

Cochran's Q

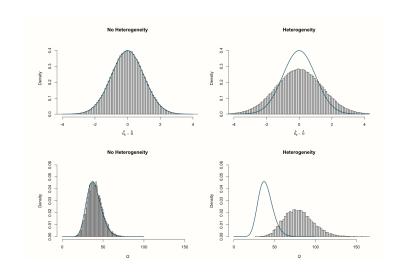
- Traditionally, Cochran's *Q* (Cochran, 1954) has been used to distinguish studies' sampling error from actual between-study heterogeneity.
- Q uses the deviation of each study's observed effect $\hat{\theta}_k$ from the summary effect $\hat{\theta}$ weighted by the inverse of the study's variance, w_k :

$$Q = \sum_{k=1}^{K} w_k (\hat{\theta}_k - \hat{\theta})^2$$

• We assume that, if variability in observed effect sizes is only caused by sampling error (i.e., the EE-model holds), then Q will follow a χ^2 distribution with K-1 degrees of freedom, where K is the number of studies.

$$Q \sim \chi_{K-1}^2$$

→ If *Q* exceeds this expected value, we are more and more likely to conclude "significant" excess variability (=heterogeneity).

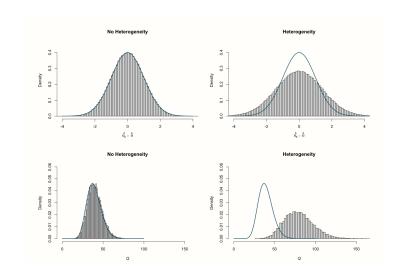




Cochran's Q

Problems With Q & the Q-Test

- Q increases both when the number of studies K, and when the precision (i.e. the sample size of a study) increases.
- Therefore, *Q* and whether it is significant highly depends on the size of the meta-analysis, and thus its statistical power.
- → We should not only rely on the significance of a Q-test when assessing heterogeneity.
- → Sometimes, meta-analysts decide whether to apply a fixed-effect or random-effects model based on the significance of the *Q*-test. This is also highly discouraged



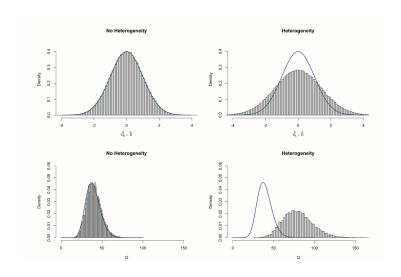


Higgins and Thompson's I^2

- I² is another metric to quantify the between-study heterogeneity in meta-analyses
- In its "classic" definition, I^2 quantifies, in percent, how much the observed value of Q exceeds the expected Q value when there is no heterogeneity (i.e., K-1):

$$I^2 = \max\left\{0, \frac{Q - (K - 1)}{Q}\right\}$$

- The popularity of this statistic may be associated with the fact that there is an (in)famous "rule of thumb" how to interpret it (Higgins & Thompson 2002):
 - $I^2 = 25\%$: low heterogeneity
 - $I^2 = 50\%$: moderate heterogeneity
 - $I^2 = 75\%$: substantial heterogeneity.

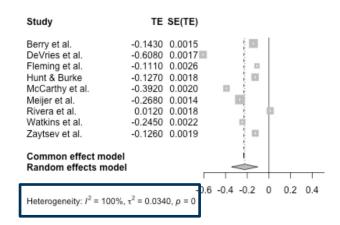




Higgins and Thompson's I^2

Problems of I^2

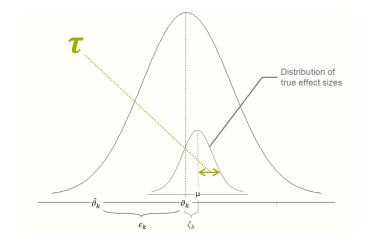
- I^2 is <u>not</u> sensitive to changes in the **number of studies** in the analysis.
- However, it is still a relative measure of heterogeneity, and its value heavily depends on the precision of the included studies
- I^2 is the percentage of variability not caused by sampling error.
- If our studies become increasingly large, the sampling error tends to zero, while at the same time, I² tends to 100% – simply because the studies have a greater sample size.



$$I^2 = \max\left\{0, \frac{Q - (K - 1)}{Q}\right\}$$



- Only τ^2 is insensitive to <u>both</u> the number and precision of the included studies
- τ (square root of τ^2) can be interpreted as the **standard** deviation of the true effect size distribution
- But this is often difficult to communicate to stakeholders without training in meta-analysis

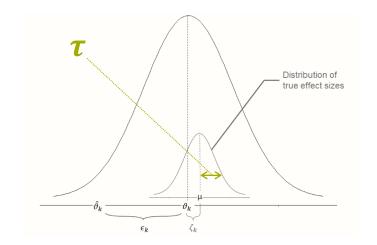




- Solution: calculate prediction intervals (PI), which directly express the impact that τ^2 has on the scale of the effect size
- Indicates the interval into which effect sizes of new studies are expected to fall, based on present evidence:

Upper PI:
$$\hat{\mu} + t_{K-1,0.975} \sqrt{\widehat{SE}_{\mu}^2 + \hat{\tau}^2}$$

Lower PI:
$$\hat{\mu} - t_{K-1,0.975} \sqrt{\widehat{SE}_{\mu}^2 + \hat{\tau}^2}$$

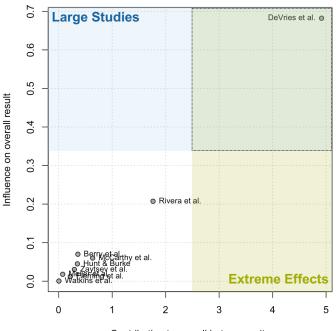


- PIs can be much wider than the CI.
- As $\tau^2 \to 0$, the PI and CI become increasingly identical.

Outliers & Model Diagnostics



- It is possible that very large studies, or studies with very extreme effect sizes may dominate the overall effect
- It is advisable to run sensitivity analyses excluding such studies
- There are no iron-clad rules on how to define outliers in a meta-analysis (but Viechtbauer & Cheung, 2010 provide helpful diagnostics)
- An intuitive way to identify outliers is through a
 Baujat plot



Contribution to overall heterogeneity