

Supplementary Material

A Missing proofs from Section 2

A.1 Proof of Theorem 2.2

Proof. Fix a feature $f \in F$ and a feature $\tilde{f} \notin F$. Recall we used \mathbf{a}^j to denote the j th column of the test matrix \mathbf{A} . For each row $i \in [t]$, define the random variable $X_i := a_{if}s_i - a_{i\tilde{f}}s_i$, which is the contribution of the i th test to the difference $\rho(f) - \rho(\tilde{f})$. In particular,

$$\rho(f) - \rho(\tilde{f}) = \langle \mathbf{a}^f, \mathbf{s} \rangle - \langle \mathbf{a}^{\tilde{f}}, \mathbf{s} \rangle = \sum_{i=1}^t X_i.$$

The variables X_i are identically and independently distributed. We first estimate $\mathbb{E}[X_i]$. Let T_i denote the i th test, i.e. $T_i = \{j \mid a_{ij} = 1\}$. Then, it is easy to see that $\mathbb{E}[X_i \mid \text{both } f, \tilde{f} \in T_i] = 0$, and $\mathbb{E}[X_i \mid \text{both } f, \tilde{f} \notin T_i] = 0$. Thus, letting $q = 1 - p$ to shorten the notations, we have

$$\begin{aligned} & \mathbb{E}[X_i] \\ &= \mathbb{E}[X_i \mid f \in T_i, \tilde{f} \notin T_i] \cdot \mathbb{P}[f \in T_i, \tilde{f} \notin T_i] + \\ & \quad \mathbb{E}[X_i \mid f \notin T_i, \tilde{f} \in T_i] \cdot \mathbb{P}[f \notin T_i, \tilde{f} \in T_i] \\ &= pq\mathbb{E}[X_i \mid f \in T_i, \tilde{f} \notin T_i] + pq\mathbb{E}[X_i \mid f \notin T_i, \tilde{f} \in T_i] \\ &= pq \left(\sum_{T \subseteq [N] - \{f, \tilde{f}\}} s(T_i) \cdot \mathbb{P}[T_i = T \cup \{f\} \mid f \in T_i, \tilde{f} \notin T_i] \right) \\ & \quad - pq \left(\sum_{T \subseteq [N] - \{f, \tilde{f}\}} s(T_i) \cdot \mathbb{P}[T_i = T \cup \{\tilde{f}\} \mid f \notin T_i, \tilde{f} \in T_i] \right) \\ &= pq \left(\sum_{T \subseteq [N] - \{f, \tilde{f}\}} s(T \cup \{f\}) \cdot p^{|T|} q^{N-2-|T|} \right) \\ & \quad - pq \left(\sum_{T \subseteq [N] - \{f, \tilde{f}\}} s(T \cup \{\tilde{f}\}) \cdot p^{|T|} q^{N-2-|T|} \right) \\ &= pq \left(\sum_{T \subseteq [N] - \{f, \tilde{f}\}} (s(T \cup \{f\}) - s(T \cup \{\tilde{f}\})) \cdot p^{|T|} q^{N-2-|T|} \right) \\ &\geq Cpq \left(\sum_{T \subseteq F - \{f, \tilde{f}\}} p^{|T|} q^{N-2-|T|} \right) = Cpq. \end{aligned}$$

Consequently, when $C \geq 0$ every term in the summation above is non-negative, implying that $\mathbb{E}[X_i] \geq 0$, which in turn implies $\mathbb{E}[\rho(f)] \geq \mathbb{E}[\rho(\tilde{f})]$. Since the X_i are i.i.d. in $[-1, 1]$, by Hoeffding's inequality [8], when $C > 0$ we have

$$\mathbb{P}[\rho(f) - \rho(\tilde{f}) \leq 0] \leq \exp \left\{ \frac{-2t^2 C^2 p^2 q^2}{4t} \right\}.$$

The probability that there is *some* pair $f \in F, \tilde{f} \notin F$ for which $\rho(f) - \rho(\tilde{f}) \leq 0$ is thus at most $d(N - d) \exp \left\{ \frac{-tC^2 p^2 q^2}{2} \right\}$. The last expression is at most δ when t satisfies (1). \square

A.2 Proof of Proposition 2.4 and Proposition 2.5

A special case of separable scoring function is a scoring function satisfying a monotonicity condition: if a subset T_2 of features has more relevant features than another subset T_1 , then $s(T_2)$ has to be better than $s(T_1)$.

Definition A.1 (Monotone scoring function). Let $C \geq 0$ be a real number. The score function $s : 2^{[N]} \rightarrow [0, 1]$ is said to be C -monotone if the following property holds: for any two subsets $T_1, T_2 \subseteq [N]$ such that $T_1 \cap F$ is a proper subset of $T_2 \cap F$, we have $s(T_2) - s(T_1) \geq C$. The 0-monotone scoring functions are called *monotone* for short.

Proposition A.2. *If s is a C -monotone scoring function, then it is a C -separable scoring function.*

Proof. Fix $f \in F$, $\tilde{f} \notin F$, and $T \subset [N] - \{f, \tilde{f}\}$. Let $T_2 = T \cup \{f\}$ and $T_1 = T \cup \{\tilde{f}\}$. Then, $T_1 \cap F \subset T_2 \cap F$. Hence, $s(T_2) - s(T_1) \geq C$, as desired. \square

From the above proposition, to show that a function is separable it is sufficient to show that it is monotone.

Proof of Proposition 2.4. Due to conditional independence, it can be checked that $s(T) = \sum_{f \in T} s(f)$. From this the claim can be easily verified. \square

Proof of Proposition 2.5. Due to a basic property of mutual information, $s(T \cup \{f\}) = I(\mathbf{X}_{T \cup \{f\}}; Y) = H(\mathbf{X}_{T \cup \{f\}}) - H(\mathbf{X}_{T \cup \{f\}}|Y) = H(\mathbf{X}_{T \cup \{f\}}) - H(\mathbf{X}_T|Y) - H(X_f|Y)$, where the last identity is due to the conditional independence assumption. Fix $f \in F$ and $\tilde{f} \notin F$. Since $H(\mathbf{X}_{T \cup \{f\}}) = H(\mathbf{X}_T) + H(X_f) - I(\mathbf{X}_T; X_f)$, we have $s(T \cup \{f\}) = I(\mathbf{X}_T; Y) + I(X_f; Y) - I(\mathbf{X}_T; X_f)$. Combine with a similar formulae for $s(T \cup \{\tilde{f}\})$, we obtain:

$$s(T \cup \{f\}) - s(T \cup \{\tilde{f}\}) = (I(X_f; Y) - I(\mathbf{X}_T; X_f)) - (I(X_{\tilde{f}}; Y) - I(\mathbf{X}_T; X_{\tilde{f}})) \geq 0,$$

which concludes the proof. \square

A.3 On eliminating irrelevant features

The rank $\rho(f)$ of a feature is proportional to the average score of all tests that the feature f participates in. If f is “lucky” enough to participate in tests that contain relevant features, its rank might be inflated. This observation leads to our second idea: we need a way to quickly eliminate features that are likely to be irrelevant.

Theorem A.3. *Let F be the set of hidden relevant features. Let $d = |F|$. Let \mathbf{A} be the random $t \times N$ test matrix obtained by setting each entry to be 1 with probability $p \in [0, 1]$ and 0 with probability $1 - p$. For an irrelevant feature $\tilde{f} \notin F$, let $U_{\tilde{f}}$ denote the total number of tests that \tilde{f} belongs, and $V_{\tilde{f}}$ the total number of tests that \tilde{f} belongs but none of the relevant features belong.*

For any $\delta \in (0, 1)$, and any β such that $0 < \beta < (1 - p)^d$, the following holds:

$$\mathbb{P}[V_{\tilde{f}} \geq \beta U_{\tilde{f}} \text{ for all } \tilde{f} \notin F] \geq 1 - \delta,$$

provided that the total number of tests is at least

$$t \geq \frac{1}{2} \cdot \frac{(1 + \beta)^2}{p^2((1 - p)^d - \beta)^2} \log((N - d)/\delta). \quad (2)$$

Proof. Let \tilde{f} be an arbitrary irrelevant feature. For each $j \in [t]$, let X_j be the indicator variable for the event that \tilde{f} is in test j , and Y_j be the indicator variable for the event that \tilde{f} belongs to the j th test but none of the relevant features are in test j . Then, $U_{\tilde{f}} = \sum_{j \in [t]} X_j$ and $V_{\tilde{f}} = \sum_{j \in [t]} Y_j$. It follows that $\mathbb{E}[Y_j - \beta X_j] = p(1 - p)^d - \beta p$. Furthermore, we have $Y_j - \beta X_j \in [-\beta, 1]$, and for $j \in [t]$ the variables $Y_j - \beta X_j$ are independent. Hence, by Hoeffding bound we have

$$\begin{aligned} \mathbb{P}[V_{\tilde{f}} < \beta U_{\tilde{f}}] &= \mathbb{P}\left[\sum_{j \in [t]} (Y_j - \beta X_j) < 0\right] \\ &\leq \exp\left\{-\frac{2t^2 p^2 ((1 - p)^d - \beta)^2}{t(1 + \beta)^2}\right\} \\ &= \exp\left\{-\frac{2tp^2 ((1 - p)^d - \beta)^2}{(1 + \beta)^2}\right\}. \end{aligned}$$

Hence, due to condition 2,

$$\mathbb{P}[V_{\tilde{f}} < \beta U_{\tilde{f}} \text{ for some } \tilde{f} \notin F] \leq (N - d) \exp\left\{-\frac{2tp^2 ((1 - p)^d - \beta)^2}{(1 + \beta)^2}\right\} \leq \delta.$$

\square

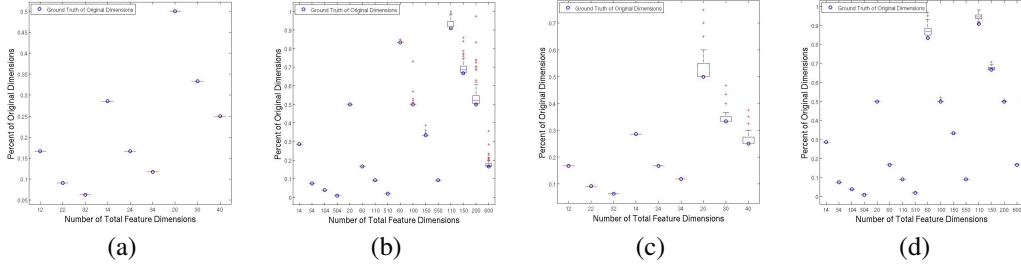


Figure 6: Box plot from synthetic data on a) the identifiability of original features with abundant data from multinomial distribution, b) the identifiability of original features with reasonable-sized data from multinomial distribution, c) the identifiability of original features with abundant data from discretized Gaussian distribution, d) the identifiability of original features with reasonable-sized data from discretized Gaussian distribution.

The above theorem is useful when we can find a score function such that the tests that contain **no** relevant have low scores, say less than some threshold θ . In that case, the natural algorithm is to first eliminate all features such that at least a β fraction of its tests score lower than θ .

To make use of the above algorithm, we need to set the parameters. For example, suppose we set $p = 1/d$. Then $(1 - p)^d = (1 - 1/d)^d$ is an increasing function in d that tends to $1/e \approx 0.37$ fairly quickly. Hence, $d \geq 4$ we can pick $\beta = 0.25$ (or more). But there is a tradeoff between β and the number of tests t , hence we do not want to pick β to be too close to $(1 - p)^d$.

As a second example, suppose we set $p = 1/(2d)$. Then, $(1 - p)^d = (1 - 1/\sqrt{d})^d \rightarrow 1/\sqrt{e} \approx 0.61$. In this case we can even pick $\beta = 1/2$.

B Additional details on synthetic experiment results

We evaluate the entire PFS idea synthetically. We generate a simple categorical binary class dataset using two multinomial distributions. Let N_o be the number of original data dimensions (*i.e.* where the data is actually dependent on). The N_n noisy dimensions are generated with uniform probability for all N_n dimensions, so the synthetic data generated is of dimension $N = N_o + N_n$. The number of trials were restricted to five in our data generation. As theorem 2.2 suggests, by setting $p = 0.5$ we only need logarithmic number of tests with respect to the number of feature dimensions, but it will lead to inaccurate score estimation when N is large and the number of data samples are small. Therefore, we first simulate a case where we have abundant samples by sampling 10,000 samples for each class with $N_o \in \{2, 4, 10\}$ and $N_n \in \{10, 20, 30\}$. We set $p = 0.5$ and $t = \lceil \frac{2}{p^2(1-p)^2} \log(N_o N_n / \delta) \rceil$ where $\delta = 0.01$. In addition, to attain a more realistic setting, we generate 1,000 samples for each class, with $N_o \in \{4, 10, 50, 100\}$ and $N_n \in \{10, 50, 100, 500\}$. We set $p = \frac{3}{N}$ so that we can get reasonable score estimate and $t = 10N$. To account for the randomness of the test, we ran every experiment 100 times; the result is shown in Figure 6(a) and (b) respectively. It is clear that most of the time all the original dimensions are contained in the top D_o ranked features, in particular, when the score can be estimated reasonably well, the top D_o features contains exactly all the original features (see figure 6 (a) and (c)).

Since not all real world data are categorical valued, we simulated another real-valued binary dataset. The N_o original data dimensions are generated from two Gaussian with mean 3 and -3 . They share the same variance, and it is uniformly sampled from the interval $(0, 1]$. The N_n noisy dimensions are generated from Gaussian with mean sample uniformly from interval $[-1, 1]$ and variance uniformly sampled from interval $(0, 1]$. We then quantize each dimension into five equal distanced bins. We use the exactly same settings as the previous experiments, and the result is illustrated in figure 6 (c) and (d), it can be observed that the performance are consistent with the last set of experiments.

C Additional results on the small and medium data sets

C.1 Accuracy and runtime results on Micro-array dataset

The Colon and Leukemia dataset are both binary class dataset that contains 62 samples with 2,000 dimensions and 72 data points with 7,070 dimensions respectively; the Lymph and NCI9 dataset both have 9 classes and

respectively contain 96 samples with 4,026 dimensions and 60 samples with 9,712 dimensions; The Lung dataset contains 73 data samples of 325 dimensions and is a 7-class dataset.

We set the maximum number of selected features to be 50. d Models were trained for each dataset with the top d features where d varies from 1 to 50, and we report the best overall *leave-one-out* classification error among all 50 combinations of features. For the wrapper method we set $p = 10/N$ and for filter method we set $p = 4/N$, where N is the dimension of data.

Table 2: Leave one out error on micro-array datasets from various methods

Method/Dataset	Colon	Leukemia	Lung	Lymph	NCI9
MIM	10	2	14	13	25
MIM (Filtered)	10	2	13	13	25
MRMR	9	2	13	7	23
MRMR (Filtered)	9	2	13	7	23
CMIM	9	1	9	9	26
CMIM (Filtered)	9	1	9	9	26
JMI	9	1	11	9	24
JMI (Filtered)	9	1	11	9	24
DISR	8	1	13	11	24
DISR (Filtered)	8	1	13	11	24
CIFE	9	3	19	26	31
CIFE (Filtered)	9	3	9	10	35
ICAP	8	3	10	8	24
ICAP (Filtered)	7	2	9	9	24
FCBF	9	4	11	6	24
FCBF (Filtered)	1	2	9	14	25
LOGO	10	2	8	12	27
LOGO (Filtered)	7	1	11	11	28
FGM	9	2	8	7	21
FGM (Filtered)	10	1	9	15	25
ours (F_3)	6	1	11	13	31
ours (W_3)	8	0	9	11	28
ours (F_{10})	6	1	11	12	28
ours (W_{10})	9	1	15	13	29

C.2 Accuracy and runtime results on the NIPS Datasets

The results on NIPS dataset from different methods are shown in table 4 below.

Note that the numbers we reported are runtimes *without* running tests in parallel. Since our tests are totally independent, the parallel speed up factor will be essentially linear in the partition size.

The NIPS and micro-array datasets experiments were all completed on a machine with I7-3930K 3.20GHZ 6-core CPU and 32GB RAM with 12 threads. The running time of different methods are listed in the following table 5. and 3.

D Additional results on the large dataset

D.1 Top-features on all relations

As we shown in Figure 2(b), the top-features extracted by our method makes intuitive sense for relations **Spouse**, **MemberOf**, and **TopMember**. Figure 8 shows the result for other relations using the same protocol we described in the body of this paper.

We see that for most relations, the top features selected by our method makes intuitive sense, which implies the effectiveness of our approach. For relations like **per:stateorprovinces_of_residence**, the top keywords are not direct indicator of the relation (although they strongly imply the relation), this is a known problem of how the training set is generated [27, 17], and is orthogonal to the feature selection process.

Table 3: Micro-array dataset runtime performance (in seconds)

Methods/Dataset	Colon	Leukemia	Lung	Lymph	NCI9
MIM	1.77	7.27	0.36	6.17	9.08
MIM (Filtered)	0.23	0.78	0.09	0.70	0.96
MRMR	10.96	50.36	2.02	41.72	57.27
MRMR (Filtered)	1.26	4.99	0.39	4.41	5.78
JMI	17.46	76.29	4.45	90.43	127.78
JMI (Filtered)	1.99	7.75	0.83	9.15	12.93
ICAP	37.80	167.25	8.73	180.59	248.40
ICAP (Filtered)	4.28	17.24	1.65	18.38	25.01
DISR	28.13	118.22	7.20	144.71	206.25
DISR (Filtered)	3.17	12.07	1.36	15.22	20.75
CMIM	1.19	3.47	1.12	5.49	2.78
CMIM (Filtered)	0.77	1.42	0.75	2.64	1.16
CIFE	37.53	166.53	8.85	185.69	259.42
CIFE (Filtered)	4.25	16.76	1.64	18.25	25.73
FCBF	14.01	87.17	34.70	3991.4	838.07
FCBF (Filtered)	2.10	18.76	5.50	114.24	158.39
LOGO	32.51	180.58	66.99	156.66	86.84
LOGO (Filtered)	14.87	27.53	52.37	102.49	53.34
FGM	1.73	3.30	4.54	86.71	142.44
FGM (Filtered)	1.15	1.15	0.83	5.33	5.61
ours (F_3)	1.01	5.35	0.25	5.01	6.61
ours (W_3)	19.72	82.87	46.21	1112.22	1599.15
ours (F_{10})	2.80	14.79	0.74	14.50	19.68
ours (W_{10})	66.71	274.12	153.23	3699.2	5233.84

Table 4: Accuracy results from different methods on NIPS datasets

Methods	Datasets			
	GISETTE		MADELON	
	BER (%)	Features (%)	BER (%)	Features (%)
MIM	3.15	9.40	12.33	2.80
MIM (Filtered)	3.08	6.42	12.33	2.80
MRMR	3.69	8.04	47.83	9.40
MRMR (Filtered)	4.58	4.62	46.17	9.20
JMI	4.02	1.94	11.28	2.00
JMI (Filtered)	4.63	5.62	11.28	2.00
ICAP	4.58	6.24	12.33	2.80
ICAP (Filtered)	4.17	4.62	12.33	2.80
DISR	3.06	7.32	10.61	1.80
DISR (Filtered)	2.92	7.02	14.22	2.60
CMIM	4.46	3.16	12.33	2.80
CMIM (Filtered)	4.82	2.22	12.33	2.80
CIFE	7.82	9.74	39.83	10.20
CIFE (Filtered)	7.80	9.62	39.33	3.60
FCBF	16.86	0.02	45.50	0.20
FCBF (Filtered)	16.86	0.02	45.50	0.20
LOGO	3.09	4.00	43.94	17.80
LOGO (Filtered)	3.40	1.38	21.11	11.60
FGM	2.15	0.70	39.50	9.00
FGM (Filtered)	2.54	1.00	39.11	1.40
ours (F_3)	4.85	9.34	22.61	4.40
ours (W_3)	2.72	6.30	10.17	2.40
ours (F_{10})	4.69	9.94	18.39	1.40
ours (W_{10})	2.89	9.18	10.50	2.40

Table 5: NIPS dataset runtime performance (in seconds)

Methods/Dataset	GISETTE	MADELON
MIM	0.79	0.04
MIM (Filtered)	0.22	0.01
MRMR	23807.11	5.39
MRMR (Filtered)	3439.96	4.68
JMI	5303.93	7.86
JMI (Filtered)	963.97	2.07
ICAP	30901.43	59.68
ICAP (Filtered)	5866.00	20.08
DISR	5533.05	12.49
DISR (Filtered)	864.76	3.16
CMIM	3162.91	12.88
CMIM (Filtered)	3030.55	11.61
CIFE	30658.80	45.52
CIFE (Filtered)	5715.51	17.72
FCBF	107.57	0.94
FCBF (Filtered)	13.65	0.44
LOGO	19303.79	18.17
LOGO (Filtered)	2441.69	9.73
FGM	36.11	6.48
FGM (Filtered)	27.87	4.09
ours (F_3)	3.32	0.23
ours (W_3)	11.03	0.32
ours (F_{10})	10.48	0.72
ours (W_{10})	36.12	1.05

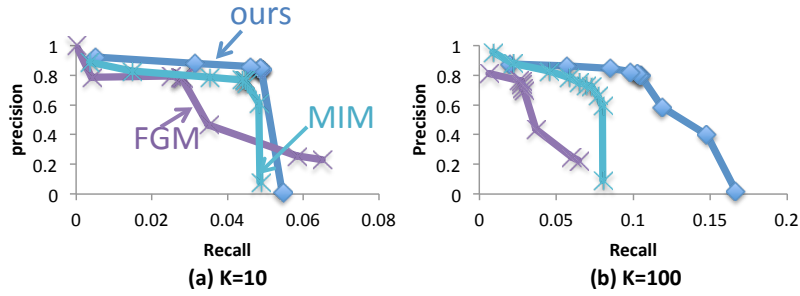


Figure 7: Precision/Recall on TAC-KBP with Number of Features $K = 10$ and $K = 100$.

D.2 On varying the number selected features

Figure 2(a) shows the Precision/Recall on TAC-KBP data set with the number of selected features $K = 1000$. Figure 7 shows the result for $K = 10$ and $K = 100$. We can see that different approaches perform similarly as $K = 1000$ case.

Relation	Keywords					
org:city_of_headquarters	based	headquarters	COXnet	directed	seized	control of
org:founded_by	founder	leader	chairman	co-founder	executive	
org:parents	employees	owned	unit	divisions	subsidiary	
org:subsidiaries	employees	articles	owned	unit	divisions	subsidiary
org:top_members_employees	head	executive	chairman	general	president	
per:children	son	father	daughter	mother	said	
per:city_of_residence	executive	president	chairman	executive	born in	
per:city_of_birth	born	ARodriguez	Peavy	told in	native	
per:city_of_death	died	died home	died hospital	killed	attack city	assassination
per:countries_of_residence	mayor	said in	born in	Democrat	told in	
per:employee_of	executive	chairman	president	professor	director	
per:member_of	leader	member	rebels	commander	iraq leader	
per:parents	son	father	daughter	mother	sons	
per:schools_attended	holds degree from	standout	graduate	student	attend	
per:siblings	brother	sister	half-brother	found along-with	pregnancy	give shops
per:spouse	wife	pictures	husband	married	widower	
per:stateorprovinces_of_residence	governor of	senator from	Republican	Democrat	Republican of	

Figure 8: Top Keywords for All 17 Relations We Considered in TAC-KBP

E Stability Experiments

Given different data samples from the same distribution, the feature selection algorithm should ideally identify the same set of features assuming there is a unique set of “true” features¹⁰. However, due to biases incurred during data sampling and the redundancy present in the data, the algorithm may end up identifying different sets of features leading to inconsistency. Kuncheva [13] presented a *consistency index* which measures the consistency between two sets, with a positive value indicating similar sets, negative value for anti-correlation and zero for random relations.

For measuring the consistency index, we take 50 bootstraps from a dataset and select feature on the bootstraps. The consistency index of the dataset from a particular method is taken as the median value from the 50 bootstraps. The box plot of consistency index from different methods on the 15 UCI datasets are shown in figure 9. In general, filter method has relatively higher stability as compared to wrapper method. The stability measure

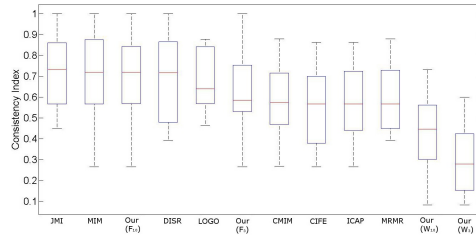


Figure 9: Consistency index across 15 UCI datasets.

of the filter method is very similar to JMI and MIM method, which is attributed to the similarity in obtaining the scores. From the stability measure of our method in figure 9, we can also observe that as we increase the number of tests, the algorithm gets more stable, which confirms the experiments we did in previous section.

¹⁰This will not hold in case there are multiple subsets of features that are equally good.