

Minimal Data Needed for Valid & Accurate Image-Based fMRI Meta-Analysis

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Abstract

Meta-analysis provides a quantitative approach to summarise the rich functional Magnetic Resonance Imaging literature (fMRI). Due to the lack of availability of image data supporting existing literature, the majority of fMRI meta-analysis are coordinate-based. However, when image data is available for each study, the optimal approach is to perform an image-based meta-analysis. A number of approaches have been proposed to perform such meta-analyses including combination of standardised statistics, just effect estimates or both effects estimates and their sampling variance. While the latter is the preferred approach in the statistical community, its properties are only guaranteed in large samples. Additionally, often only standardised estimates are shared, reducing the possible meta-analytic approaches. Finally, because the BOLD signal is non-quantitative care has to be taken in order to insure that effect estimates are expressed in the same units, especially when combining data from different software packages. Given the growing interest in data sharing in the neuroimaging community there is a need to identify what is the minimal data to be shared in order to allow for future image-based meta-analysis. In this paper, we compare the validity and the accuracy of nine meta-analytic approaches on simulated and real data.

Keywords: Meta-analysis, Neuroimaging, Mixed-effects

1. Introduction

A growing literature is focusing on the lack of statistical power in neuroimaging studies (see, e.g. [2]), feeding the debate on the validity and reproducibility of published neuroimaging results. Meta-analysis, by providing inference based on the results of previously conducted studies, provides an essential method to increase power and hence confidence in neuroimaging.

A number of methods have been proposed for neuroimaging meta-analysis (see [21] for a review). As the results of neuroimaging studies are usually conveyed by providing a table of peak coordinate and statistics, most of these meta-analyses are restricted to combining coordinate-based information. Nevertheless the best practice method is an Image-Based Meta-Analysis (IBMA) that combines the effect estimates and standard errors from each study [1].

In order for IBMA to be possible in neuroimaging, tools for sharing 3D volumes obtained as a result of a statistical analysis are needed. NeuroVault [7] is an example of one such platform which facilitates sharing of neuroimaging results data but emphasis is mainly on statistical maps. There are three evident approaches to sharing summary data from each study i :

1. the contrast estimates $\hat{\beta}_i$ and contrast variance estimates s_i^2 .
2. the contrast estimates $\hat{\beta}_i$.
3. the standardized statistic maps Z_i .

Depending on how much data is shared, different strategies can be used to combine the available results into a meta-analysis. While the first option is the best practice, leading to statistically optimal estimates [4], it requires the contrasts to be expressed with in the same units and inference relies on asymptotic results (i.e under large sample sizes). In fMRI, units will depends on the field strength [3] as well as data, model and contrast vector scaling [14] and the number of samples included in a meta-analysis is usually small.

Given the growing interest in data sharing in the neuroimaging community [19, 15], and the relative easiness of sharing and combining just (unitless) statistic maps, there is a need to identify what is the minimal data to be shared in order to allow for future IBMA.

Here we compare the use of IMBA using 9 meta-analytic approaches: 2 approaches use $\hat{\beta}_i$'s and s_i^2 's, 2 $\hat{\beta}_i$'s only and 5 Z_i 's. We compare the validity and the accuracy of the nine meta-analytic approaches on simulated and real data including 21 studies of pain.

Figure 1: False positive rates of the meta-analytic estimators under the null hypothesis for $p < 0.05$.

Section 2 provides theoretical background on each of the nine meta-analytic approaches. Experiments undertaken on simulated and real data are then described. The results are described in section 3. Discussions, including our recommendations are provide in 4 Finally, we conclude in section 5.

2. Methods

2.1. Theory

For study $i = 1, \dots, k$ we have contrast estimate $\hat{\beta}_i$, its contrast variance estimate s_i^2 (i.e. squared standard error), its equivalent Z-statistic map Z_i and its sample size n_i .

Combining contrast estimates and their standard error. The gold standard approach is to fit contrast estimates and their standard error with a hierarchical general linear model (GLM) [4], creating a third-level (level 1: subject; level 2: study; level 3: meta-analysis). The general formulation for the study-level data is:

$$\hat{\beta} = X\gamma + \epsilon \quad (1)$$

where $\gamma = [\gamma_1 \dots \gamma_p]^T$ is a vector of meta-analytic parameters to estimate, $\hat{\beta} = [\hat{\beta}_1 \dots \hat{\beta}_k]^T$ is the vector of contrast estimates, X is the $k \times p$ study-level matrix (typically $p = 1$ with just a column of ones for a one-sample test) and $\epsilon \sim \mathcal{N}(0, W)$ is the residual error term.

In the most general case of a random-effects meta-analysis, i.e. assuming non-zero between-study variance, we have $W = \text{diag}(\sigma_1^2/n_1 + \tau^2, \dots, \sigma_k^2/n_k + \tau^2)$ where τ^2 denotes the between-study variance and σ_i^2/n_i denotes the variance of the contrast estimate for study i . Eq. (1) can be solved by weighted least squares giving:

$$\hat{\gamma} = (X^T W^{-1} X)^{-1} X^T W^{-1} \hat{\beta} \quad (2)$$

$$\text{Var}(\hat{\gamma}) = (X^T W^{-1} X)^{-1} \quad (3)$$

Table S1 and S2 provides derivation of the WLS estimates and their sampling variances for one and two-sample tests. But in practice, the weight matrix W is unknown and has to be estimated from the data. Given \hat{W} a consistent estimate of W , the feasible generalized least squares (FGLS) estimator is computed as:

$$\hat{\gamma} = (X^T \hat{W}^{-1} X)^{-1} X^T \hat{W}^{-1} \hat{\beta} \quad (4)$$

$$\text{Var}(\hat{\gamma}) = (X^T \hat{W}^{-1} X)^{-1} \quad (5)$$

Here we use s_i^2 as an estimate of σ_i^2/n_i and estimate $\hat{\tau}^2$ from the data. FGLS is asymptotically efficient but its finite sample properties are unknown [8]. We used the ‘FLAME 1’ implementation available in FSL that provides inference by comparing the statistic to a Student distribution with $k - p$ degrees of freedom [26] as depicted in Table 1. This reference approach will be referred to as **Mixed-effects (MFX) GLM**.

In a fixed-effects meta-analysis, i.e. assuming no or negligible between-study variance, we have $W = \text{diag}(\sigma_1^2/n_1, \dots, \sigma_k^2/n_k)$. We obtain a feasible generalized least squares estimate by approximating σ_i^2/n_i by s_i^2 . We used the ‘Simple OLS’ implementation available in FSL that provides inference by comparing the statistic to a Student distribution with $(\sum_{i=1}^k (n_i - 1)) - p$ degrees of freedom [23] as depicted in Table 1. This reference approach will be referred to as **Fixed-effects (FFX) GLM**.

Combining contrast estimates. If the s_i^2 are unavailable, the contrast estimates $\hat{\beta}_i$ can be combined by assuming that the within-study contrast variance σ_i^2/n_i is constant ($\sigma_i^2/n_i = \sigma^2 \forall i$) or negligible in comparison to the between-study variance ($\sigma_i^2/n_i \ll \tau^2$). Then $W = \text{diag}(\sigma_C^2, \dots, \sigma_C^2)$ where σ_C^2 combines the within and between-study variances, i.e. $\sigma_C^2 \simeq \tau^2$ or $\sigma_C^2 \simeq \tau^2 + \sigma^2$ (note, however, in this setting we do not separately estimate τ^2 or σ^2). Under these assumptions, Eq. (1) can be solved by ordinary least squares giving:

$$\hat{\gamma} = (X^T X)^{-1} X^T \hat{\beta} \quad (6)$$

$$\text{Var}(\hat{\gamma}) = (X^T X)^{-1} \sigma_C^2 \quad (7)$$

Given $\hat{\sigma}_C^2$ the unbiased sample variance, we obtain the statistics presented in Table 1 for one sample tests. This approach will be referred to **Random-effects (RFX) GLM** in the following. Inference can be carried out by comparing

	Meta-analytic statistic	Nominal H_0 distrib.	Inputs	Assumptions
MFX GLM	$(\sum \kappa_i \hat{\beta}_i) / \sqrt{\sum_{i=1}^k \kappa_i}$ with $\kappa_i = 1/(\hat{\tau}^2 + s_i^2)$	\mathcal{T}_{k-1}	$\hat{\beta}_i, s_i^2$	IGE, large sample.
RFX GLM	$(\sum_{i=1}^k \frac{\hat{\beta}_i}{\sqrt{k}}) / \widehat{\sigma_C^2}$	\mathcal{T}_{k-1}	$\hat{\beta}_i$	IGE; $\tau^2 + \sigma_i^2 = \sigma_C^2 \forall i$
Contrast Perm.	$(\sum_{i=1}^k \frac{\hat{\beta}_i}{\sqrt{k}}) / \widehat{\sigma_C^2}$	Empirical	$\hat{\beta}_i$	ISE.
FFX GLM	$(\sum_{i=1}^k \frac{\hat{\beta}_i}{s_i^2}) / \sqrt{\sum_{i=1}^k 1/s_i^2}$	$\mathcal{T}_{(\sum_{i=1}^k n_i - 1) - 1}$	$\hat{\beta}_i, s_i^2$	$\tau^2 = 0$, large sample.
Fisher	$-2 \sum_i \ln P_i$	$\chi_{(2k)}^2$	Z_i	$\tau^2 = 0$
Stouffer	$\sqrt{k} \times \frac{1}{k} \sum_i Z_i$	$\mathcal{N}(0, 1)$	Z_i	$\tau^2 = 0$
Weighted Z	$\frac{1}{\sqrt{\sum_i n_i}} \sum_i \sqrt{n_i} Z_i$	$\mathcal{N}(0, 1)$	Z_i, n_i	$\tau^2 = 0$
Z RFX	$(\sum_{i=1}^k Z_i) / \sqrt{k} \hat{\sigma}$	\mathcal{T}_{k-1}	Z_i	IGE; $1 + \tau^2 / \sigma_i^2$ cst.
Z Perm.	$(\sum_{i=1}^k Z_i) / \sqrt{k}$	Empirical	Z_i	ISE.

Table 1: Statistics for one-sample meta-analysis tests and their sampling distributions under the null hypothesis H_0 . Empirical null distributions are determined using permutations with sign flipping. IGE=Independent Gaussian Errors, ISE=Independent Symmetric Errors. Note: $P_i = \Phi(-Z_i)$, $\widehat{\sigma_C^2}$ is the unbiased sample variance.

the RFX GLM statistic to a Student distribution with $k - 1$ degrees of freedom, this result holds asymptotically as well as in small samples [8].

As an alternative to parametric approaches, non-parametric inference [9, 16] can be performed by comparing the one-sample RFX GLM T-statistic to the distribution obtained with “sign flipping”, i.e. randomly multiplying each study’s data by 1 or -1, justified by an assumption of independent studies and symmetrically distributed random error. For two-sample tests, the non-parametric distribution can be obtained by random permutation of the group labels. This approach will be referred to as **Contrast permutation**.

Combining standardised statistics. When only test statistic images are available there are a several alternative approaches available. **Fisher’s** meta-analysis provide a statistic to combine the associated p-values [5]. **Stouffer’s** approach combines directly the standardised statistic [22]. In [28] following [11], the author proposed a weighted method that weights each study’s Z_i by the square root of its sample size [3,7]. This approach will be referred to as **Weighted Z**. All these meta-analytic statistics assumes no or negligible between-study variance and are only suited for one-sample tests. The corresponding statistics and nominal null distributions are presented in Table 1.

As suggested in [1], to get an RFX-like method based on Stouffer’s approach, the standardised statistical estimates Z_i can be combined in an OLS analysis. The corresponding estimate, referred as **Z RFX** is also provided in 1. Non-parametric inference [9, 16] can also be obtained by sign flipping on the Z_i ’s. This approach will be referred to as **Z permutation**.

2.2. Approximations

In practice, all the methods are either approximate in small samples or rely on assumptions that might not be verified in the context of neuroimaging meta-analyses. Methods based on FGLS (MFX GLM and FFX GLM) have approximate parametric null distributions. The nominal distributions of RFX GLM and two-sample contrast permutations are under the (unrealistic) assumption of homogeneous standard errors over studies; even if all studies are ‘clean’ and conducted at the same center, variation in sample size will induce differences in s_i^2 ’s. The fixed-effects methods (Fisher, Stouffer, wieghted Z and FFX GLM) assume homogeneity across studies, i.e. zero between-study variance. All contrast methods (MFX GLM, RFX GLM, Contrast permutation and FFX GLM) require the contrasts to be expressed with in the same units. Finally, RFX Stouffer and Z RFX are ad-hoc methods and their validity has to be investigated.

2.3. Experiments

2.3.1. Validity on null data

We used Monte Carlo simulations to empirically investigate the validity of each estimator. We simulated a set of contrast estimates $\hat{\beta}_i$ and contrast variance estimates s_i^2 according to:

$$\hat{\beta}_i \sim \mathcal{N}(0, \frac{\sigma_i^2}{n} + \tau^2) \quad (8)$$

$$s_i^2 \sim \frac{\sigma_i^2}{n} \frac{\chi_{(n-1)}^2}{n-1} \quad (9)$$

where σ_i^2 was either constant across studies (homoscedasticity) and taken from $\{n \times [0.25, 0.5, 1, 2, 4]\}$ or varying across studies (heteroscedasticity) such as $\max(\sigma_i^2) = \alpha \min(\sigma_i^2)$ with $\alpha \in \{2, 4, 8, 16\}$ and $\text{mean}(\sigma_i^2) = 1$, allowing for 5 different values of σ_i^2 that were linearly spaced and repeated as many times as needed for the specified number of studies k . The between-study variance τ^2 was set to zero (homogeneity) or 1 (heterogeneity). We looked at four different number of studies per meta-analysis $k \in \{5, 10, 25, 50\}$ and two number of subjects $n \in \{20, 100\}$. We set the default number of subjects per studies to $n = 20$ which is a common sample size in existing neuroimaging studies [18]. A total of 144 parameter sets ($9\sigma_i^2 \times 2\tau^2 \times 4k \times 2n_i$) was therefore tested and a total of 1 026 000 realisations was performed for each parameter set.

Three types of analyses were computed: a one-sample meta-analysis testing significance of the mean effect in a group of k studies, a two-sample meta-analysis testing significance in mean differences between two groups of k studies each and an unbalanced two-sample meta-analysis testing significance in mean differences between two groups of $2/5 \times k$ and $8/5 \times k$ respectively. Two-sample analysis assumed a common between-study variance across groups.

We conducted those simulations to evaluate the validity of each estimator in small samples and under violations of their assumptions, such as inhomogeneity of contrast variances s_i^2 or presence of non-negligible between-study variance. Furthermore, we studied the robustness of contrast-based methods to the presence of mismatched units across studies. To simulate units mismatch, each contrast estimates $\hat{\beta}_i$ and contrast variance estimates s_i^2 was replaced by a rescaled version: $\hat{\beta}_i^* = \hat{\beta}_i a_i$ and $s_i^{*2} = s_i^2 a_i^2$. 2 types of unit mismatched were investigated:

- Mismatched contrast vectors: a_i linearly sampled between 0.4 and 1.6 (mean 1).
- Mismatched data scaling algorithms (simulating data from different analysis software): either 20% or 50% of the studies were rescaled with a factor $a_i = 2$ other studies keeping the original scaling.

In the case of two-sample tests, the same mismatch was applied to both groups.

2.3.2. Accuracy on real data

We then used Receiver-Operating-Characteristics (ROC) curves to assess the sensibility of each of the nine meta-analytic estimators on real data. We used a dataset of 21 studies of pain that were initially runned using FSL. This dataset is available at: <http://neurovault.org/collections/XRJCFIYG/>.

Comparability of contrast estimates depends on equivalent scaling of the data, models, and contrast vectors. Data scaling was consistently performed by FSL, setting median brain intensity to 10,000; model were all created by FSL's Feat tool; and contrasts were constructed to preserve units, with sum of positive elements equal to 1, sum of negative elements equal to -1.

We used Neurosynth [27]'s automated large scale coordinate-based meta-analysis of pain as ground truth of areas that should present an effect (cf. <http://neurosynth.org/analyses/terms/pain/>). For each meta-analytic approach, we estimated the true positive rate over a range of thresholds and combined these values to the false positive rates computed on simulated data to create the ROC curves.

The code used to compute the real data analysis is available at: https://github.com/cmaumet/zmeta_rocs.

2.3.3. Software and computing resources

Simulated and real data meta-analyses were computed with Matlab R2016b. The code used to compute the simulations is available at: https://github.com/cmaumet/zmeta_buster and used the IBMA toolbox [12], SPM (RRID:SCR_007037) [6], FSL (RRID:SCR_002823) [10] and SnPM (RRID:SCR_002092) [16]. Simulations were computed on a High Performance Computing cluster provided by the Statistics Department at the University of Warwick. All other analyses were computed on a Mac Book Pro laptop.

Figures displayed in this manuscript were computed with R [20] and used ggplot2 [24] and cowplot [25] packages. The code used to compute the figures displayed in this manuscript is available at: <https://github.com/cmaumet/zmeta> in the form of R jupyter notebooks [17].

3. Results

3.1. Robustness to violation of model assumptions

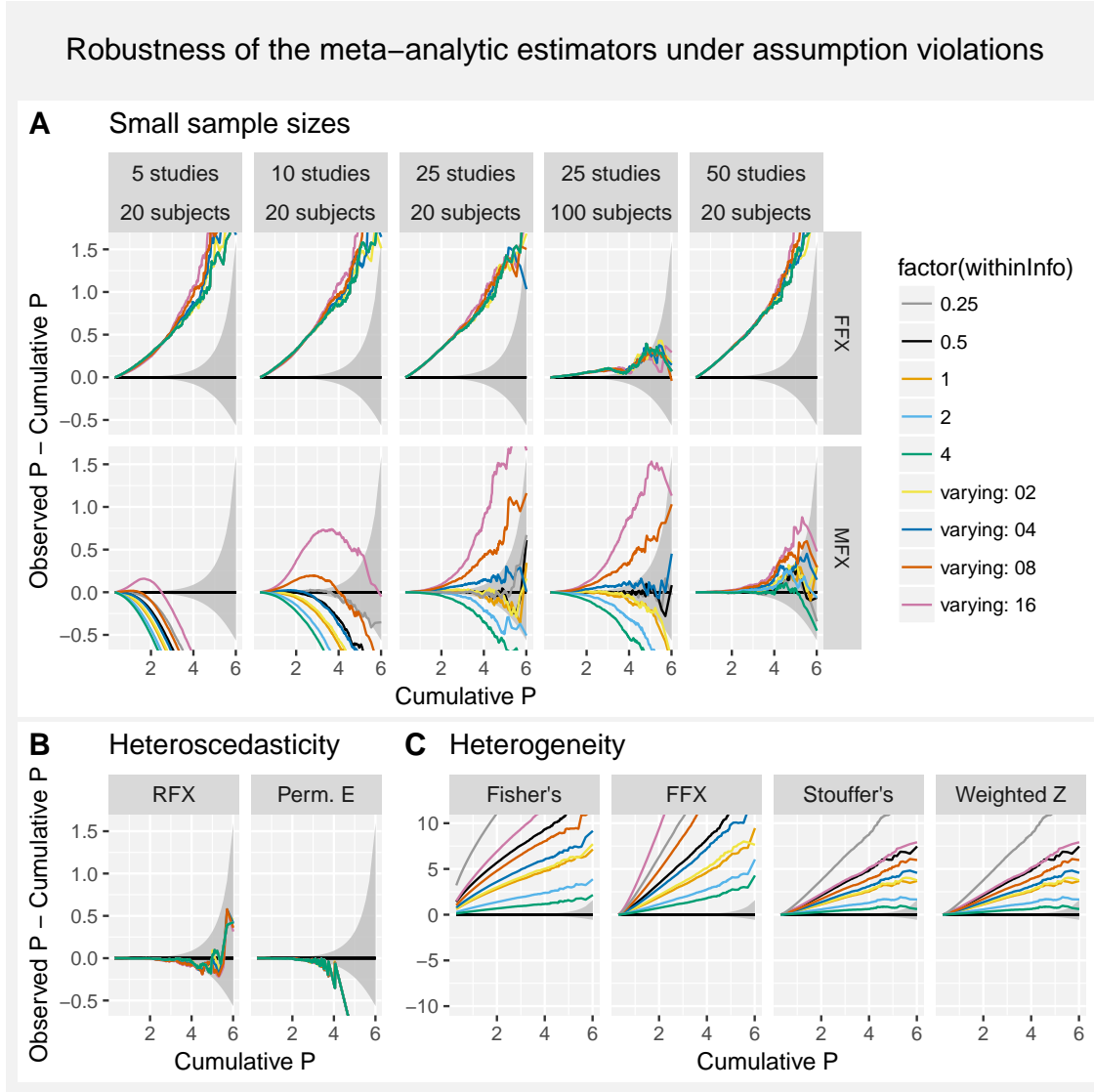


Figure 2: Deviation from theoretical P-values in one-sample tests under violations of the underlying model assumptions: small sample sizes (A), heteroscedasticity (B) and heterogeneity (C). P-values are displayed using a negative log₁₀ scale.

Fig. 2A presents the one-sample simulation results in small samples, i.e. under small number of studies or small number of subjects. We focus here on methods for which validity is only guaranteed in large samples: FFX GLM and MFX GLM, under ideal conditions otherwise (i.e. $\tau^2=0$ for FFX GLM and $\tau^2=1$ for MFX GLM). When the number of subjects is small, FFX GLM is invalid for all within-study variances investigated, regardless of the number of studies included in the meta-analysis. On the other hand MFX GLM is conservative for small number of studies and constant within-study variance. More surprisingly, while MFX GLM is valid for constant within-study variances it is invalid in the presence of large variations in the within-study variances, regardless of the number of subjects included in each study.

Fig. 2B presents the one-sample simulation results under heteroscedasticity (σ_i^2 varying across studies). We focus here on methods for which validity is only guaranteed under homoscedasticity: RFX and Contrast permutation, in a sample of 25 studies with 20 subjects each under ideal conditions otherwise (i.e. $\tau^2=1$). RFX GLM and contrast permutation are robust to heteroscedasticity for all settings studied. RFX GLM is closer to nominal. For small P-values, Contrast Permutation is conservative as expected due to the discrete nature of its distribution.

Fig. 2C presents the one-sample simulation results under heterogeneity ($\tau^2 > 0$). We focus here on methods for which validity is only guaranteed under homogeneity: Fisher, Stouffer, Weighted Z and FFX GLM, with a sample of $k = 25$

studies with $n_i = 20$ subjects each. All fixed-effects methods are invalid under heterogeneity.

Similar behaviours are observed for two-sample tests (cf. Supplementary Fig. S1 and Fig. S2).

3.2. Robustness to units mismatch

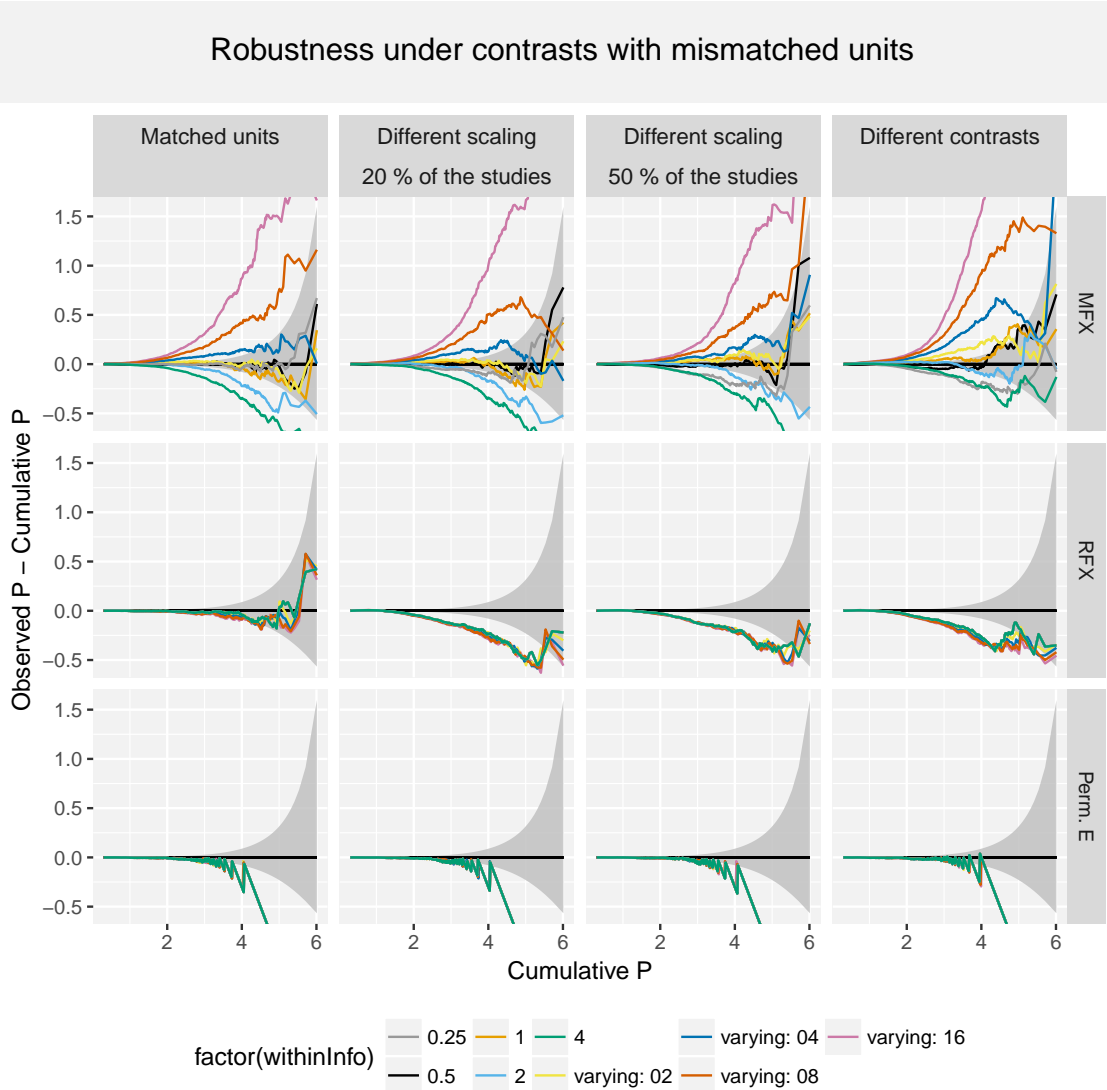


Figure 3: Deviation from theoretical P-values in one-sample tests under violations of the underlying model assumptions: small sample sizes (A), heteroscedasticity (B) and heterogeneity (C). Deviation from theoretical Z in two-sample tests with unit mismatch, under ideal circumstances for each statistical approach ($\tau^2 = 1$ and $k = 5, 25, 50$ with matched (“nominal”) or mismatched (“different scaling target”, “different scaling algorithm”, “different contrast vector scaling”) units.

Fig. 3 presents the simulation results under unit mismatches for one-sample tests. When different scaling algorithm are used (Fig. 3, 2nd and 3rd columns), e.g. with different neuroimaging software packages (provided that differences in scaling targets have been accounted for), Contrast Permutation has a behaviour that is close to nominal. RFX GLM is valid but conservative. MFX GLM is robust to the presence of unit mismatches when the studies are homoscedastic. In the presence of strong heteroscedasticity, MFX GLM remains invalid as when the units are matched due to finite sample inaccuracies (cf. previous section). In the presence of slight heteroscedasticity, unit mismatches can cause invalidity of otherwise valid MFX GLM. When the contrast are scaled differently (Fig. 3, 4th column), we observe a very similar pattern than for different scaling algorithm. Similar behaviours are observed for two-sample tests (cf. Supplementary Fig. S3 and Fig. S4).

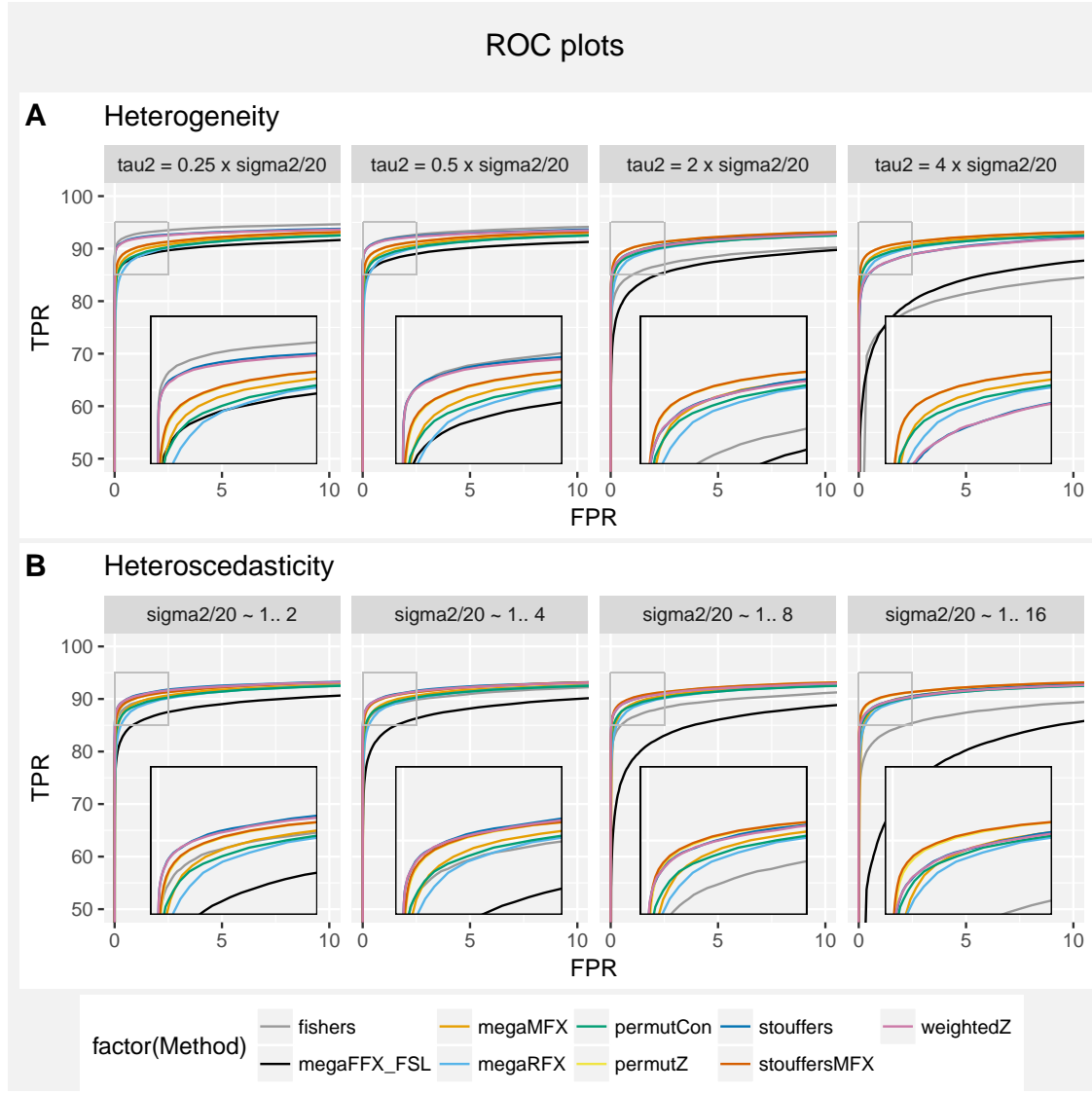


Figure 4: ROC curves of the meta-analytic estimators where true positive rate were computed using a real data meta-analysis of 21 studies of pain and false positive rates using simulated data under various levels of heterogeneity (A) and heteroscedasticity (B).

3.3. Real data

Fig. 4A. displays the ROC curves for all the meta-analytic estimators under varying levels of heterogeneity. As expected, the fixed-effects approaches are the most affected by heterogeneity. Hence, Fisher's method is both the most sensitive method under low heterogeneity and the less sensitive under large heterogeneity. Random-effects approaches are relatively insensitive to the level of heterogeneity. Amongst random-effects approaches the most optimal are Stouffers RFX and Z Permutation that display nearly identical ROC curves, followed by MFX GLM, Contrast Permutation and finally RFX GLM. Differences between MFX GLM and RFX GLM are likely to be non-significant as they both had p-values within the 95% confidence interval.

Fig. 4B. displays the ROC curves for all the meta-analytic estimators under varying levels of heteroscedasticity. Again, the fixed-effects approaches are the most sensitive to heteroscedasticity. This can be explained by the fact that under high heteroscedasticity, some studies will present a low (or high) within-study variance, relatively increasing the between-study variance by comparison to the within-study variance. Amongst the random-effects approaches the most optimal are again Stouffer MFX and Z permutation that display nearly identical ROC curves.

4. Discussions

With the growing availability of summary image data for published neuroimaging studies, image-based meta-analysis becomes feasible. Here, we investigated the validity and accuracy of nine meta-analytic estimators under conditions that are typically observed in fMRI and which might invalidate the underlying assumptions of each method.

As expected, fixed-effects methods were shown to be invalid in the presence of heterogeneity. On the other hand, homoscedastic methods were shown to be robust to heteroscedasticity which is in line with published literature on group fMRI statistics [13]. More surprisingly, MFX GLM was showed to be invalid in the presence of large heteroscedasticity due to its approximations not being accurate in small samples.

In the presence of mismatched units, GLM RFX appeared conservative, while contrast permutation provided the best behaviour, closest to nominal. As confirmed by our real data analysis, we do expect heterogeneity to be present in meta-analytic studies due for instance to variations in the analytic procedure including varying experimental designs, analysis workflows or even due to different imaging instruments. The dataset we used in our real data analysis was created within a single lab and using the same neuroimaging analysis software (FSL), our estimates of heteroscedasticity and heterogeneity are likely to be lower than would be typically observed in a dataset pulling more heterogeneous studies. Similarly, we do expect heteroscedasticity to be present for the same reasons as well as due to varying sample sizes across studies. Finally meta-analysis sample sizes are still relatively small.

Given the still relatively small sample sizes that can be achieved in IBMA as of today, we recommend using RFX GLM, Contrast Permutation, Stouffers MFX or Z Permutation that do not rely on small sample approximations and are robust to both heterogeneity and heteroscedasticity. Unit mismatch across contrasts is likely to be an issue as, even when statistic maps are shared in support of a publication, it is very rare for it to be accompanied by metadata describing full details on the analysis. Although z-based meta-analyses are suboptimal [4] until full reporting is routinely done, we suggest to use Z-based methods that are insensitive to units.

Because true areas of activations in real data are unknown, we relied on an external source to compute the ground truth activations: the Neurosynth platform which provides very large-scale automatic coordinate-based meta-analysis of the literature. Because this ground truth was determined using a coordinate-based (and not image-based) meta-analysis, it is very likely to be missing some true activated areas with small effect sizes (that would effectively not pass the threshold at the level of a single study). Our estimated true positive rates are therefore likely to be overestimations.

Because true areas of no activations in real data are unknown, we relied on simulated data to compute the false positive rates. We only had a single real dataset for meta-analysis and therefore only computed true positive rates once and combined them with the different false positive rates computed by simulations. In practice the sensibility is very likely to vary under different level of heterogeneity, heteroscedasticity and one would need a larger real dataset to fully investigate sensitivity. The most faithful are the ones that correspond to the settings that are closer to the one observed in the real dataset, i.e. is $\sigma^2/20 \approx 1 \dots 16$.

Finally, in our simulations we investigated heteroscedasticity due to varying within-study variances but not due to varying sample sizes. We do expect the results to be consistent (ignoring the X^2 dof) but would have to be double checked.

5. Conclusion

We have compared nine meta-analytic approaches in the context of one-sample test. Through simulations, we found the expected invalidity of fixed-effects approaches in the presence of study heterogeneity, but also of FFX GLM even with no between-study variation. In a real dataset of 21 studies of pain, there was evidence for substantial between-study variation that supports the use of random-effects meta-analytic statistics. When only contrast estimates are available, RFX GLM was valid. When only standardised estimates are available, permutation is the preferred option as the one providing the most faithful results.

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7. References

- [1] Meta-analysis of neuroimaging data: a comparison of image-based and coordinate-based pooling of studies. *NeuroImage*, 45(3):810–23, 2009.

- [2] Katherine S Button, John P a Ioannidis, Claire Mokrysz, Brian a Nosek, Jonathan Flint, Emma S J Robinson, and Marcus R Munafò. Power failure: why small sample size undermines the reliability of neuroscience. *Nature reviews. Neuroscience*, 14(5):365–76, 2013.
- [3] Gang Chen, Paul A Taylor, and Robert W Cox. Is the Statistic Value All We Should Care about in Neuroimaging? *bioRxiv*, (September):064212, 2016.
- [4] P. Cummings. Meta-analysis based on standardized effects is unreliable. *Archives of pediatrics & adolescent medicine*, 158(6):595–7, 2004.
- [5] R.A. Fisher. *Statistical Methods for Research Workers*. Oliver and Boyd, Edinburgh, 1932.
- [6] K.J. Friston, J. Ashburner, S.J. Kiebel, T.E. Nichols, and W.D. Penny, editors. *Statistical Parametric Mapping: The Analysis of Functional Brain Images*. Academic Press, 2007.
- [7] Krzysztof J. Gorgolewski, Gael Varoquaux, Gabriel Rivera, Yannick Schwarz, Satrajit S. Ghosh, Camille Maumet, Vanessa V. Sochat, Thomas E. Nichols, Russell A. Poldrack, Jean-Baptiste Poline, Tal Yarkoni, and Daniel S. Margulies. NeuroVault.org: a web-based repository for collecting and sharing unthresholded statistical maps of the human brain. *Frontiers in Neuroinformatics*, 9(8), April 2015.
- [8] WH William H . Greene. *Econometric analysis*, volume 97. 2012.
- [9] A. P. Holmes, R. C. Blair, G. J.D. Watson, and I. Ford. Nonparametric Analysis of Statistic Images from Functional Mapping Experiments. *Journal of Cerebral Blood Flow and Metabolism*, 16:7–22, 1996.
- [10] Mark Jenkinson, Christian F. Beckmann, Timothy E.J. Behrens, Mark W. Woolrich, and Stephen M. Smith. Fsl. *NeuroImage*, 62(2):782 – 790, 2012. 20 YEARS OF fMRI.
- [11] T. Liptak. On the combination of independent tests. *Magyar Tud. Akad. Mat. Kutato Int. Kozl.*, 3:171–197, 1958.
- [12] Camille Maumet and Thomas E. Nichols. IBMA: An SPM toolbox for NeuroImaging Image-Based Meta-Analysis. In *7th INCF Congress of Neuroinformatics*, volume 8, Leiden, Netherlands, August 2014.
- [13] J. A. Mumford and T. E. Nichols. Simple group fMRI modeling and inference. *NeuroImage*, 47(4):1469–75, 2009.
- [14] Thomas E. Nichols. Spm plot units, 2012.
- [15] Thomas E Nichols, Samir Das, Simon B Eickhoff, Alan C Evans, Tristan Glatard, Michael Hanke, Nikolaus Kriegeskorte, Michael P Milham, Russell A Poldrack, Jean-baptiste Poline, Erika Proal, Bertrand Thirion, David C Van Essen, Tonya White, and B T Thomas Yeo. Best practices in data analysis and sharing in neuroimaging using MRI. *Nature Neuroscience*, 20(3):299–303, feb 2017.
- [16] Thomas E. Nichols and Andrew P Holmes. Nonparametric permutation tests for functional neuroimaging: a primer with examples. *Human brain mapping*, 15(1):1–25, jan 2002.
- [17] Fernando Pérez and Brian E. Granger. IPython: a system for interactive scientific computing. *Computing in Science and Engineering*, 9(3):21–29, May 2007.
- [18] Russell A Poldrack, Chris I Baker, Joke Durnez, Krzysztof J Gorgolewski, Paul M Matthews, Marcus R Munafò, Thomas E Nichols, Jean-baptiste Poline, Edward Vul, and Tal Yarkoni. Scanning the horizon : towards. *Nature Publishing Group*, 2017.
- [19] Russell A. Poldrack and Krzysztof J Gorgolewski. Making big data open : data sharing in neuroimaging. *Nature neuroscience*, 17(11), 2014.
- [20] R Core Team. *R: A Language and Environment for Statistical Computing*. R Foundation for Statistical Computing, Vienna, Austria, 2016.
- [21] J. Radua and D. Mataix-Cols. Meta-analytic methods for neuroimaging data explained. *Biology of mood & anxiety disorders*, 2(1):6, 2012.
- [22] S. Stouffer, L. DeVinney, and E. Suchmen. *The American Soldier: Adjustment During Army Life*, volume 1. Princeton University Press, Princeton, NJ, 1949.

- [23] unknown. Feat/userguide - fslwiki, 2017.
- [24] Hadley Wickham. *ggplot2: Elegant Graphics for Data Analysis*. Springer-Verlag New York, 2009.
- [25] Claus O. Wilke. *cowplot: Streamlined Plot Theme and Plot Annotations for 'ggplot2'*, 2016. R package version 0.7.0.
- [26] Mark W. Woolrich, Timothy E J Behrens, Christian F. Beckmann, Mark Jenkinson, and Stephen M Smith. Multilevel linear modelling for FMRI group analysis using Bayesian inference. *NeuroImage*, 21(4):1732–47, apr 2004.
- [27] Tal Yarkoni, Russell A. Poldrack, Thomas E. Nichols, David C Van Essen, Tor D Wager, and David C Van Essen. Large-scale automated synthesis of human functional neuroimaging data. *Nature methods*, 8(8):665–70, aug 2011.
- [28] D V Zaykin. Optimally weighted Z-test is a powerful method for combining probabilities in meta-analysis. *Journal of evolutionary biology*, 24(8):1836–41, 2011.

	$\hat{\gamma}_1$	$\text{Var}(\hat{\gamma}_1)$
Random effects	$\left(\sum \eta_i \hat{\beta}_i \right) / \left(\sum \eta_i \right)$ with $\eta_i = 1/(\tau^2 + \sigma_i^2/n_i)$	$1/\sum \eta_i$
Fixed effects	$\left(\sum \hat{\beta}_i \times n_i/\sigma_i^2 \right) / \left(\sum n_i/\sigma_i^2 \right)$	$1/(\sum n_i/\sigma_i^2)$

Table S1: One-sample weighted least squares (WLS) estimates and their sampling distributions for random-effects and fixed-effects meta-analyses. The FGLS estimates and assumed sampling distributions are obtained by substituting: $\tau^2 \leftarrow \hat{\tau}^2$ and $\sigma_i^2/n_i \leftarrow s_i^2$.

	$\hat{\gamma}_1 - \hat{\gamma}_2$	$\text{Var}(\hat{\gamma}_1 - \hat{\gamma}_2)$
Random effects	$\left(\sum_{i \in G_1} \eta_i \hat{\beta}_i \right) / \left(\sum_{i \in G_1} \eta_i \right) - \left(\sum_{i \in G_2} \eta_i \hat{\beta}_i \right) / \left(\sum_{i \in G_2} \eta_i \right)$ with $\eta_i = 1/(\tau^2 + \sigma_i^2/n_i)$	$1/\sum_{i \in G_1} \eta_i + 1/\sum_{i \in G_2} \eta_i$
Fixed effects	$\left(\sum_{i \in G_1} \hat{\beta}_i \times n_i/\sigma_i^2 \right) / \left(\sum_{i \in G_1} n_i/\sigma_i^2 \right) - \left(\sum_{i \in G_2} \hat{\beta}_i \times n_i/\sigma_i^2 \right) / \left(\sum_{i \in G_2} n_i/\sigma_i^2 \right)$	$1/\left(\sum_{i \in G_1} n_i/\sigma_i^2 \right) + 1/\left(\sum_{i \in G_2} n_i/\sigma_i^2 \right)$

Table S2: Two-sample weighted least squares (WLS) estimates and their sampling distributions for random-effects and fixed-effects meta-analyses. The FGLS estimates and assumed sampling distributions are obtained by substituting: $\tau^2 \leftarrow \hat{\tau}^2$ and $\sigma_i^2/n_i \leftarrow s_i^2$.

	$\hat{\gamma}_1$	$\text{Var}(\hat{\gamma}_1)$
RFX GLM	$\sum \hat{\beta}_i/k$	σ_C^2/k
Z MFX	$\sum Z_i/k$	σ_C^2/k

Table S3: For the meta-analytic approaches based on Ordinary Least Squares (OLS), one-sample meta-analytic estimates and sampling variances. Note that σ_C^2 will be different for each row of this table.

Robustness of the meta-analytic estimators under assumption violations

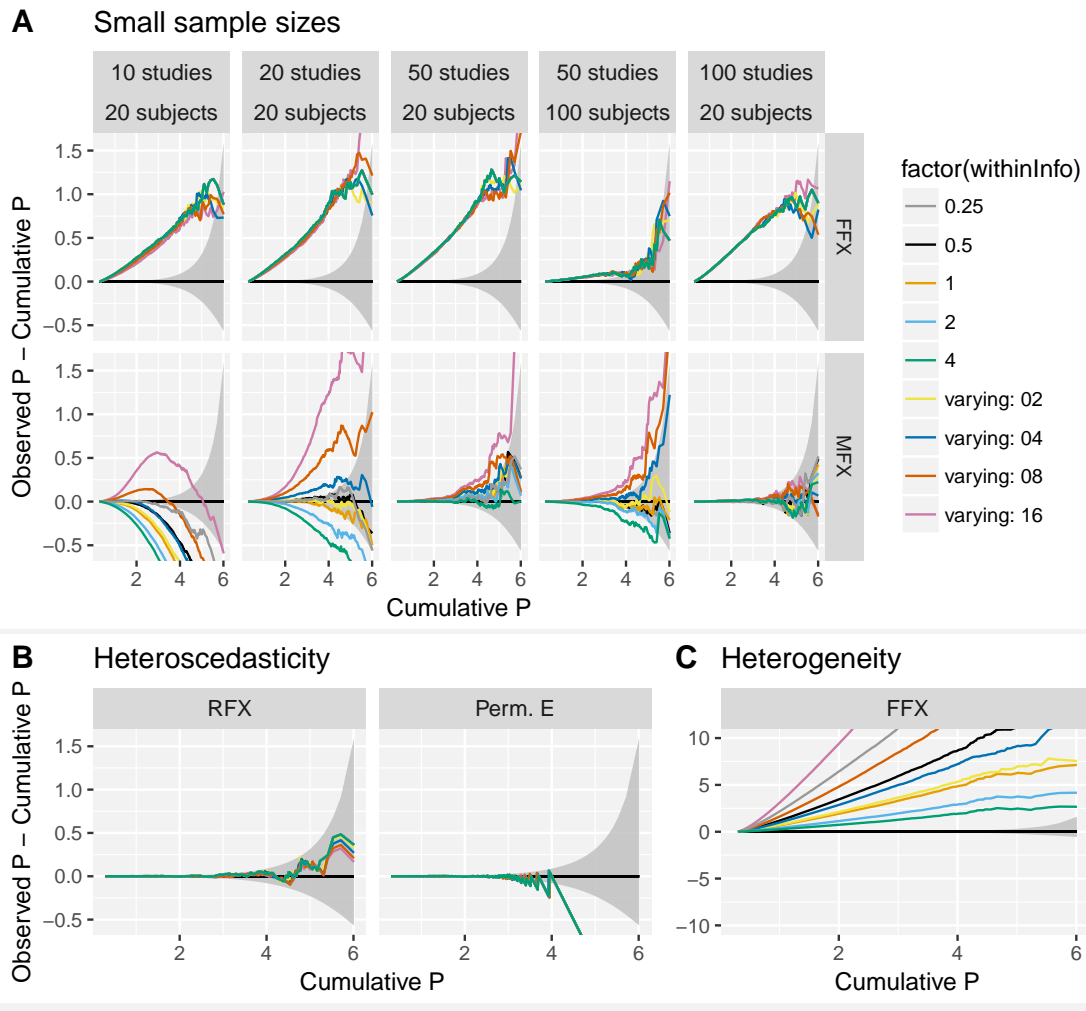


Figure S1: Deviation from theoretical P-values in two-sample tests under violations of the underlying model assumptions: small sample sizes (A), heteroscedasticity (B) and heterogeneity (C). P-values are displayed using a negative \log_{10} scale.

Robustness of the meta-analytic estimators under assumption violations

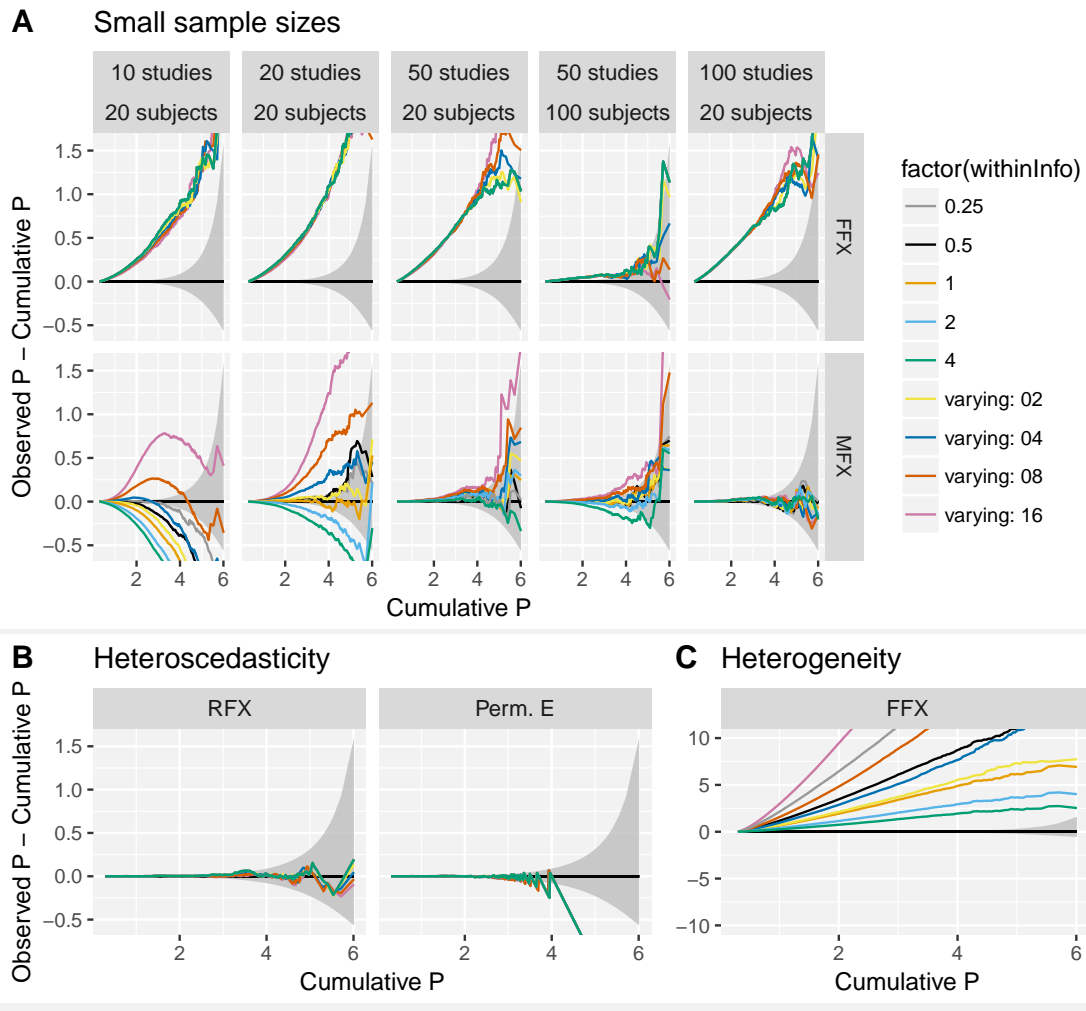


Figure S2: Deviation from theoretical P-values in unbalanced two-sample tests under violations of the underlying model assumptions: small sample sizes (A), heteroscedasticity (B) and heterogeneity (C). P-values are displayed using a negative \log_{10} scale.

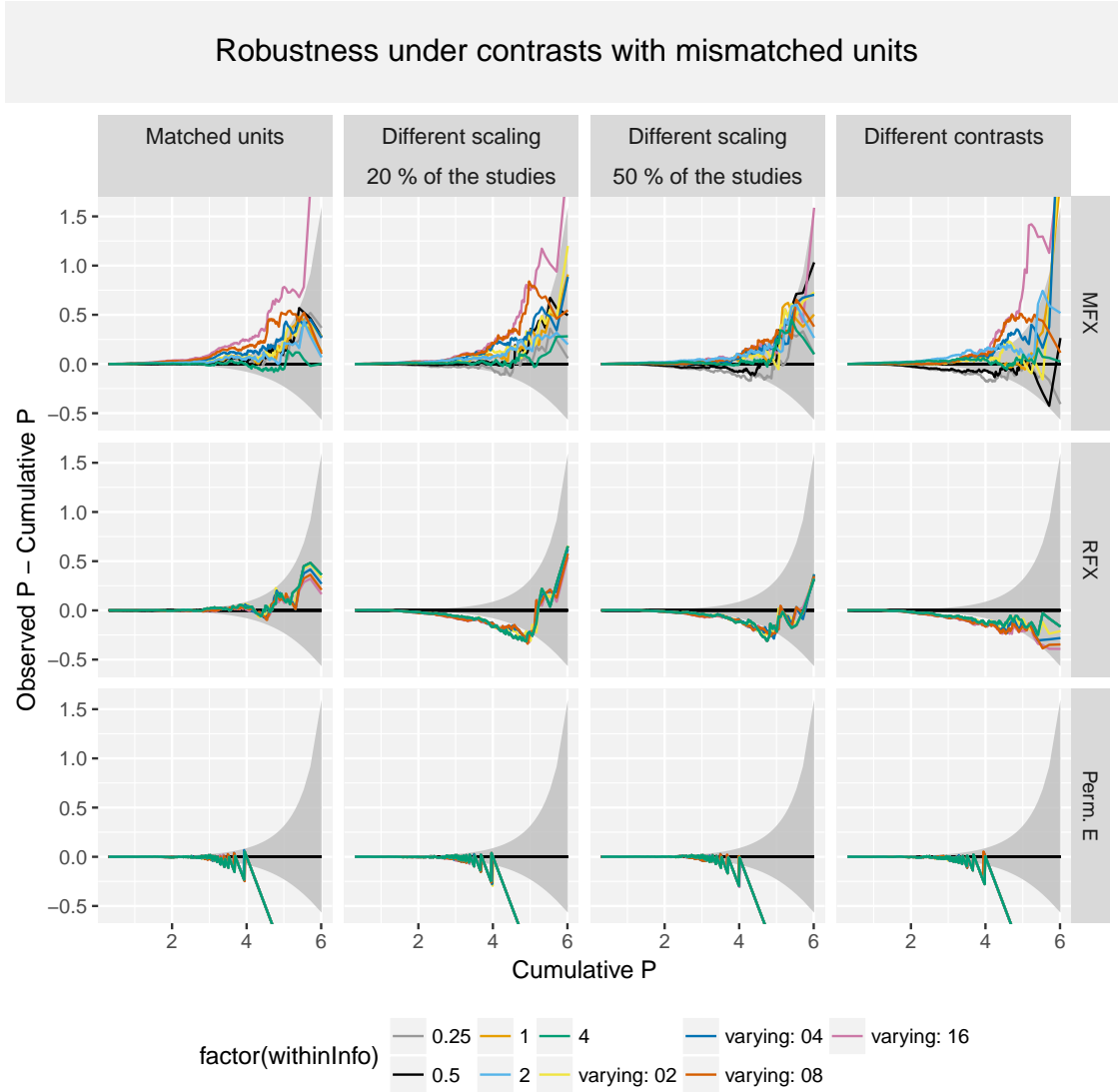


Figure S3: Deviation from theoretical P-values in one-sample tests under violations of the underlying model assumptions: small sample sizes (A), heteroscedasticity (B) and heterogeneity (C). Deviation from theoretical Z in two-sample tests with unit mismatch, under ideal circumstances for each statistical approach ($\tau^2 = 1$ and $k = 5, 25, 50$ with matched (“nominal”) or mismatched (“different scaling target”, “different scaling algorithm”, “different contrast vector scaling”) units).

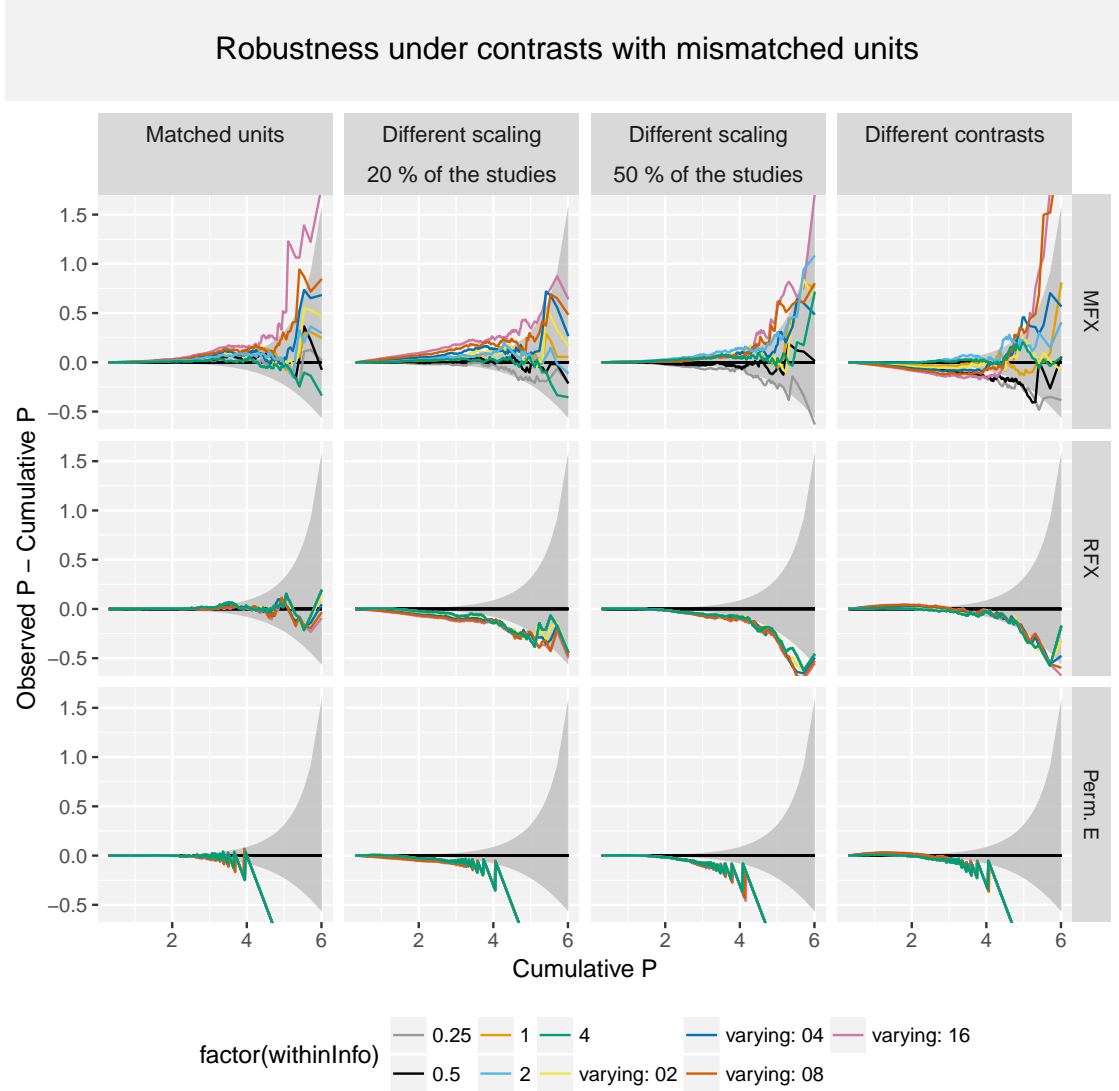


Figure S4: Deviation from theoretical P-values in one-sample tests under violations of the underlying model assumptions: small sample sizes (A), heteroscedasticity (B) and heterogeneity (C). Deviation from theoretical Z in two-sample tests with unit mismatch, under ideal circumstances for each statistical approach ($\tau^2 = 1$ and $k = 5, 25, 50$ with matched (“nominal”) or mismatched (“different scaling target”, “different scaling algorithm”, “different contrast vector scaling”) units).