

# Better-Than-Chance Classification for Signal Detection

Jonathan Rosenblatt      Roei Gilron      Roy Mukamel

August 13, 2016

## Abstract

[TODO]

## 1 Introduction

A common workflow in neuroimaging consists of fitting a classifier, and estimating its predictive accuracy using cross validation. Given that the cross validated accuracy is a random quantity, it is then common to test if the cross validated accuracy is significantly better than chance using a permutation test. Examples in the neuroscientific literature include Golland and Fischl [2003], Pereira et al. [2009], Varoquaux et al. [2016], and especially the recently popularized *multivariate pattern analysis* (MVPA) framework of Kriegeskorte et al. [2006]. This practice is also observed in very high profile publications in the genetics literature: Golub et al. [1999], Slonim et al. [2000], Radmacher et al. [2002], Mukherjee et al. [2003], Juan and Iba [2004], Jiang et al. [2008].

To fix ideas, we will adhere to a concrete example. In Gilron et al. [2016], the authors seek to detect brain regions which encode differences between vocal and non-vocal stimuli. Following the MVPA workflow, the localization problem is cast as a supervised learning problem: if the type of the stimulus can be predicted from the spatial activation pattern significantly better than chance, then a region is declared to encode vocal/non-vocal information. We call this an *accuracy test*, a.k.a. *class prediction*, or *pattern discrimination*.

This same signal detection task can be also approached as a two-group multivariate test. Inferring that a region encodes vocal/non-vocal information, is essentially inferring that the spatial distribution of brain activations is different given a vocal/non-vocal stimulus. As put in Pereira et al. [2009]:

26 ... the problem of deciding whether the classifier learned to dis-  
 27 criminate the classes can be subsumed into the more general ques-  
 28 tion as to whether there is evidence that the underlying distribu-  
 29 tions of each class are equal or not.

30 A practitioner may then call upon a two-group population test such as  
 31 Hotelling’s  $T^2$  [Anderson, 2003]. Alternatively, if the size of a brain re-  
 32 gion is large compared to the number of observations, so that the spatial  
 33 covariance cannot be fully estimated, then a high dimensional version of  
 34 Hotelling’s test can be called upon, such as in Schäfer and Strimmer [2005]  
 35 or Srivastava [2007]. For brevity, and in contrast to *accuracy tests*, we will  
 36 call any two-sample multivariate tests simply *population tests*, also termed  
 37 *class comparisons*. [TODO: rename to parameter test?]

38 At this point, it becomes unclear which is preferable: a population test or  
 39 an accuracy test? The former with a heritage dating back to Hotelling [1931],  
 40 and the latter being extremely popular, as the 959 citations<sup>1</sup> of Kriegeskorte  
 41 et al. [2006] suggest.

42 The comparison between location and accuracy tests was precisely the  
 43 goal of Ramdas et al. [2016], who compared the  $T^2$  population test to the  
 44 accuracy of *Fisher’s linear discriminant analysis* classifier (LDA). By com-  
 45 paring the rates of convergence of the powers to 1, Ramdas et al. [2016]  
 46 concluded that accuracy and population tests are rate equivalent.

47 Asymptotic relative efficiency measures (ARE) are typically used by statis-  
 48 ticians to compare between rate-equivalent test statistics [van der Vaart,  
 49 1998]. Ramdas et al. [2016] derive the asymptotic power functions of the  
 50 two test statistics, which allows to compute the ARE between Hotelling’s  $T^2$   
 51 (location) test and Fisher’s LDA (accuracy) test. Theorem 14.7 of van der  
 52 Vaart [1998] relates asymptotic power functions to ARE. Using the results of  
 53 Ramdas et al. [2016] we deduce that the ARE is lower bounded by  $2\pi \approx 6.3$ .  
 54 This means that Fisher’s LDA requires at least 6.3 more samples to achieve  
 55 the same (asymptotic) power than the  $T^2$  test. In this light, the accuracy  
 56 test is remarkably inefficient compared to the population test. For compar-  
 57 ison, the t-test is only 1.04 more (asymptotically) efficient than Wilcoxon’s  
 58 rank-sum test [Lehmann, 2009], so that an ARE of 6.3 is strong evidence in  
 59 favor of the population test.

60 Before discarding accuracy tests as inefficient, we recall that Ramdas  
 61 et al. [2016] analyzed a *half-sample* holdout. The authors conjectured that a  
 62 leave-one-out approach, which makes more efficient use of the data, may have  
 63 better performance. Also, the analysis in Ramdas et al. [2016] is asymptotic.  
 64 This eschews the discrete nature of the accuracy statistic, which will be

---

<sup>1</sup>GoogleScholar. Accessed on Aug 4, 2016.

65 shown to have crucial impact. Since typical sample sizes in neuroscience are  
 66 not large, we seek to study which test is to be preferred in finite samples?  
 67 Our conclusion will be quite simple: *population tests almost always have more*  
 68 *power than accuracy tests.*

69 Our statement rests upon the observation that with typical sample sizes,  
 70 the accuracy test statistic is highly discrete. Permutation testing with dis-  
 71 crete test statistics are known to be conservative [Hemerik and Goeman,  
 72 2014], since they are insensitive to mild perturbations of the data, and they  
 73 cannot exhaust the permissible false positive rate. The degree of discretiza-  
 74 tion is governed by the number of samples. In our neuroscience example  
 75 from Gilron et al. [2016], the classification is performed based on 40 trials,  
 76 so that the test statistic may assume only 40 possible values. This number  
 77 of examples is not unusual if considering this is the number of trial-repeats,  
 78 or the number of subjects in an neuroimaging study.

79 The discretization effect is aggravated if the test statistic is highly concen-  
 80 trated. For an intuition consider the usage of a the *resubstitution accuracy*  
 81 as a test statistic. This statistic simply means that the accuracy is not cross  
 82 validated. If the data is high dimensional, the resubstitution accuracy will be  
 83 very high due to over fitting. In a very high dimensional model, the resubsti-  
 84 tution accuracy will be 1 for the observed data [McLachlan, 1976, Theorem  
 85 1], but also for any permutation. The concentration of resubstitution accu-  
 86 racy near 1, and its discreteness, render this test completely useless, with a  
 87 power tending to 0 for any (fixed) effect size, as the dimension of the model  
 88 grows.

89 To compare the power of accuracy tests and population tests in finite sam-  
 90 ples, we perform a simulation study of a battery of test statistics. We start  
 91 with formalizing the problem in Section 2. The main findings are reported  
 92 in Sections 4 and 5. A discussion follows in Section 6.

## 93 2 Problem setup

94 Let  $y \in \mathcal{Y}$  be a class encoding. Let  $x \in \mathcal{X}$  be a  $p$  dimensional feature vector.  
 95 In our vocal/non-vocal example we have  $\mathcal{Y} = \{-1, 1\}$  and  $p$ , the number of  
 96 voxels in a brain region so that  $\mathcal{X} = \mathbb{R}^{27}$ .

97 Given  $n$  pairs of  $(x_i, y_i)$ , typically assumed i.i.d., a population test amounts  
 98 to testing whether  $x|y = 1$  has the the same distribution as  $x|y = -1$ . I.e.,  
 99 we test if the multivariate voxel activation pattern has the same distribution  
 100 when given a vocal stimulus, as when given a non-vocal stimulus.

An accuracy test amounts to learning a predictive model  $\hat{f}(x)$  from some  
 assumed model class  $\hat{f} \in \mathcal{F}$ . The prediction accuracy, denoted  $\mathcal{E}_{\hat{f}}$ , is de-

defined as the probability of a given classifier  $\hat{f}$  of making a correct prediction. Denoting by  $I(A)$  the indicator function of the event  $A$ , we get

$$\mathcal{E}_{\hat{f}} := \mathbf{E} \left[ I(\hat{f}(x) = y) \right] \quad (1)$$

when given a randomly drawn data point,  $(x, y)$ . A statistically significant “better than chance” estimate of  $\mathcal{E}_{\hat{f}}$  is evidence that the classes are distinct.

## 2.1 Candidate Tests

The design of a permutation test using the prediction accuracy, requires the following design choices:

1. Is the statistic cross validated or not?
2. For a V-fold cross validated test statistic:
  - (a) Should the data be refolded in each permutation?
  - (b) Should the data folding be balanced (a.k.a. stratified)?
  - (c) How many folds?
3. How to estimate accuracy?

We will now address these questions while bearing in mind that unlike the typical supervised learning setup, we are not interested in an unbiased estimate of the prediction error, but rather in the mere detection of a difference between two groups.

**Cross validate or not?** Since we are merely interested in detecting a difference between classes, a biased error estimate is not an issue provided that bias is consistent over all permutations. The underlying intuition is that if the exact same computation is performed over all permutations, then a permutation test will be “fair”, i.e., will not inflate the false positive rate. We will thus be considering both cross validated accuracies, and resubstitution accuracies as our test statistics.

**Balanced folding?** The standard practice when cross validating is to constrain the data folds to be balanced (i.e. stratified) [e.g. Ojala and Garriga, 2010]. This means that each fold has the same number of examples from each class. We will report results with both balanced and unbalanced data foldings, only to discover, it does not really matter.

128 **Refolding?** The standard practice in neuroimaging is to refold the data  
129 after each permutation, so that data folds are balanced after each label per-  
130 mutation. We will adhere, even though it can be circumvented by permuting  
131 features instead of labels, as done by Golland et al. [2005].

132 **How many folds?** Different authors suggest different rules for the number  
133 of folds. We will be varying the number of folds, and ultimately discover that  
134 the power *decreases with the number of folds*.

**How to estimate accuracy?** Given a predictor  $\hat{f}$ , a natural accuracy test statistic is its accuracy  $\mathcal{E}_{\hat{f}}$ . Since low accuracies, even 0, are evidence that the classes are separated, can consider the departure from chance level,  $|\mathcal{E}_{\hat{f}} - 0.5|$ , as the test statistic. For unbalanced classes, chance level is not 0.5, but rather the probability of the majority class, we denote by  $\hat{p}_{max}$ . This suggests the following test statistic  $|\mathcal{E}_{\hat{f}} - \hat{p}_{max}|$ . Since we will be aggregating these statistics over random data sets where  $\hat{p}_{max}$  may vary, it seems appropriate to standardize the scale of this statistic. We thus propose the z-scored accuracy statistic:

$$|\mathcal{E}_{\hat{f}} - \hat{p}_{max}| / \sqrt{\hat{p}_{max}(1 - \hat{p}_{max})}. \quad (2)$$

135 The of tests we will be comparing is collected for convenience in Table 1.

Name	Basis	CV	Accuracy	Parameters
Hotelling	Hotelling	–	–	–
Hotelling.shrink	Hotelling	–	–	–
lda.CV.1	LDA	V-fold	accuracy	–
lda.CV.2	LDA	V-fold	z-accuracy	–
lda.noCV.1	LDA	–	accuracy	–
lda.noCV.2	LDA	–	z-accuracy	–
sd	SD	–	–	–
svm.CV.1	SVM	V-fold	accuracy	cost=1e1
svm.CV.2	SVM	V-fold	accuracy	cost=1e-1
svm.CV.3	SVM	V-fold	z-accuracy	cost=1e1
svm.CV.4	SVM	V-fold	z-accuracy	cost=1e-1
svm.noCV.1	SVM	–	accuracy	cost=1e1
svm.noCV.2	SVM	–	accuracy	cost=1e-1
svm.noCV.3	SVM	–	z-accuracy	cost=1e1
svm.noCV.4	SVM	–	z-accuracy	cost=1e-1

Table 1: This table collects the various test statistics we will be studying. Three are population tests: Hotelling, Hotelling.shrink, and sd. *Hotelling* is the classical two-group  $T^2$  statistic. *Hotelling.shrink* is a high dimensional version with the regularized covariance in Schäfer and Strimmer [2005]. *sd* is another high dimensional version of the  $T^2$ , from Srivastava et al. [2013]. The rest of the tests are variations of the linear SVM, and Fisher’s LDA, with varying accuracy measures, cross validated or not, and varying tuning parameters. For example, *svm.CV.4* is a linear SVM implemented with the *svm* R function, the cost parameter set at 0.1, and using the cross validated z-scored accuracy in Eq. 2. Another example is *lda.noCV.1*, which is Fisher’s LDA, returning the resubstitution accuracy.

136

### 137 3 Controlling the False Positive Rate

138 Figure 1 demonstrates that all of the tests considered conserve the desired  
139 0.05 false positive rate, up to varying levels of conservatism. This can be  
140 seen from the fact that the probability of rejection is no larger than 0.05 in  
141 the absence of any effect, encoded by a red circle. This is true, in particular  
142 if: (a) the folds are balanced or not, (b) the tuning parameters of some test  
143 statistic are varied, (d) the number of folds is varied. We also observe that  
144 the most conservative tests are the resubstitution accuracy statistics. We  
145 return to this matter in the Discussion.

Figure 1: The power of a permutation test with various test statistics. The power on the  $x$  axis. Effect are color and shape coded. The various statistics on the  $y$  axis. Their details are given in Table 1. Effects vary over 0 (red circle), 0.25 (green triangle), and 0.5 (blue square). Simulation details in Appendix B. Cross-validation was performed with balanced and unbalanced data folding. See sub-captions.



(a) Unbalanced.

(b) Balanced.

## 4 Power

Having established that all of the tests in our battery control the false positive rate, it remains to be seen if they have similar power—especially when comparing population tests to accuracy tests. From the simulation results reported in Appendix C we collect the following insights:

1. population tests have more power than accuracy tests in all our configurations.
2. The conservativeness decays as the sample grows (Figures 9a, 9b and 10a)
3. For heavy tailed distributions (Figure 8b), the extra power of the location test vanishes.
4. The presence of correlations between coordinates reduces the signal to noise ratio (SNR), thus reduces power. More importantly, in the presence of correlations the effect of regularization is amplified, increasing the power difference between regularized and non-regularized test

161 statistics. Put differently- in low SNR regimes, regularization proves  
162 crucial (Figure 10b).

163 5. The z-scoring of the accuracies was introduced to deal with unbalanced  
164 foldings. If the z-scoring has any effect at all, it merely kills power.

165 6. Both accuracy and population tests are inappropriate for scale alter-  
166 natives (Figure 8a). This was to be expected and is reported mostly as  
167 a sanity check.

168 7. Balanced folding only affects the z-scored accuracy, in the opposite  
169 direction than we anticipated.

170 8. Increasing the SVM’s cost parameter, which reduces the number of  
171 support vectors entering the classifier, reduces power.

172 The major insight from simulations is that the use of accuracy tests for  
173 signal detection is underpowered compared to population tests. We now  
174 verify this finding on a neuroimaging dataset.

## 175 5 Neuroimaging Example

176 Figure 2 is an application of both a location and an accuracy test to the data  
177 of Pernet et al. [2015]. The authors of Pernet et al. [2015] collected fMRI  
178 data while subjects were exposed to the sounds of human speech (vocal),  
179 and other non-vocal sounds. Each subject was exposed to 20 sounds of each  
180 type, totaling in  $n = 40$  trials in each scan. The study was rather large and  
181 consisted of about 200 subjects. The data was kindly made available by the  
182 authors at the OpenfMRI website<sup>2</sup>.

183 We perform group inference using within-subject permutations along the  
184 analysis pipeline of Stelzer et al. [2013], which was also reported in Gilron  
185 et al. [2016]. For completeness, the pipeline is described in Appendix A. To  
186 demonstrate our point, we compare the *sd* population test with the *svm.cv.1*  
187 accuracy test.

188 In agreement with our simulation results, the population test (*sd*) discov-  
189 ers more brain regions when compared to an accuracy test (*svm.cv.1*). The  
190 former discovers 1,232 regions, while the latter only 441, as depicted in Fig-  
191 ure 2. We emphasize that both test statistics were compared with the same  
192 permutation scheme, and the same error controls, so that any difference in  
193 detections is due to their different power.

---

<sup>2</sup><https://openfmri.org/>



194 Having established that accuracy tests are typically underpowered for sig-  
 195 nal detection compared to population tests, we wish to identify the conditions  
 196 under which this will occur, and discuss practical implications.



*Figure 2:* Brain regions encoding information discriminating between vocal and non-vocal stimuli. Map reports the centers of 27-voxel sized spherical regions, as discovered by an accuracy test (*svm.cv.1*), and a population test (*sd*). *svm.cv.1* was computed using 5-fold cross validation, and a cost parameter of 1. Region-wise significance was determined using the permutation scheme of Stelzer et al. [2013], followed by region-wise  $FDR \leq 0.05$  control using the Benjamini-Hochberg procedure [Benjamini and Hochberg, 1995]. Number of permutations equals 400. The population test detect 1,232 regions, and the accuracy test 441, 399 of which are common to both. For the details of the analysis see Appendix A and Gilron et al. [2016].

## 197 6 Discussion

198 We have set out to understand which of the tests is more powerful: the  
 199 accuracy test or the population test. No amount of simulations can replace  
 200 the insight provided by a good closed-form analytic result. The finite sample  
 201 power of permutation tests is a formidable mathematical problem, so we  
 202 currently content ourselves with simulations. We have concluded that the  
 203 population tests are typically preferable. Their high dimensional versions,  
 204 such as Srivastava [2007] and Schäfer and Strimmer [2005], are particularly  
 205 well suited for neuroimaging problems such as MVPA. We attribute this

206 to several phenomena: (a) Discretization introduced in finite samples by  
207 the accuracy test statistic. (b) Inefficient use of the data for the validation  
208 holdout set. (c) Regularization crucial in high dimensional problems.

209 The presence of heavy tails shrinks the power advantage of the population  
210 tests over accuracy tests. Our empirical example suggests that even if the  
211 population test does not necessarily dominate the accuracy test in power,  
212 empirically, it does have an advantage.

213 The degree of discretization is governed by the sample size. For this  
214 reason, an asymptotic analysis such as Ramdas et al. [2016] may uncover the  
215 holdout inefficiency, but will not uncover the discretization effect.

216 The practical advice for the practitioner, is that for the purpose of signal  
217 detection, there is typically a population test that is more powerful than  
218 an accuracy test. There is also a good chance that it would be easier to  
219 implement, and faster to run, since no cross validation will be involved.

## 220 6.1 Ease of implementation

221 A very important consideration is the ease of implementation. The need for  
222 cross validation of the accuracy test greatly increases its computational com-  
223 plexity. Moreover, anyone who has actually implemented tests with discrete  
224 statistics, will attest they are more prone to programming errors. This is  
225 because their unforgiveness to the type of inequalities used. Indeed, mistak-  
226 enly replacing a weak inequality with a strong inequality in one’s program  
227 may considerably change the results. This is not the case for continuous test  
228 statistics.

## 229 6.2 Reservations

230 Some reservations to the generality of our findings are in order. Firstly,  
231 not all accuracy tests are concerned with signal detection. Consider brain  
232 decoding for machine interfaces, or clinical diagnosis, where the presence of  
233 a medical condition is predicted from imaging data [e.g. Olivetti et al., 2012,  
234 Wager et al., 2013]. In those examples, the purpose of the test is not to  
235 detect a difference between classes, but to actually test the performance of a  
236 particular classifier.

237 Secondly, it may be argued that accuracy tests permits the separation  
238 between classes in high dimensions, such as in *reproducing kernel Hilbert*  
239 *spaces* (RKHS) by using non-linear predictors. This is a false argument—  
240 accuracy test do not have any more flexibility than population tests. Indeed,  
241 it is possible to test for location in the same dimension the classifier is learned.  
242 Gretton et al. [2012] is an example where the test for location is performed

in the RKHS of the data. It is also possible to test for the equality of two multivariate distributions [TODO: cite vogelstein]. On the other hand, based on our reported neuroimaging example, and others, we find that a population test in the original feature space is indeed a simple and powerful approach to signal detection.

### 6.3 A good accuracy test

For the cases a population test cannot replace an accuracy test, we collect some conclusions and best practices from our simulations. We give particular emphasis in this section to V-fold cross validation due to its popularity, but note that sampling the test set with replacement is actually preferable, as we discuss in Section 6.4.

**Sample size.** The conservativeness of accuracy tests decrease with sample size.

**Permute features.** Permuting features is easier than permuting labels. It allows to preserve balanced folds after a permutation without refolding. Although we did not find a power difference between balanced and unbalanced foldings.

**Use less folds.** For V-fold CV, power decreases as the number of folds increases. This is quite interesting since two phenomena compete as the number of folds increase: (a) the train set is larger so that better accuracies are achievable. (b) The test set is smaller so that the accuracy estimate is more variable. The decrease in power with increase fold number suggests that the latter dominates the former. Put differently: it is easier to detect a small stable departure from chance level, than a large but unstable one.

**Resubstitution accuracy in low dimension.** Resubstitution accuracy is useful in low dimension. In high dimension, the power loss is considerable compared to a cross validated approach. We attribute this to the compounding of discretization and concentration effects: the difference between the sampling distribution of the resubstitution accuracy is simply indistinguishable under the null and under the alternative. In low dimensional problems, the discretization is less impactful, and the computational burden of cross validation can be avoided by using the resubstitution accuracy. There is a fundamental difference between V-folding and resubstitution. The latter should not be thought of as the limit of the former.

277 **Regularize** Regularizing the accuracy test proves very useful in high di-  
 278 mensional problems. Put differently: reducing variance by adding some bias  
 279 is very useful to detect better-than-chance classification.

280 **Don't z-score.** There is no gain in z-scoring the accuracy scores. Our  
 281 motivating rational was clearly flawed. [TODO: why?]

## 282 6.4 Smoothing accuracy estimates

283 It may be possible to alleviate the effect of discretization by appropriate  
 284 cross-validation. The discreteness of the accuracy statistic is governed by  
 285 the number of examples in the union (over all validation iterations) of test  
 286 sets. For V-fold CV, for instance, this number is simply the sample size. This  
 287 suggests that the accuracy can be “smoothed” by allowing the test sample to  
 288 be drawn with replacement. The *bootstrap* may seem like a good candidate  
 289 approach since it samples examples with replacement. It does so, however,  
 290 for the train set, and not the test set. An algorithm that samples test sets  
 291 with replacement is the *leave-one-out bootstrap estimator* (bLOO) and its  
 292 derivation– the *0.632 bootstrap estimator* (b0.632) [Hastie et al., 2003, Sec  
 293 7.11].

**Definition 1** (bLOO). Denoting by  $C^{(i)}$  the index set of bootstrap samples,  
 $b$ , where observation  $i$  is not in the train set, and by  $\hat{f}^b$  the classifier fitted to  
 the  $b$ 'th bootstrap training sample, then the *leave-one-out bootstrap* estimate  
 is defined as:

$$\mathcal{E}_{bLOO} := \frac{1}{n} \sum_{i=1}^n \frac{1}{|C^{(i)}|} \sum_{b \in C^{(i)}} I(\hat{f}^b(x_i) = y_i).$$

Equivalently, denoting by  $S^{(b)}$  the indexes of observations,  $i$ , that are not in  
 the bootstrap train sample  $b$ ,

$$\mathcal{E}_{bLOO} = \frac{1}{B} \sum_{b=1}^B \frac{1}{|S^{(b)}|} \sum_{i \in S^{(b)}} I(\hat{f}^b(x_i) = y_i).$$

**Definition 2** (b0.632). Denoting by  $\mathcal{E}_{resub}$  the resubstitution accuracy esti-  
 mate, the b0.632 accuracy estimator,  $\mathcal{E}_{0.632}$ , is defined as

$$\mathcal{E}_{0.632} := 0.368 \mathcal{E}_{resub} + 0.632 \mathcal{E}_{bLOO}.$$

294 Simulation results reported in Figure 3 with naming conventions in Ta-  
 295 ble 2. It can be seen that selecting test sets with replacement does increase

the power, when compared to V-fold cross validation, but still falls short from the power of population tests. It can also be seen that power increases with the number of bootstrap replications, itself reducing the level of discretization. The type of bootstrap, bLOO versus b0.632, does not change the power.

Name	Basis	Type	B	Accuracy	Parameters
lda.Boot.1	LDA	b0.632	10	accuracy	—
lda.Boot.2	LDA	bLOO	10	accuracy	—
svm.Boot.1	SVM	b0.632	10	accuracy	cost=1e1
svm.Boot.2	SVM	bLOO	10	accuracy	cost=1e1
svm.Boot.3	SVM	b0.632	50	accuracy	cost=1e1
svm.Boot.4	SVM	bLOO	50	accuracy	cost=1e1

Table 2: The same as Table 1 for bootstrapped accuracy estimates. bLOO and b0.632 are defined in definitions 1 and 2 respectively.  $B$  denotes the number of Bootstrap samples.

301

## 6.5 High dimensional classifiers

Inspecting Figure 1a (for instance), it can be seen that Hotelling’s  $T^2$  test has similar power to accuracy tests. It should thus be argued that the real advantage of the population tests is due to their adaptation to high dimension by regularization (*sd* and *Hotelling.shrink*), and not only to discretization. To study this, we call upon several regularized classifiers, designed for high dimensional problems. In the spirit of the regularized covariance of *Hotelling.shrink*, we try an  $l_2$  regularized svm Friedman et al. [2010], and shrinkage based LDA [Pang et al., 2009, Ramey et al., 2016]. In the spirit of the diagonalized covariance of *sd*, we try a diagonalized LDA [Dudoit et al., 2002], which can be thought of a method intersecting Fisher’s LDA and Naive Bayes.

Simulation results reported in Figure 4 with naming conventions in Table 3. It can be seen that regularizing a classifier in high dimension, just like a parameter test, improves power. It can also be seen that (regularized) parameter tests are still more powerful than (regularized) accuracy tests. This was to be expected, since we already saw in (e.g. Figure 1a) that the unregularized parameter test, *Hotelling*, is slightly more powerful than the regularized accuracy test, *svm.CV.1* for instance.

We can compound regularization in this section with the bootstrapping



*Figure 3: Bootstrap*— The power of a permutation test with various test statistics. The power on the  $x$  axis. Effect are color and shape coded. The various statistics on the  $y$  axis. Their details are given in tables 1 and 2. Effects vary over 0 (red circle), 0.25 (green triangle), and 0.5 (blue square). Simulation details in Appendix B.

322 from Section 6.4, to improve finite sample power of the accuracy tests. This  
323 is done in the *svm.highdim.2* test, which still falls short from the power of the  
324 location tests, but is a much more powerful accuracy test than the original  
325 non-regularized, V-fold validated, version of *svm.CV.1*.

Name	Basis	CV	Accuracy	Parameters
svm.highdim.1	SVM	V-fold	accuracy	cost=1e-1
svm.highdim.2	SVM	B=50	accuracy	cost=1e-1
lda.highdim.1	LDA	V-fold	accuracy	—
lda.highdim.2	LDA	V-fold	accuracy	—
lda.highdim.3	LDA	V-fold	accuracy	—

Table 3: The same as Table 1 for regularized (high dimensional) predictors. *svm.highdim.1* is an  $l_2$  regularized SVM Friedman et al. [2010]. *lda.highdim.1* is the Diagonal Linear Discriminant Analysis of Dudoit et al. [2002]. *lda.highdim.2* is the High-Dimensional Regularized Discriminant Analysis of Ramey et al. [2016]. *lda.highdim.3* is the Shrinkage-based Diagonal Linear Discriminant Analysis of Pang et al. [2009].

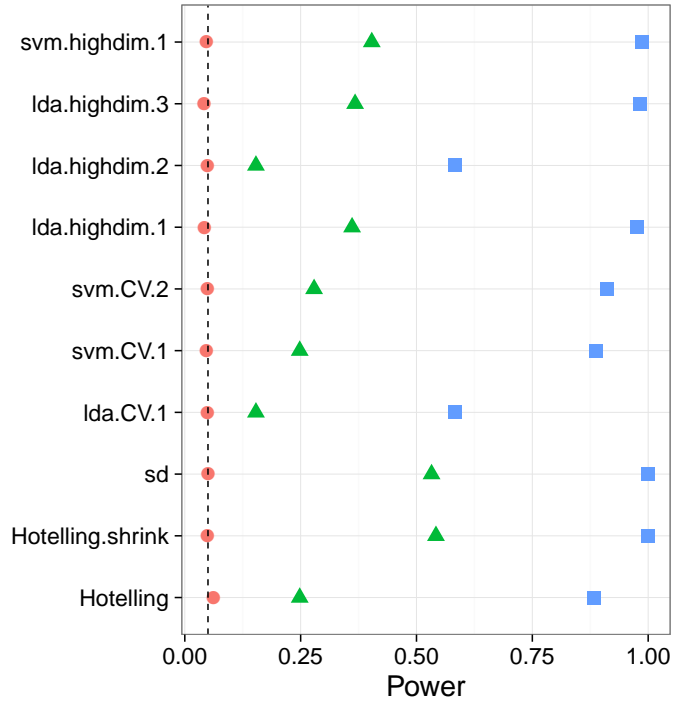


Figure 4: **HighDim Classifier**— The power of a permutation test with various test statistics. The power on the  $x$  axis. Effect are color and shape coded. The various statistics on the  $y$  axis. Their details are given in tables 1 and 3. Effects vary over 0 (red circle), 0.25 (green triangle), and 0.5 (blue square). Simulation details in Appendix B.

## 327 6.6 Related Literature

328 Olivetti et al. [2012] and Olivetti et al. [2014] looked into the problem of  
329 choosing a good accuracy test. They propose a new test they call an *inde-*  
330 *pendence test*, and demonstrate by simulation that it has more power than  
331 other accuracy tests, and can deal with non-balanced data sets. We did not  
332 include this test in the battery we compared, but we note the following: (a)  
333 The independence test of Olivetti et al. [2012] relies on a discrete test statis-  
334 tic. It may thus be improved with the methods discussed in this section,  
335 before the application of Olivetti et al. [2012]’s independence test. (b) In  
336 contrast with the underlying motivation of Olivetti et al. [2012]’s indepen-  
337 dence test, we did not find that balancing the data folds is crucial for an  
338 accuracy test.

339 Golland et al. [2005] study accuracy tests using simulation, neuroimaging  
340 data, genetic data, and analytically. Their analytic results formalize our in-  
341 tuition from Section 1 on the effect of concentration of the accuracy statistic:  
342 The finite Vapnik–Chervonenkis (VC) dimension requirement [Golland and  
343 Fischl, 2003, Sec 4.3] prevents the permutation p-value from (asymptotically)  
344 concentrating near 1. Like ourselves, they also find that the power increases  
345 with the size of the test set (Figure 4, middle). This is seen in their Figure 4,  
346 where the size of the test-set,  $K$ , governs the discretization. Since they per-  
347 mutate features, not labels, then all their permutation samples are balanced,  
348 and there is no issue of refolding.

349 Golland et al. [2005] simulate the power of accuracy tests by sampling  
350 from a Gaussian mixture family of models, and not from a location family  
351 as our own simulations. Under their model  $(x_i|y_i = 1) \sim p\mathcal{N}(\mu_1, I) +$   
352  $(1 - p)\mathcal{N}(\mu_2, I)$  and  $(x_i|y_i = -1) \sim (1 - p)\mathcal{N}(\mu_1, I) + p\mathcal{N}(\mu_2, I)$ . Varying  $p$   
353 interpolates between the null distribution ( $p = 0.5$ ) and a location shift model  
354 ( $p = 0$ ). We now perform the same simulation as Golland et al. [2005], after  
355 reparametrizing  $p$  so that  $p = 0$  corresponds to the null model, and in the  
356 same dimensionality as our previous simulations. We find that also in this  
357 mixture class of models a population test has more power than an accuracy  
358 test (Figure 5).

## 359 6.7 Epilogue

360 Given all the above, we find the popularity of accuracy tests quite puzzling.  
361 We believe this is due to a reversal of the inference cascade. Researchers first  
362 fit a classifier, and then ask if the classes are any different. Were they to  
363 start by asking if classes are any different, and only then try to classify, then  
364 population tests would naturally arise as the preferred method. As put by



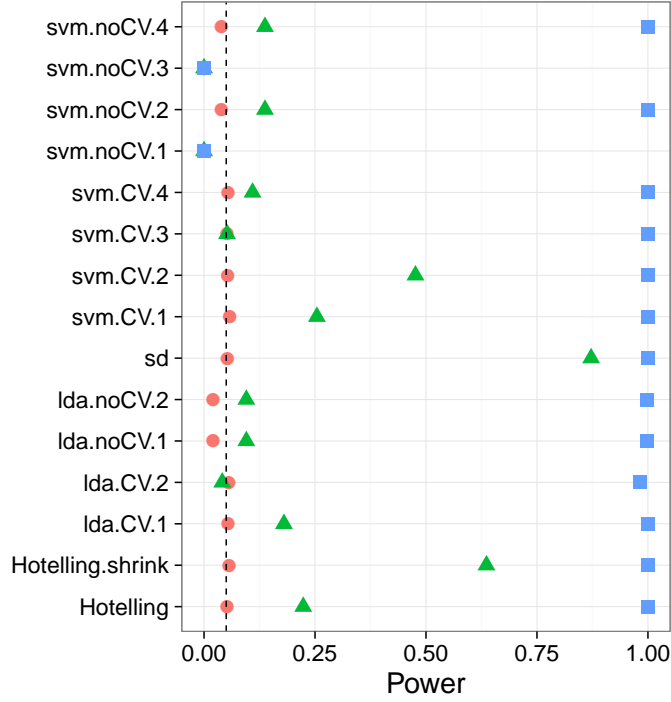


Figure 5: **Mixture**—  $\mathbf{x}_i = \chi_i \mu + \eta_i$ ;  $\chi_i = \{-1, 1\}$  and  $Prob(\chi_i = 1) = (1/2 - p)^{y_i^*} (1/2 + p)^{1-y_i^*}$ .  $\mu$  is a  $p$ -vector with  $3/\sqrt{p}$  in all coordinates. The effect,  $p$ , is color and shape coded and varies over 0 (red circle),  $1/4$  (green triangle) and  $1/2$  (blue square).

365 Ramdas et al. [2016]:

366 The recent popularity of machine learning has resulted in the ex-  
 367 tensive teaching and use of prediction in theoretical and applied  
 368 communities and the relative lack of awareness or popularity of  
 369 the topic of Neyman-Pearson style hypothesis testing in the com-  
 370 puter science and related “data science” communities.

371 And more simply by Frank Harrell in the CrossValidated Q&A site<sup>3</sup>:

372 ... your use of proportion classified correctly as your accuracy  
 373 score. This is a discontinuous improper scoring rule that can be  
 374 easily manipulated because it is arbitrary and insensitive.

## 375 7 Acknowledgments

<sup>3</sup>[http://stats.stackexchange.com/questions/17408/  
 how-to-assess-statistical-significance-of-the-accuracy-of-a-classifier.](http://stats.stackexchange.com/questions/17408/how-to-assess-statistical-significance-of-the-accuracy-of-a-classifier)

## References

- T. W. Anderson. *An Introduction to Multivariate Statistical Analysis*. Wiley-Interscience, Hoboken, NJ, 3 edition edition, July 2003. ISBN 978-0-471-36091-9.
- Y. Benjamini and Y. Hochberg. Controlling the false discovery rate: a practical and powerful approach to multiple testing. *JOURNAL-ROYAL STATISTICAL SOCIETY SERIES B*, 57:289–289, 1995.
- S. Dudoit, J. Fridlyand, and T. P. Speed. Comparison of Discrimination Methods for the Classification of Tumors Using Gene Expression Data. *Journal of the American Statistical Association*, 97(457):77–87, Mar. 2002. ISSN 0162-1459. doi: 10.1198/016214502753479248.
- J. Friedman, T. Hastie, and R. Tibshirani. Regularization Paths for Generalized Linear Models via Coordinate Descent. *Journal of Statistical Software*, 33(1):1–22, 2010.
- R. Gilron, J. Rosenblatt, O. Koyejo, R. A. Poldrack, and R. Mukamel. Quantifying spatial pattern similarity in multivariate analysis using functional anisotropy. *arXiv:1605.03482 [q-bio]*, May 2016.
- P. Golland and B. Fischl. Permutation tests for classification: towards statistical significance in image-based studies. In *IPMI*, volume 3, pages 330–341. Springer, 2003.
- P. Golland, F. Liang, S. Mukherjee, and D. Panchenko. Permutation Tests for Classification. In P. Auer and R. Meir, editors, *Learning Theory*, number 3559 in Lecture Notes in Computer Science, pages 501–515. Springer Berlin Heidelberg, June 2005. ISBN 978-3-540-26556-6 978-3-540-31892-7. doi: 10.1007/11503415\_34.
- T. R. Golub, D. K. Slonim, P. Tamayo, C. Huard, M. Gaasenbeek, J. P. Mesirov, H. Coller, M. L. Loh, J. R. Downing, M. A. Caligiuri, C. D. Bloomfield, and E. S. Lander. Molecular Classification of Cancer: Class Discovery and Class Prediction by Gene Expression Monitoring. *Science*, 286(5439):531–537, Oct. 1999. ISSN 0036-8075, 1095-9203. doi: 10.1126/science.286.5439.531.
- A. Gretton, K. M. Borgwardt, M. J. Rasch, B. Schölkopf, and A. Smola. A Kernel Two-sample Test. *J. Mach. Learn. Res.*, 13:723–773, Mar. 2012. ISSN 1532-4435.

- 410 T. Hastie, R. Tibshirani, and J. Friedman. *The Elements of Statistical Learning*. Springer, July 2003. ISBN 0-387-95284-5.  
411
- 412 J. Hemerik and J. Goeman. Exact testing with random permutations.  
413 *arXiv:1411.7565 [math, stat]*, Nov. 2014.
- 414 H. Hotelling. The Generalization of Student’s Ratio. *The Annals of Mathematical Statistics*, 2(3):360–378, Aug. 1931. ISSN 0003-4851, 2168-8990.  
415 doi: 10.1214/aoms/1177732979.  
416
- 417 W. Jiang, S. Varma, and R. Simon. Calculating confidence intervals for  
418 prediction error in microarray classification using resampling. *Statistical Applications in Genetics and Molecular Biology*, 7(1), 2008.  
419
- 420 L. Juan and H. Iba. Prediction of tumor outcome based on gene expression  
421 data. *Wuhan University Journal of Natural Sciences*, 9(2):177–182, Mar.  
422 2004. ISSN 1007-1202, 1993-4998. doi: 10.1007/BF02830598.
- 423 N. Kriegeskorte, R. Goebel, and P. Bandettini. Information-based functional  
424 brain mapping. *Proceedings of the National Academy of Sciences of the United States of America*, 103(10):3863–3868, July 2006. ISSN 0027-8424,  
425 1091-6490. doi: 10.1073/pnas.0600244103.  
426
- 427 E. L. Lehmann. Parametric versus nonparametrics: two alternative method-  
428 ologies. *Journal of Nonparametric Statistics*, 21(4):397–405, 2009. ISSN  
429 1048-5252. doi: 10.1080/10485250902842727.
- 430 G. J. McLachlan. The bias of the apparent error rate in discriminant analysis.  
431 *Biometrika*, 63(2):239–244, Jan. 1976. ISSN 0006-3444, 1464-3510. doi:  
432 10.1093/biomet/63.2.239.
- 433 S. Mukherjee, P. Tamayo, S. Rogers, R. Rifkin, A. Engle, C. Campbell,  
434 T. R. Golub, and J. P. Mesirov. Estimating dataset size requirements  
435 for classifying DNA microarray data. *Journal of Computational Biology: A Journal of Computational Molecular Cell Biology*, 10(2):119–142, 2003.  
436 ISSN 1066-5277. doi: 10.1089/106652703321825928.  
437
- 438 M. Ojala and G. C. Garriga. Permutation Tests for Studying Classifier Perfor-  
439 mance. *Journal of Machine Learning Research*, 11(Jun):1833–1863, 2010.  
440 ISSN ISSN 1533-7928.
- 441 E. Olivetti, S. Greiner, and P. Avesani. Induction in Neuroscience with  
442 Classification: Issues and Solutions. In G. Langs, I. Rish, M. Grosse-  
443 Wentrup, and B. Murphy, editors, *Machine Learning and Interpretation*

- 444 *in Neuroimaging*, number 7263 in Lecture Notes in Computer Science,  
 445 pages 42–50. Springer Berlin Heidelberg, 2012. ISBN 978-3-642-34712-2  
 446 978-3-642-34713-9. doi: 10.1007/978-3-642-34713-9\_6.
- 447 E. Olivetti, S. Greiner, and P. Avesani. Statistical independence for the  
 448 evaluation of classifier-based diagnosis. *Brain Informatics*, 2(1):13–19, Dec.  
 449 2014. ISSN 2198-4018, 2198-4026. doi: 10.1007/s40708-014-0007-6.
- 450 H. Pang, T. Tong, and H. Zhao. Shrinkage-based Diagonal Discriminant  
 451 Analysis and Its Applications in High-Dimensional Data. *Biometrics*, 65  
 452 (4):1021–1029, Dec. 2009. ISSN 1541-0420. doi: 10.1111/j.1541-0420.2009.  
 453 01200.x.
- 454 F. Pereira, T. Mitchell, and M. Botvinick. Machine learning classifiers and  
 455 fMRI: A tutorial overview. *NeuroImage*, 45(1, Supplement 1):S199–S209,  
 456 Mar. 2009. ISSN 1053-8119. doi: 10.1016/j.neuroimage.2008.11.007.
- 457 C. R. Pernet, P. McAleer, M. Latinus, K. J. Gorgolewski, I. Charest, P. E. G.  
 458 Bestelmeyer, R. H. Watson, D. Fleming, F. Crabbe, M. Valdes-Sosa, and  
 459 P. Belin. The human voice areas: Spatial organization and inter-individual  
 460 variability in temporal and extra-temporal cortices. *NeuroImage*, 119:164–  
 461 174, Oct. 2015. ISSN 1053-8119. doi: 10.1016/j.neuroimage.2015.06.050.
- 462 M. D. Radmacher, L. M. McShane, and R. Simon. A Paradigm for  
 463 Class Prediction Using Gene Expression Profiles. *Journal of Computa-*  
 464 *tional Biology*, 9(3):505–511, June 2002. ISSN 1066-5277. doi: 10.1089/  
 465 106652702760138592.
- 466 A. Ramdas, A. Singh, and L. Wasserman. Classification Accuracy as a Proxy  
 467 for Two Sample Testing. *arXiv:1602.02210 [cs, math, stat]*, Feb. 2016.
- 468 J. A. Ramey, C. K. Stein, P. D. Young, and D. M. Young. High-Dimensional  
 469 Regularized Discriminant Analysis. *arXiv preprint arXiv:1602.01182*,  
 470 2016.
- 471 J. Schäfer and K. Strimmer. A Shrinkage Approach to Large-Scale Covariance  
 472 Matrix Estimation and Implications for Functional Genomics. *Statistical*  
 473 *Applications in Genetics and Molecular Biology*, 4(1), Jan. 2005. ISSN  
 474 1544-6115. doi: 10.2202/1544-6115.1175.
- 475 D. K. Slonim, P. Tamayo, J. P. Mesirov, T. R. Golub, and E. S. Lander. Class  
 476 Prediction and Discovery Using Gene Expression Data. In *Proceedings of*  
 477 *the Fourth Annual International Conference on Computational Molecular*

- 478 *Biology*, RECOMB '00, pages 263–272, New York, NY, USA, 2000. ACM.  
479 ISBN 978-1-58113-186-4. doi: 10.1145/332306.332564.
- 480 M. S. Srivastava. Multivariate Theory for Analyzing High Dimensional Data.  
481 *Journal of the Japan Statistical Society*, 37(1):53–86, 2007. doi: 10.14490/  
482 jjss.37.53.
- 483 M. S. Srivastava, S. Katayama, and Y. Kano. A two sample test in high  
484 dimensional data. *Journal of Multivariate Analysis*, 114:349–358, Feb.  
485 2013. ISSN 0047-259X. doi: 10.1016/j.jmva.2012.08.014.
- 486 J. Stelzer, Y. Chen, and R. Turner. Statistical inference and multiple test-  
487 ing correction in classification-based multi-voxel pattern analysis (MVPA):  
488 Random permutations and cluster size control. *NeuroImage*, 65:69–82, Jan.  
489 2013. ISSN 1053-8119. doi: 10.1016/j.neuroimage.2012.09.063.
- 490 A. W. van der Vaart. *Asymptotic Statistics*. Cambridge University Press,  
491 Cambridge, UK ; New York, NY, USA, Oct. 1998. ISBN 978-0-521-49603-  
492 2.
- 493 G. Varoquaux, P. R. Raamana, D. Engemann, A. Hoyos-Idrobo, Y. Schwartz,  
494 and B. Thirion. Assessing and tuning brain decoders: cross-validation,  
495 caveats, and guidelines. working paper or preprint, June 2016.
- 496 T. D. Wager, L. Y. Atlas, M. A. Lindquist, M. Roy, C.-W. Woo, and E. Kross.  
497 An fMRI-Based Neurologic Signature of Physical Pain. *New England Jour-  
498 nal of Medicine*, 368(15):1388–1397, Apr. 2013. ISSN 0028-4793. doi:  
499 10.1056/NEJMoa1204471.

## 500 A Analysis pipeline

501 Here is the analysis pipeline of Stelzer et al. [2013] we for the auditory data in  
 502 Gilron et al. [2016]. Denoting by  $i = 1, \dots, I$  the subject index,  $v = 1, \dots, V$   
 503 the voxel index, and  $s = 1, \dots, S$  the permutation index. Since regions<sup>4</sup> are  
 504 centered around a unique voxel, the voxel index  $v$  also serves as a unique  
 505 region index. Algorithm 1 computes a region-wise test statistic, which is  
 506 compared to its permutation null distribution computed by Algorithm 2.

**Algorithm 1:** Compute a group parametric map.

**Data:** fMRI scans, and experimental design.  
**Result:** Brain map of group statistics:  $\{\bar{T}_v\}_{v=1}^V$

```

1 for  $v \in 1, \dots, V$  do
2   for  $i \in 1, \dots, I$  do
3      $T_{i,v} \leftarrow$  test statistic for subject  $i$  in a region centered at  $v$ .
4    $\bar{T}_v \leftarrow \frac{1}{I} \sum_{i=1}^I T_{i,v}$ .
```

**Algorithm 2:** Compute a permutation p-value map.

**Data:** fMRI scans of 20 subjects, experimental design.  
**Result:** Brain map of permutation p-values:  $\{p_v\}_{v=1}^V$

```

1 for  $s \in 1, \dots, S$  do
2   permute labels;
3    $\bar{T}_v^s \leftarrow$  parametric map
```

---

<sup>4</sup>*searchlight* or *sphere* in the MVPA parlance

## 509 B Simulation Details

510 The following details are common to all the reported simulations, unless  
511 stated otherwise in a figure’s caption. The R code for the simulations can be  
512 found in [TODO].

513 Each simulation is based on 4,000 replications. In each replication, we  
514 generate  $n$  i.i.d. samples from a shift model  $\mathbf{x}_i = \mu \mathbf{y}_i^* + \eta_i$ . Where  $y_i^* = \{0, 1\}$   
515 is the class of subject  $i$  in dummy coding. Recalling that  $y_i = \{-1, 1\}$  is the  
516 class in effect coding, then clearly  $y_i = 2y_i^* - 1$ . The noise is distributed as  
517  $\eta_i \sim \mathcal{N}_p(0, \Sigma)$ . The sample size  $n = 40$ . The dimension of the data is  $p = 23$ .  
518 The covariance  $\Sigma = I$ . Effects, i.e. shifts  $\mu$ , are equal coordinate  $p$ -vectors  
519 with coordinates that vary over  $\mu \in \{0, 1/4, 1/2\}$ .

520 Having generated the data, we compute each of the test statistics in Ta-  
521 ble 1. For test statistics that require data folding, we used 8 folds. We then  
522 compute a permutation p-value by permuting the class labels, and recomput-  
523 ing each test statistic. We perform 400 such permutations. We then reject  
524 the  $\mu_i = 0$  null hypothesis if the permutation p-value is smaller than 0.05.  
525 The reported power is the proportion of replication where the permutation  
526 p-value falls below 0.05.

## C Simulation Results

Figure 6: Simulation details in Appendix B except the changes in the sub-captions.

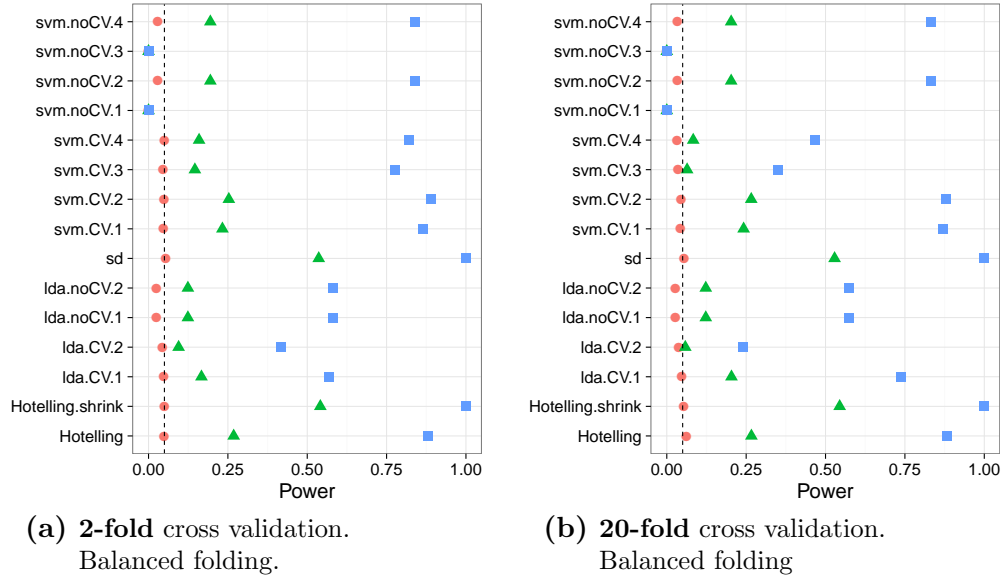




Figure 7: Simulation details in Appendix B except the changes in the sub-captions.



Figure 8: Simulation details in Appendix B except the changes in the sub-captions.

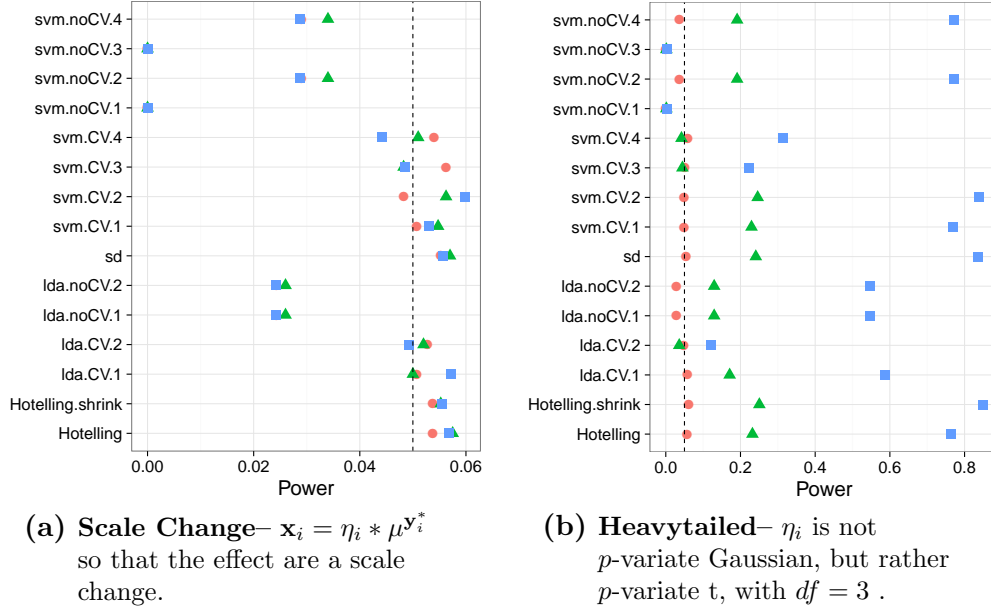
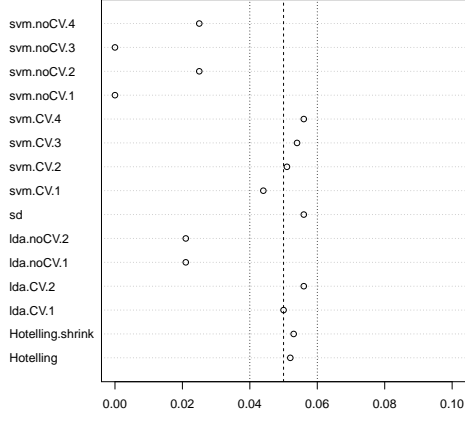
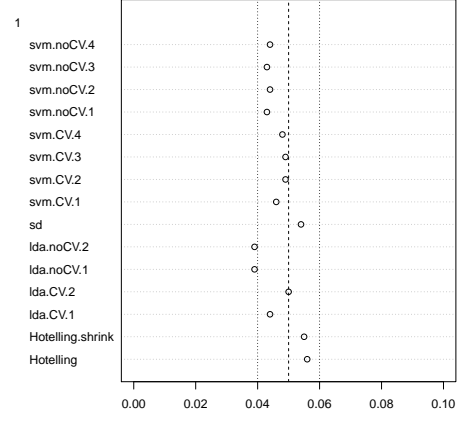


Figure 9: Simulation details in Appendix B except the changes in the sub-captions.

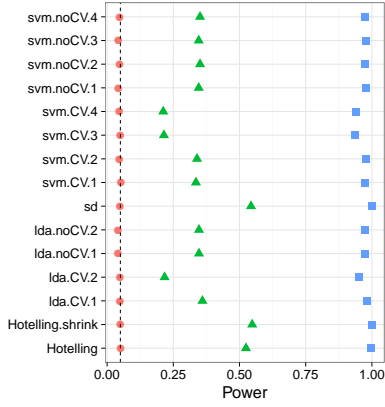


(a) Low-Dimension— False positive rates for  $n = 40$ .

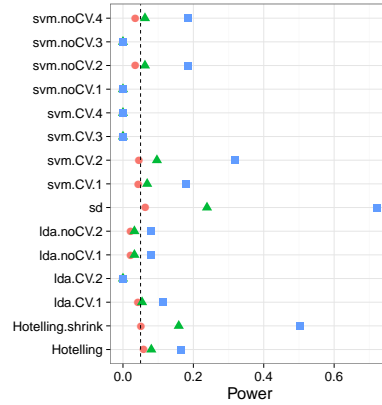


(b) High-Dimension— False positive rates for  $n = 400$ .

Figure 10: Simulation details in Appendix B except the changes in the sub-captions.



(a) High-Dimension, local alternative—  $n = 400$ ,  $\mu \in \frac{1}{\sqrt{10}} \times \{0, 1/4, 1/2\}$ .



(b) AR(1) dependence—  $\Sigma_{k,l} = \rho^{|k-l|}$ ;  $\rho = 0.8$ .