

Georg Heinze

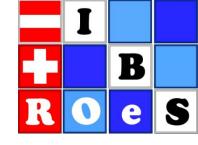
Section for Clinical Biometrics

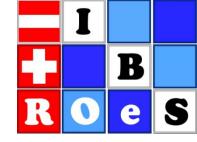
Center for Medical Statistics, Informatics and Intelligent Systems (CeMSIIS) Medical University of Vienna

- The data flood:
- Many simultaneous tests
- New hypotheses beyond $\mu 1=\mu 2$
- Prediction with >>n explanatory variables

- A variety of new platforms: genomics, methylation, copy numbers, transcriptomics, NGSeq, proteomics, metabolomics, miRNA
- Impossibility to check assumptions

- The levees:
- FDR, cost-effective sequential designs
- New test statistics (moderated t, LIM, adaptive trimmed t, ...)
- Regularized estimation,
- cross-validated tuning,
- double-cross-validated internal validation
- (do not optimize and validate with the same data!)
- Adopt methods to platforms,
- Keep analysis pipelines traceable
- Robust methods that do not make strong assumptions (c')

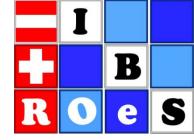




• Lists of differential expressed genes:

Please don't take them for granted;
use concepts like
locFDR or
Odds of differential expression
to weight genes in further gene enrichment analysis

Statistical models: serve as a (simplified) description of reality.
 They should help us to deduce general principles in biological associations.
 Incorporate existing knowledge (eg, confounders, pathways) where possible,
 Don't let machines do this artistic task alone!



- Why care about omics?
- They may provide useful answers to understand:

Why patients are different,

Why some patients have better prognosis than others,

The etiology of diseases, and How to improve prognosis,

How to find new (perhaps personalized) therapy targets.