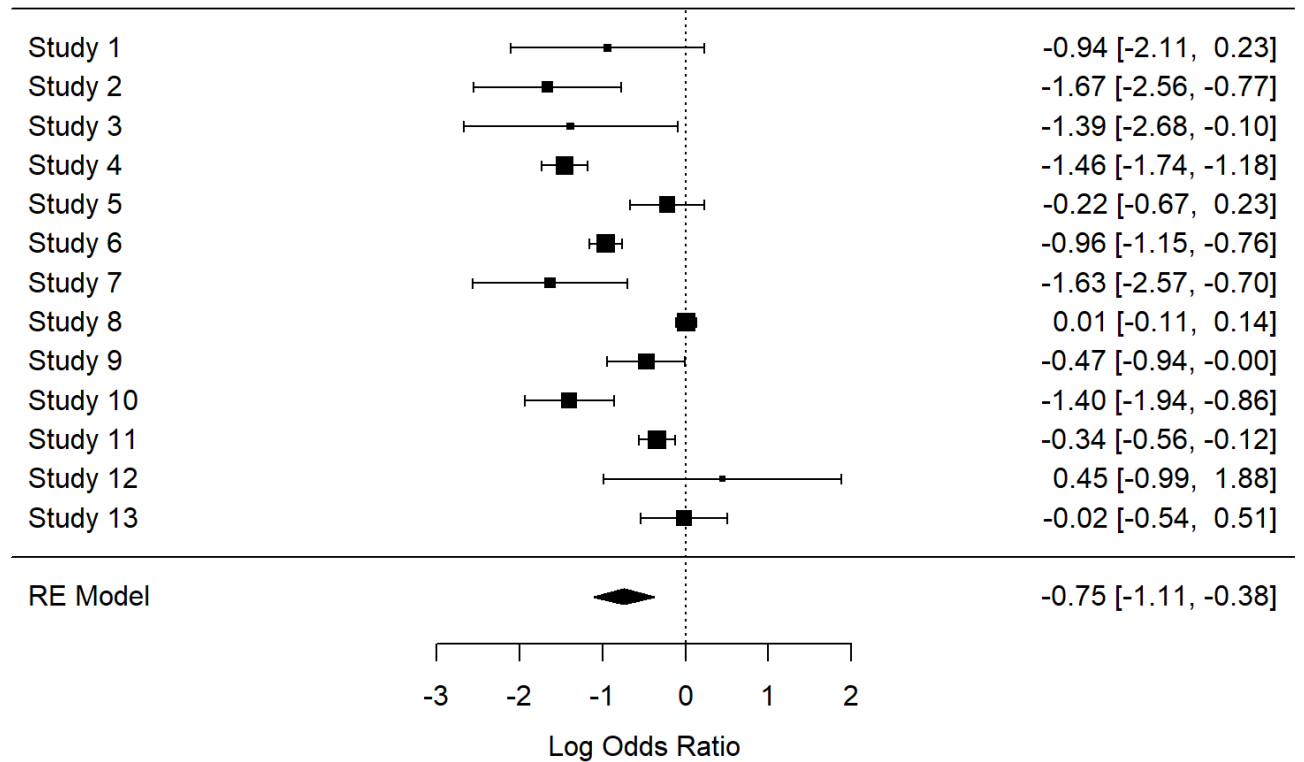
```
rEffs<-rma(yi=yi, vi=vi, data=dat.bcg) #yi is the effect size measures, vi is their sampling variance
rEffs #NB will use log OR
```

```
##
## Random-Effects Model (k = 13; tau^2 estimator: REML)
##
## tau^2 (estimated amount of total heterogeneity): 0.3378 (SE = 0.1784)
## tau (square root of estimated tau^2 value):      0.5812
## I^2 (total heterogeneity / total variability):    92.07%
## H^2 (total variability / sampling variability):   12.61
##
## Test for Heterogeneity:
## Q(df = 12) = 163.1649, p-val < .0001
##
## Model Results:
##
## estimate      se      zval      pval      ci.lb      ci.ub
## -0.7452  0.1860  -4.0057  <.0001  -1.1098  -0.3806  ***
##
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

- Significant pooled effect [exponentiate coefficients using `exp()` to convert to ORs]
- Substantial heterogeneity

Visualise using forest plot

```
forest(rEffs)
```



Include moderators in model

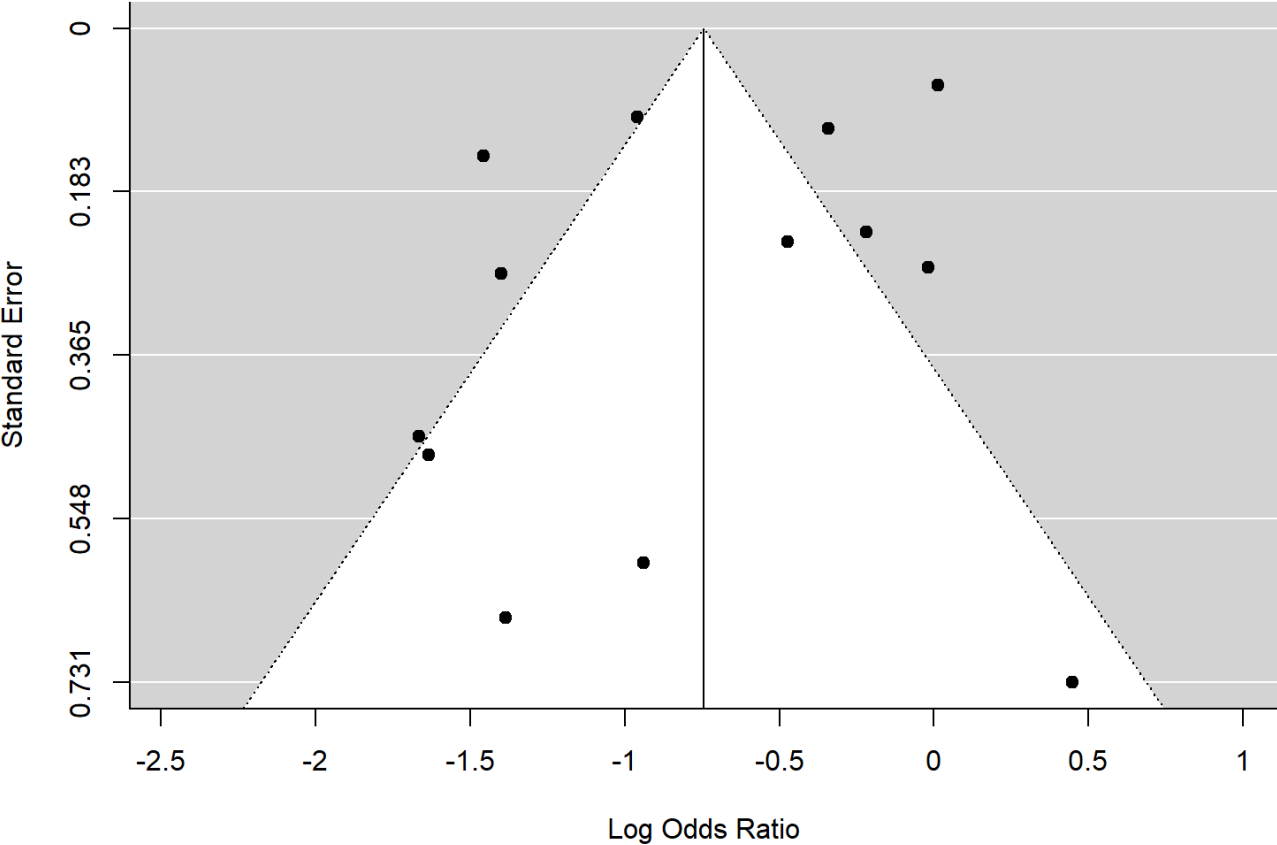
```
mEffs<- rma(yi=yi, vi=vi, mods= ~ablat+year, data=dat.bcg) #ablat and year are moderators; specified lm()-style
mEffs
```

```
##
## Mixed-Effects Model (k = 13; tau^2 estimator: REML)
##
## tau^2 (estimated amount of residual heterogeneity):      0.0913 (SE = 0.0745)
## tau (square root of estimated tau^2 value):            0.3022
## I^2 (residual heterogeneity / unaccounted variability): 67.29%
## H^2 (unaccounted variability / sampling variability):    3.06
## R^2 (amount of heterogeneity accounted for):            72.96%
##
## Test for Residual Heterogeneity:
## QE(df = 10) = 25.0121, p-val = 0.0053
##
## Test of Moderators (coefficients 2:3):
## QM(df = 2) = 16.2533, p-val = 0.0003
##
## Model Results:
##
##      estimate      se      zval      pval      ci.lb      ci.ub
## intrcpt -10.5347  27.3739  -0.3848   0.7004  -64.1865   43.1172
## ablat    -0.0288   0.0095  -3.0311   0.0024   -0.0475   -0.0102 **
## year      0.0055   0.0138   0.3949   0.6929   -0.0216    0.0325
##
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

- Sig effect of latitude
- No sig effect of study year

Examine publication bias

```
funnel(rEffs)
```



Statistical evaluation of publication bias

```
regtest(rEfs, model='lm') #regress effect sizes on standard error
```

```
##  
## Regression Test for Funnel Plot Asymmetry  
##  
## model:      weighted regression with multiplicative dispersion  
## predictor: standard error  
##  
## test for funnel plot asymmetry: t = -1.5070, df = 11, p = 0.1600
```

- regtest() tells us no significant relation between standard error and effect size

Trim and fill method

```
trimfill(rEfts) #trim & fill method
```

```
##
## Estimated number of missing studies on the right side: 0 (SE = 2.3309)
##
## Random-Effects Model (k = 13; tau^2 estimator: REML)
##
## tau^2 (estimated amount of total heterogeneity): 0.3378 (SE = 0.1784)
## tau (square root of estimated tau^2 value):      0.5812
## I^2 (total heterogeneity / total variability):    92.07%
## H^2 (total variability / sampling variability):   12.61
##
## Test for Heterogeneity:
## Q(df = 12) = 163.1649, p-val < .0001
##
## Model Results:
##
## estimate      se      zval      pval      ci.lb      ci.ub
## -0.7452  0.1860  -4.0057  <.0001  -1.1098  -0.3806  ***
##
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

- Trim and fill method suggests that accounting for publication bias would not make result non-significant

Pitfalls and limitations

- Garbage in, garbage out
- Typically able to include fewer studies than a systematic review
- Comparing apples and oranges
- File-drawer problem
- Subjectivity
- Structured/mechanical approach does not lend itself well to all research questions

Summary

- Meta-analysis provides quantitative summary across multiple studies
- Can give greater weight to more precise studies
- Can answer questions that individual studies can't
 - *Presence of publication bias*
 - *Sources of heterogeneity*
- Doesn't solve problem of poor primary study quality/publication bias etc.