

Feature Screening via Distance Correlation Learning

Runze LI, Wei ZHONG, and Liping ZHU

This article is concerned with screening features in ultrahigh-dimensional data analysis, which has become increasingly important in diverse scientific fields. We develop a sure independence screening procedure based on the distance correlation (DC-SIS). The DC-SIS can be implemented as easily as the sure independence screening (SIS) procedure based on the Pearson correlation proposed by Fan and Lv. However, the DC-SIS can significantly improve the SIS. Fan and Lv established the sure screening property for the SIS based on linear models, but the sure screening property is valid for the DC-SIS under more general settings, including linear models. Furthermore, the implementation of the DC-SIS does not require model specification (e.g., linear model or generalized linear model) for responses or predictors. This is a very appealing property in ultrahigh-dimensional data analysis. Moreover, the DC-SIS can be used directly to screen grouped predictor variables and multivariate response variables. We establish the sure screening property for the DC-SIS, and conduct simulations to examine its finite sample performance. A numerical comparison indicates that the DC-SIS performs much better than the SIS in various models. We also illustrate the DC-SIS through a real-data example.

KEY WORDS: Sure independence screening; Sure screening property; Ultrahigh dimensionality; Variable selection.

1. INTRODUCTION

Various regularization methods have been proposed for feature selection in high-dimensional data analysis, which has become increasingly frequent and important in various research fields. These methods include, but are not limited to, the LASSO (Tibshirani 1996), the smoothly clipped absolute deviation (SCAD) (Fan and Li 2001; Kim, Choi, and Oh 2008; Zou and Li 2008), the least angle regression (LARS) algorithm (Efron et al. 2004), the elastic net (Zou and Hastie 2005; Zou and Zhang 2009), the adaptive LASSO (Zou 2006), and the Dantzig selector (Candes and Tao 2007). All these methods allow the number of predictors to be greater than the sample size, and perform quite well for high-dimensional data.

With the advent of modern technology for data collection, researchers are able to collect ultrahigh-dimensional data at relatively low cost in diverse fields of scientific research. The aforementioned regularization methods may not perform well for ultrahigh-dimensional data due to the simultaneous challenges of computational expediency, statistical accuracy, and algorithmic stability (Fan, Samworth, and Wu 2009). These challenges call for new statistical modeling techniques for ultrahigh-dimensional data. Fan and Lv (2008) proposed the sure independence screening (SIS) and showed that the Pearson correlation ranking procedure possesses a sure screening

property for linear regressions with Gaussian predictors and responses. That is, all truly important predictors can be selected with probability approaching one as the sample size diverges to ∞ . Hall and Miller (2009) extended the Pearson correlation learning by considering polynomial transformations of predictors. To rank the importance of each predictor, they suggested a bootstrap procedure. Fan, Samworth, and Wu (2009) and Fan and Song (2010) proposed a more general version of independent learning, which ranks the maximum marginal likelihood estimators or the maximum marginal likelihood for generalized linear models. Fan, Feng, and Song (2011) considered nonparametric independence screening in sparse ultrahigh-dimensional additive models. They suggested estimating the nonparametric components marginally with spline approximation, and ranking the importance of predictors using the magnitude of nonparametric components. They also demonstrated that this procedure possesses the sure screening property with vanishing false selection rate. Zhu et al. (2011) proposed a sure independent ranking and screening (SIRS) procedure to screen significant predictors in multi-index models. They further showed that under linearity condition assumption on the predictor vector, the SIRS enjoys the ranking consistency property (i.e., the SIRS can rank the important predictors at the top asymptotically). Ji and Jin (2012) proposed the two-stage method: screening by Univariate thresholding and cleaning by Penalized least squares for Selecting variables, namely UPS. They further theoretically demonstrated that under certain settings, the UPS can outperform the LASSO and subset selection, both of which are one-stage approaches. This motivates us to develop more effective screening procedures using two-stage approaches.

In this article, we propose a new feature screening procedure for ultrahigh-dimensional data based on distance correlation (DC). Székely, Rizzo, and Bakirov (2007) and Székely and Rizzo (2009) showed that the DC of two random vectors equals to zero if and only if these two random vectors are independent. Furthermore, the DC of two univariate normal random variables is a strictly increasing function of the absolute value of

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the Pearson correlation of these two normal random variables. These two remarkable properties motivate us to use the DC for feature screening in ultrahigh-dimensional data. We refer to our SIS procedure based on the DC as the DC-SIS. The DC-SIS can be implemented as easily as the SIS. It is equivalent to the SIS when both the response and the predictor variables are normally distributed. However, the DC-SIS has appealing features that existing screening procedures, including SIS, do not possess. For instance, none of the aforementioned screening procedures can handle grouped predictors or multivariate responses. The proposed DC-SIS can be directly employed for screening grouped variables, and can be directly used for ultrahigh-dimensional data with multivariate responses. Feature screening for multivariate responses and/or grouped predictors is of great interest in pathway analyses. As in Chen et al. (2011), pathway here means sets of proteins that are relevant to specific biological functions without regard to the state of knowledge concerning the interplay among such protein. Since proteins may work interactively to perform various biological functions, pathway analyses complement the marginal association analyses for individual protein, and aim to detect a priori defined set of proteins that are associated with phenotypes of interest. There is a surged interest in pathway analyses in the recent literature (Ashburner et al. 2000; Mootha et al. 2003; Subramanian et al. 2005; Tian et al. 2005; Bild et al. 2006; Efron and Tibshirani 2007; Jones et al. 2008). Thus, it is of importance to develop feature screening procedures for multivariate responses and/or grouped predictors.

We systematically study the theoretical properties of the DC-SIS, and prove that the DC-SIS possesses the sure screening property in the terminology of Fan and Lv (2008) under very general model settings, including linear regression models, for which Fan and Lv (2008) established the sure screening property of the SIS. The sure screening property is a desirable property for feature screening in ultrahigh-dimensional data. Even importantly, the DC-SIS can be used for screening features without specifying a regression model between the response and the predictors. Compared with the model-based screening procedures (Fan and Lv 2008; Fan, Samworth, and Wu 2009; Wang 2009; Fan and Song 2010; Fan, Feng, and Song 2011), the DC-SIS is a model-free screening procedure. This virtue makes the proposed procedure robust to model misspecification. This is a very appealing feature of the proposed procedure in that it may be very difficult to specify an appropriate regression model for the response and the predictors with little information about the actual model in ultrahigh-dimensional data.

We conduct Monte Carlo simulation studies to numerically compare the DC-SIS with the SIS and the SIRS. Our simulation results indicate that the DC-SIS can significantly outperform the SIS and the SIRS under many model settings. We also assess the performance of the DC-SIS as a grouped variable screener, and the simulation results show that the DC-SIS performs very well. We further examine the performance of the DC-SIS for feature screening in ultrahigh-dimensional data with multivariate responses; simulation results demonstrate that screening features for multiple responses jointly may have a dramatic advantage over screening features with each response separately.

The rest of this article is organized as follows. In Section 2, we develop the DC-SIS for feature screening and establish its sure screening property. In Section 3, we examine the finite sample

performance of the DC-SIS via Monte Carlo simulations. We also illustrate the proposed methodology through a real-data example. This article concludes with a brief discussion in Section 4. All technical proofs are given in Appendices A and B.

2. INDEPENDENCE SCREENING USING DC

2.1 Some Preliminaries

Székely, Rizzo, and Bakirov (2007) advocated using the DC for measuring dependence between two random vectors. To be precise, let $\phi_{\mathbf{u}}(\mathbf{t})$ and $\phi_{\mathbf{v}}(\mathbf{s})$ be the respective characteristic functions of the random vectors \mathbf{u} and \mathbf{v} , and $\phi_{\mathbf{u},\mathbf{v}}(\mathbf{t}, \mathbf{s})$ be the joint characteristic function of \mathbf{u} and \mathbf{v} . They defined the distance covariance between \mathbf{u} and \mathbf{v} with finite first moments to be the nonnegative number $\text{dcov}(\mathbf{u}, \mathbf{v})$ given by

$$\text{dcov}^2(\mathbf{u}, \mathbf{v}) = \int_{R^{d_u+d_v}} \|\phi_{\mathbf{u},\mathbf{v}}(\mathbf{t}, \mathbf{s}) - \phi_{\mathbf{u}}(\mathbf{t})\phi_{\mathbf{v}}(\mathbf{s})\|^2 w(\mathbf{t}, \mathbf{s}) d\mathbf{t} d\mathbf{s}, \quad (2.1)$$

where d_u and d_v are the dimensions of \mathbf{u} and \mathbf{v} , respectively, and

$$w(\mathbf{t}, \mathbf{s}) = \{c_{d_u} c_{d_v} \|\mathbf{t}\|_{d_u}^{1+d_u} \|\mathbf{s}\|_{d_v}^{1+d_v}\}^{-1},$$

with $c_d = \pi^{(1+d)/2} / \Gamma\{(1+d)/2\}$. Throughout this article, $\|\mathbf{a}\|_d$ stands for the Euclidean norm of $\mathbf{a} \in \mathbb{R}^d$, and $\|\phi\|^2 = \phi\bar{\phi}$ for a complex-valued function ϕ , with $\bar{\phi}$ being the conjugate of ϕ . The DC between \mathbf{u} and \mathbf{v} with finite first moments is defined as

$$\text{dcorr}(\mathbf{u}, \mathbf{v}) = \frac{\text{dcov}(\mathbf{u}, \mathbf{v})}{\sqrt{\text{dcov}(\mathbf{u}, \mathbf{u})\text{dcov}(\mathbf{v}, \mathbf{v})}}. \quad (2.2)$$

Székely, Rizzo, and Bakirov (2007) systematically studied the theoretical properties of the DC.

Two remarkable properties of the DC motivate us to use it in a feature screening procedure. The first one is the relationship between the DC and the Pearson correlation coefficient. For two univariate normal random variables U and V , with the Pearson correlation coefficient ρ , Székely, Rizzo, and Bakirov (2007) and Székely and Rizzo (2009) showed that

$$\text{dcorr}(U, V) = \left\{ \frac{\rho \arcsin(\rho) + \sqrt{1-\rho^2} - \rho \arcsin(\rho/2) - \sqrt{4-\rho^2+1}}{1+\pi/3-\sqrt{3}} \right\}^{1/2}, \quad (2.3)$$

which is strictly increasing in $|\rho|$. This property implies that the DC-based feature screening procedure is equivalent to the marginal Pearson correlation learning for linear regression with normally distributed predictors and random error. In such a situation, Fan and Lv (2008) showed that the Pearson correlation learning has the sure screening property.

The second remarkable property of the DC is $\text{dcorr}(\mathbf{u}, \mathbf{v}) = 0$ if and only if \mathbf{u} and \mathbf{v} are independent (Székely, Rizzo, and Bakirov 2007). We note that two univariate random variables U and V are independent if and only if U and $T(V)$, a strictly monotone transformation of V , are independent. This implies that a DC-based feature screening procedure can be more effective than the marginal Pearson correlation learning in the presence of nonlinear relationship between U and V . We will demonstrate in the next section that a DC-based screening procedure

is a model-free procedure in that one does not need to specify a model structure between the predictors and the response.

Székely, Rizzo, and Bakirov (2007, remark 3) stated that

$$\text{dcov}^2(\mathbf{u}, \mathbf{v}) = S_1 + S_2 - 2S_3,$$

where S_j , $j = 1, 2$, and 3 , are defined as:

$$\begin{aligned} S_1 &= E\{\|\mathbf{u} - \tilde{\mathbf{u}}\|_{d_u} \|\mathbf{v} - \tilde{\mathbf{v}}\|_{d_v}\}, \\ S_2 &= E\{\|\mathbf{u} - \tilde{\mathbf{u}}\|_{d_u}\} E\{\|\mathbf{v} - \tilde{\mathbf{v}}\|_{d_v}\}, \quad \text{and} \\ S_3 &= E\{E(\|\mathbf{u} - \tilde{\mathbf{u}}\|_{d_u} | \mathbf{u}) E(\|\mathbf{v} - \tilde{\mathbf{v}}\|_{d_v} | \mathbf{v})\}, \end{aligned} \tag{2.4}$$

where $(\tilde{\mathbf{u}}, \tilde{\mathbf{v}})$ is an independent copy of (\mathbf{u}, \mathbf{v}) .

Suppose that $\{(\mathbf{u}_i, \mathbf{v}_i), i = 1, \dots, n\}$ is a random sample from the population (\mathbf{u}, \mathbf{v}) . Székely, Rizzo, and Bakirov (2007) proposed to estimate S_1 , S_2 , and S_3 through the usual moment estimation. To be precise,

$$\begin{aligned} \hat{S}_1 &= \frac{1}{n^2} \sum_{i=1}^n \sum_{j=1}^n \|\mathbf{u}_i - \mathbf{u}_j\|_{d_u} \|\mathbf{v}_i - \mathbf{v}_j\|_{d_v}, \\ \hat{S}_2 &= \frac{1}{n^2} \sum_{i=1}^n \sum_{j=1}^n \|\mathbf{u}_i - \mathbf{u}_j\|_{d_u} \frac{1}{n^2} \sum_{i=1}^n \sum_{j=1}^n \|\mathbf{v}_i - \mathbf{v}_j\|_{d_v}, \quad \text{and} \\ \hat{S}_3 &= \frac{1}{n^3} \sum_{i=1}^n \sum_{j=1}^n \sum_{l=1}^n \|\mathbf{u}_i - \mathbf{u}_l\|_{d_u} \|\mathbf{v}_j - \mathbf{v}_l\|_{d_v}. \end{aligned}$$

Thus, a natural estimator of $\text{dcov}^2(\mathbf{u}, \mathbf{v})$ is given by

$$\widehat{\text{dcov}}^2(\mathbf{u}, \mathbf{v}) = \hat{S}_1 + \hat{S}_2 - 2\hat{S}_3.$$

Similarly, we can define the sample distance covariances $\widehat{\text{dcov}}(\mathbf{u}, \mathbf{u})$ and $\widehat{\text{dcov}}(\mathbf{v}, \mathbf{v})$. Accordingly, the sample DC between \mathbf{u} and \mathbf{v} can be defined by

$$\widehat{\text{dcorr}}(\mathbf{u}, \mathbf{v}) = \frac{\widehat{\text{dcov}}(\mathbf{u}, \mathbf{v})}{\sqrt{\widehat{\text{dcov}}(\mathbf{u}, \mathbf{u})\widehat{\text{dcov}}(\mathbf{v}, \mathbf{v})}}.$$

2.2 An Independence Ranking and Screening Procedure

In this section, we propose an independence screening procedure built upon the DC. Let $\mathbf{y} = (Y_1, \dots, Y_q)^T$ be the response vector with support Ψ_y , and $\mathbf{x} = (X_1, \dots, X_p)^T$ be the predictor vector. We regard q as a fixed number in this context. In an ultrahigh-dimensional setting, the dimensionality p greatly exceeds the sample size n . It is thus natural to assume that only a small number of predictors are relevant to \mathbf{y} . Denote by $F(\mathbf{y} | \mathbf{x})$ the conditional distribution function of \mathbf{y} given \mathbf{x} . Without specifying a regression model, we define the index set of the active and inactive predictors by

$$\begin{aligned} \mathcal{D} &= \{k : F(\mathbf{y} | \mathbf{x}) \text{ functionally depends on } X_k \text{ for some } \mathbf{y} \in \Psi_y\}, \\ \mathcal{I} &= \{k : F(\mathbf{y} | \mathbf{x}) \text{ does not functionally depend on } X_k \text{ for any } \mathbf{y} \in \Psi_y\}. \end{aligned} \tag{2.5}$$

We further write $\mathbf{x}_{\mathcal{D}} = \{X_k : k \in \mathcal{D}\}$ and $\mathbf{x}_{\mathcal{I}} = \{X_k : k \in \mathcal{I}\}$, and refer to $\mathbf{x}_{\mathcal{D}}$ as an *active* predictor vector and its complement $\mathbf{x}_{\mathcal{I}}$ as an *inactive* predictor vector. The index subset \mathcal{D} of all active predictors or, equivalently, the index subset \mathcal{I} of all inactive predictors, is the objective of our primary interest. Definition

(2.5) implies that $\mathbf{y} \perp\!\!\!\perp \mathbf{x}_{\mathcal{I}} | \mathbf{x}_{\mathcal{D}}$, where $\perp\!\!\!\perp$ denotes statistical independence. That is, given $\mathbf{x}_{\mathcal{D}}$, the remaining predictors $\mathbf{x}_{\mathcal{I}}$ are independent of \mathbf{y} . Thus, the inactive predictors $\mathbf{x}_{\mathcal{I}}$ are redundant when the active predictors $\mathbf{x}_{\mathcal{D}}$ are known.

For ease of presentation, we write

$$\omega_k = \text{dcorr}^2(X_k, \mathbf{y}), \quad \text{and} \quad \hat{\omega}_k = \widehat{\text{dcorr}}^2(X_k, \mathbf{y}),$$

for $k = 1, \dots, p$, based on a random sample $\{\mathbf{x}_i, \mathbf{y}_i\}$, $i = 1, \dots, n$. We consider using ω_k as a marginal utility to rank the importance of X_k at the population level. We use the DC because it allows for arbitrary regression relationship of \mathbf{y} onto \mathbf{x} , regardless of whether it is linear or nonlinear. The DC also permits univariate and multivariate responses, regardless of whether it is continuous, discrete, or categorical. In addition, it allows for groupwise predictors. Thus, this DC-based screening procedure is completely model-free. We select a set of important predictors with large $\hat{\omega}_k$. That is, we define

$$\hat{\mathcal{D}}^* = \{k : \hat{\omega}_k \geq cn^{-\kappa}, \text{ for } 1 \leq k \leq p\},$$

where c and κ are prespecified threshold values, which will be defined in Condition (C2) in the subsequent section.

2.3 Theoretical Properties

Next, we study the theoretical properties of the proposed independence screening procedure built upon the DC. The following conditions are imposed to facilitate the technical proofs, although they may not be the weakest ones.

(C1) Both \mathbf{x} and \mathbf{y} satisfy the subexponential tail probability uniformly in p . That is, there exists a positive constant s_0 such that for all $0 < s \leq 2s_0$,

$$\begin{aligned} \sup_p \max_{1 \leq k \leq p} E\{\exp(s\|X_k\|_1^2)\} &< \infty, \quad \text{and} \\ E\{\exp(s\|\mathbf{y}\|_q^2)\} &< \infty. \end{aligned}$$

(C2) The minimum DC of active predictors satisfies

$$\min_{k \in \mathcal{D}} \omega_k \geq 2cn^{-\kappa},$$

for some constants $c > 0$ and $0 \leq \kappa < 1/2$.

Condition (C1) follows immediately when \mathbf{x} and \mathbf{y} are bounded uniformly, or when they have a multivariate normal distribution. The normality assumption has been widely used in the area of ultrahigh-dimensional data analysis to facilitate the technical derivations; see, for example, Fan and Lv (2008) and Wang (2009).

Next we explore Condition (C2). When \mathbf{x} and \mathbf{y} have multivariate normal distribution, (2.3) gives an explicit relationship between the DC and the squared Pearson correlation. For simplicity, we write $\text{dcorr}(X_k, \mathbf{y}) = T_0(|\rho(X_k, \mathbf{y})|)$, where $T_0(\cdot)$ is strictly increasing given in (2.3). In this situation, Condition (C2) requires essentially that $\min_{k \in \mathcal{D}} |\rho(X_k, \mathbf{y})| \geq T_{\text{inv}}(2cn^{-\kappa})$, where $T_{\text{inv}}(\cdot)$ is the inverse function of $T_0(\cdot)$. This is parallel to condition 3 of Fan and Lv (2008) where it is assumed that $\min_{k \in \mathcal{D}} |\rho(X_k, \mathbf{y})| \geq 2cn^{-\kappa}$. This intuitive illustration implies that Condition (C2) requires that the marginal DC of active predictors cannot be too small, which is similar to condition 3 of Fan and Lv (2008). We remark here that, although we illustrate the

intuition by assuming that \mathbf{x} and \mathbf{y} are multivariate normal, we do not require this assumption explicitly in our context. The following theorem establishes the sure screening property for the DC-SIS procedure.

Theorem 1. Under Condition (C1), for any $0 < \gamma < 1/2 - \kappa$, there exist positive constants $c_1 > 0$ and $c_2 > 0$ such that

$$\Pr\left(\max_{1 \leq k \leq p} |\widehat{\omega}_k - \omega_k| \geq cn^{-\kappa}\right) \leq O(p[\exp\{-c_1 n^{1-2(\kappa+\gamma)}\} + n \exp\{-c_2 n^\gamma\}]). \quad (2.6)$$

Under Conditions (C1) and (C2), we have that

$$\Pr(\mathcal{D} \subseteq \widehat{\mathcal{D}}^*) \geq 1 - O(s_n[\exp\{-c_1 n^{1-2(\kappa+\gamma)}\} + n \exp\{-c_2 n^\gamma\}]), \quad (2.7)$$

where s_n is the cardinality of \mathcal{D} .

The sure screening property holds for the DC-SIS under milder conditions than that for the SIS (Fan and Lv 2008) in that we do not require the regression function of \mathbf{y} onto \mathbf{x} to be linear. Thus, the DC-SIS provides a unified alternative to existing model-based sure screening procedures. Compared with the SIRS, the DC-SIS can effectively handle grouped predictors and multivariate responses.

To balance the two terms on the right-hand side of (2.6), we choose the optimal order $\gamma = (1 - 2\kappa)/3$; then, the first part of Theorem 1 becomes

$$\Pr\left(\max_{1 \leq k \leq p} |\widehat{\omega}_k - \omega_k| \geq cn^{-\kappa}\right) \leq O(p[\exp\{-c_1 n^{(1-2\kappa)/3}\}]),$$

for some constant $c_1 > 0$, indicating that we can handle the non-polynomial (NP) dimensionality of order $\log p = o(n^{(1-2\kappa)/3})$. If we further assume that X_k and \mathbf{y} are bounded uniformly in p , then we can obtain without much difficulty that

$$\Pr\left(\max_{1 \leq k \leq p} |\widehat{\omega}_k - \omega_k| \geq cn^{-\kappa}\right) \leq O(p[\exp\{-c_1 n^{1-2\kappa}\}]).$$

In this case, we can handle the NP dimensionality $\log p = o(n^{1-2\kappa})$.

3. NUMERICAL STUDIES

In this section, we assess the performance of the DC-SIS by Monte Carlo simulation. Our simulation studies were conducted using R code. We further illustrate the proposed screening procedure with an empirical analysis of a real-data example.

In Examples 1–3, we generate $\mathbf{x} = (X_1, X_2, \dots, X_p)^T$ from normal distribution with zero mean and covariance matrix $\Sigma = (\sigma_{ij})_{p \times p}$, and the error term ε from standard normal distribution $\mathcal{N}(0, 1)$. We consider two covariance matrices to assess the performance of the DC-SIS and to compare with existing methods: (1) $\sigma_{ij} = 0.8^{|i-j|}$ and (2) $\sigma_{ij} = 0.5^{|i-j|}$. We fix the sample size n to be 200 and vary the dimension p from 2000 to 5000. We repeat each experiment 500 times, and evaluate the performance through the following three criteria:

- (1) \mathcal{S} : the minimum model size to include all active predictors. We report the 5%, 25%, 50%, 75%, and 95% quantiles of \mathcal{S} out of 500 replications.
- (2) \mathcal{P}_s : the proportion that an individual active predictor is selected for a given model size d in the 500 replications.

- (3) \mathcal{P}_a : the proportion that all active predictors are selected for a given model size d in the 500 replications.

The \mathcal{S} is used to measure the model complexity of the resulting model of an underlying screening procedure. The closer to the minimum model size the \mathcal{S} is, the better the screening procedure is. The sure screening property ensures that \mathcal{P}_s and \mathcal{P}_a are both close to one when the estimated model size d is sufficiently large. We choose d to be $d_1 = \lceil n/\log n \rceil$, $d_2 = 2\lceil n/\log n \rceil$, and $d_3 = 3\lceil n/\log n \rceil$ throughout our simulations to empirically examine the effect of the cutoff, where $\lceil a \rceil$ denotes the integer part of a .

Example 1. This example is designed to compare the finite sample performance of the DC-SIS with the SIS (Fan and Lv 2008) and the SIRS (Zhu et al. 2011). In this example, we generate the response from the following four models:

- (1.a): $Y = c_1\beta_1 X_1 + c_2\beta_2 X_2 + c_3\beta_3 \mathbf{1}(X_{12} < 0) + c_4\beta_4 X_{22} + \varepsilon,$
- (1.b): $Y = c_1\beta_1 X_1 X_2 + c_3\beta_2 \mathbf{1}(X_{12} < 0) + c_4\beta_3 X_{22} + \varepsilon,$
- (1.c): $Y = c_1\beta_1 X_1 X_2 + c_3\beta_2 \mathbf{1}(X_{12} < 0) X_{22} + \varepsilon,$
- (1.d): $Y = c_1\beta_1 X_1 + c_2\beta_2 X_2 + c_3\beta_3 \mathbf{1}(X_{12} < 0) + \exp(c_4 |X_{22}|)\varepsilon,$

where $\mathbf{1}(X_{12} < 0)$ is an indicator function.

The regression functions $E(Y | \mathbf{x})$ in models (1.a)–(1.d) are all nonlinear in X_{12} . In addition, models (1.b) and (1.c) contain an interaction term $X_1 X_2$, and model (1.d) is heteroscedastic. Following Fan and Lv (2008), we choose $\beta_j = (-1)^U (a + |Z|)$ for $j = 1, 2, 3$, and 4, where $a = 4 \log n / \sqrt{n}$, $U \sim \text{Bernoulli}(0.4)$ and $Z \sim \mathcal{N}(0, 1)$. We set $(c_1, c_2, c_3, c_4) = (2, 0.5, 3, 2)$ in this example to challenge the feature screening procedures under consideration. For each independence screening procedure, we compute the associated marginal utility between each predictor X_k and the response Y . That is, we regard $\mathbf{x} = (X_1, \dots, X_p)^T \in \mathbb{R}^p$ as the predictor vector in this example.

Tables 1 and 2 present the simulation results for \mathcal{S} , \mathcal{P}_s , and \mathcal{P}_a . The performances of the DC-SIS, SIS, and SIRS are quite similar in model (1.a), indicating that the SIS has a robust performance if the working linear model does not deviate far from the underlying true model. The DC-SIS outperforms the SIS and the SIRS significantly in models (1.b)–(1.d). Both the SIS and the SIRS have little chance to identify the important predictors X_1 and X_2 in models (1.b) and (1.c), and X_{22} in model (1.d).

Example 2. We illustrate that the DC-SIS can be directly used for screening grouped predictors. In many regression problems, some predictors can be naturally grouped. The most common example that contains group variables is the multifactor analysis of variance (ANOVA) problem, in which each factor may have several levels and can be expressed through a group of dummy variables. The goal of ANOVA is to select important main effects and interactions for accurate predictions, which amounts to the selection of groups of dummy variables. To demonstrate the practicability of the DC-SIS, we adopt the following model:

$$Y = c_1\beta_1 X_1 + c_2\beta_2 X_2 + c_3\beta_3 \{\mathbf{1}(X_{12} < q_1) + 1.5 \times \mathbf{1}(q_1 \leq X_{12} < q_2) + 2 \times \mathbf{1}(X_{12} \geq q_3)\} + c_4\beta_4 X_{22} + \varepsilon,$$

Table 1. The 5%, 25%, 50%, 75%, and 95% quantiles of the minimum model size S out of 500 replications in Example 1

S	SIS					SIRS					DC-SIS				
	5%	25%	50%	75%	95%	5%	25%	50%	75%	95%	5%	25%	50%	75%	95%
Case 1: $p = 2000$ and $\sigma_{ij} = 0.5^{ i-j }$															
(1.a)	4.0	4.0	5.0	7.0	21.2	4.0	4.0	5.0	7.0	45.1	4.0	4.0	4.0	6.0	18.0
(1.b)	68.0	578.5	1180.5	1634.5	1938.0	232.9	871.5	1386.0	1725.2	1942.4	5.0	9.0	24.5	73.0	345.1
(1.c)	395.9	1037.2	1438.0	1745.0	1945.1	238.5	805.0	1320.0	1697.0	1946.0	6.0	10.0	22.0	59.0	324.1
(1.d)	130.5	611.2	1166.0	1637.0	1936.5	42.0	304.2	797.0	1432.2	1846.1	4.0	5.0	9.0	41.0	336.2
Case 2: $p = 2000$ and $\sigma_{ij} = 0.8^{ i-j }$															
(1.a)	5.0	9.0	16.0	97.0	729.4	5.0	9.0	18.0	112.8	957.1	4.0	7.0	11.0	31.2	507.2
(1.b)	26.0	283.2	852.0	1541.2	1919.0	103.9	603.0	1174.0	1699.2	1968.0	5.0	8.0	11.0	17.0	98.0
(1.c)	224.5	775.2	1249.5	1670.0	1951.1	118.6	573.2	1201.5	1685.2	1955.0	7.0	10.0	15.0	38.0	198.3
(1.d)	79.0	583.8	1107.5	1626.2	1930.0	50.9	300.5	728.0	1368.2	1900.1	4.0	7.0	17.0	73.2	653.1
Case 3: $p = 5000$ and $\sigma_{ij} = 0.5^{ i-j }$															
(1.a)	4.0	4.0	5.0	6.0	59.0	4.0	4.0	5.0	7.0	88.4	4.0	4.0	4.0	6.0	34.1
(1.b)	165.1	1112.5	2729.0	3997.2	4851.5	560.8	1913.0	3249.0	4329.0	4869.1	5.0	11.8	45.0	168.8	956.7
(1.c)	1183.7	2712.0	3604.5	4380.2	4885.0	440.4	1949.0	3205.5	4242.8	4883.1	7.0	17.0	53.0	179.5	732.0
(1.d)	259.9	1338.5	2808.5	3990.8	4764.9	118.7	823.2	1833.5	3314.5	4706.1	4.0	5.0	15.0	77.2	848.2
Case 4: $p = 5000$ and $\sigma_{ij} = 0.8^{ i-j }$															
(1.a)	5.0	10.0	26.5	251.5	2522.7	5.0	10.0	28.0	324.8	3246.4	5.0	8.0	14.0	69.0	1455.1
(1.b)	40.7	639.8	2072.0	3803.8	4801.7	215.7	1677.8	3010.0	4352.2	4934.1	5.0	8.0	11.0	21.0	162.0
(1.c)	479.2	1884.8	3347.5	4298.5	4875.2	297.7	1359.2	2738.5	4072.5	4877.6	8.0	12.0	22.0	83.0	657.9
(1.d)	307.0	1544.0	2832.5	4026.2	4785.2	148.2	672.0	1874.0	3330.0	4665.2	4.0	7.0	21.0	165.2	1330.0

where $q_1, q_2,$ and q_3 are the 25%, 50%, and 75% quantiles of X_{12} , respectively. The variables X with the coefficients c_i 's and β_i 's are the same as those in Example 1. We write

$$\tilde{\mathbf{x}}_{12} = \{\mathbf{1}(X_{12} < q_1), \mathbf{1}(q_1 \leq X_{12} < q_2), \mathbf{1}(q_2 \leq X_{12} < q_3)\}^T$$

These three variables naturally become a group. The predictor vector in this example becomes $\mathbf{x} = (X_1, \dots, X_{11}, \tilde{\mathbf{x}}_{12}, X_{13}, \dots, X_p)^T \in \mathbb{R}^{p+2}$. We remark here that the marginal utility of the grouped variable $\tilde{\mathbf{x}}_{12}$ is defined by

$$\widehat{\omega}_{12} = \widehat{\text{dcorr}}^2(\tilde{\mathbf{x}}_{12}, Y).$$

The 5%, 25%, 50%, 75%, and 95% percentiles of the minimum model size S are summarized in Table 3. These percentiles indicate that with very high probability, the minimum model size S to ensure the inclusion of all active predictors is small. Note that $\lceil n/\log(n) \rceil = 37$. Thus, almost all \mathcal{P}_s and \mathcal{P}_a s of the DC-SIS equal 100%. All active predictors, including the grouped variable $\tilde{\mathbf{x}}_{12}$, can almost perfectly be selected into the resulting model across all three different model sizes. Hence, the DC-SIS is efficient to select the grouped predictors.

Example 3. In this example, we investigate the performance of the DC-SIS with multivariate responses. The SIS proposed by Fan and Lv (2008) cannot be directly applied for such settings. In contrast, the DC-SIS is ready for screening the active predictors by the nature of DC. In this example, we generate $\mathbf{y} = (Y_1, Y_2)^T$ from normal distribution with mean zero and covariance matrix $\Sigma_{\mathbf{y}|\mathbf{x}} = (\sigma_{x,ij})_{2 \times 2}$, where $\sigma_{x,11} = \sigma_{x,22} = 1$ and $\sigma_{x,12} = \sigma_{x,21} = \sigma(\mathbf{x})$. We consider two scenarios for the correlation function $\sigma(\mathbf{x})$:

(3.a):

$$\sigma(\mathbf{x}) = \sin(\boldsymbol{\beta}_1^T \mathbf{x}),$$

where $\boldsymbol{\beta}_1 = (0.8, 0.6, 0, \dots, 0)^T$.

(3.b):

$$\sigma(\mathbf{x}) = \frac{\{\exp(\boldsymbol{\beta}_2^T \mathbf{x}) - 1\}}{\{\exp(\boldsymbol{\beta}_2^T \mathbf{x}) + 1\}},$$

where $\boldsymbol{\beta}_2 = (2 - U_1, 2 - U_2, 2 - U_3, 2 - U_4, 0, \dots, 0)^T$

with U_i 's being independent and identically distributed (iid) according to uniform distribution Uniform[0, 1].

Tables 4 and 5 present the simulation results. Table 4 implies that the DC-SIS performs reasonably well for both models (3.a) and (3.b) in terms of model complexity. Table 5 indicates that the proportions that the active predictors are selected into the model are close to one, which supports the assertion that the DC-SIS processes the sure screening property. It implies that the DC-SIS can identify the active predictors contained in correlations between multivariate responses. This may be potentially useful in gene coexpression analysis.

Example 4. The cardiomyopathy microarray dataset was once analyzed by Segal, Dahlquist, and Conklin (2003) and Hall and Miller (2009). The goal is to identify the most influential genes for overexpression of a G protein-coupled receptor (Ro1) in mice. The response Y is the Ro1 expression level, and the predictors X_k 's are other gene expression levels. Compared with the sample size $n = 30$ in this dataset, the dimension $p = 6319$ is very large.

The DC-SIS procedure ranks two genes, labeled as Msa.2134.0 and Msa.2877.0, at the top. The scatterplots of

Table 2. The proportions of \mathcal{P}_s and \mathcal{P}_a in Example 1. The user-specified model sizes are $d_1 = \lceil n/\log n \rceil$, $d_2 = 2\lceil n/\log n \rceil$, and $d_3 = 3\lceil n/\log n \rceil$

Model	Size	SIS					SIRS					DC-SIS				
		\mathcal{P}_s				\mathcal{P}_a	\mathcal{P}_s				\mathcal{P}_a	\mathcal{P}_s				\mathcal{P}_a
		X_1	X_2	X_{12}	X_{22}	All	X_1	X_2	X_{12}	X_{22}	All	X_1	X_2	X_{12}	X_{22}	All
Case 1: $p = 2000$ and $\sigma_{ij} = 0.5^{i-j}$																
(1.a)	d_1	1.00	1.00	0.96	1.00	0.96	1.00	1.00	0.95	1.00	0.94	1.00	1.00	0.97	1.00	0.96
	d_2	1.00	1.00	0.98	1.00	0.97	1.00	1.00	0.96	1.00	0.96	1.00	1.00	0.98	1.00	0.98
	d_3	1.00	1.00	0.98	1.00	0.98	1.00	1.00	0.97	1.00	0.97	1.00	1.00	0.99	1.00	0.98
(1.b)	d_1	0.08	0.07	0.97	1.00	0.03	0.02	0.03	0.98	1.00	0.00	0.72	0.70	0.99	1.00	0.58
	d_2	0.12	0.13	0.98	1.00	0.06	0.05	0.05	0.99	1.00	0.01	0.85	0.84	1.00	1.00	0.76
	d_3	0.15	0.17	0.99	1.00	0.07	0.06	0.06	0.99	1.00	0.01	0.89	0.88	1.00	1.00	0.82
(1.c)	d_1	0.12	0.13	0.01	0.99	0.00	0.04	0.03	0.51	1.00	0.01	0.93	0.93	0.77	1.00	0.65
	d_2	0.17	0.18	0.03	0.99	0.00	0.07	0.05	0.67	1.00	0.01	0.97	0.96	0.84	1.00	0.79
	d_3	0.21	0.21	0.05	0.99	0.00	0.09	0.08	0.75	1.00	0.02	0.98	0.97	0.89	1.00	0.84
(1.d)	d_1	0.42	0.22	0.14	0.42	0.02	1.00	0.98	0.87	0.05	0.04	1.00	0.91	0.81	0.99	0.73
	d_2	0.48	0.29	0.22	0.50	0.03	1.00	0.99	0.91	0.10	0.09	1.00	0.94	0.87	1.00	0.82
	d_3	0.56	0.32	0.26	0.54	0.04	1.00	0.99	0.93	0.12	0.11	1.00	0.96	0.92	1.00	0.88
Case 2: $p = 2000$ and $\sigma_{ij} = 0.8^{i-j}$																
(1.a)	d_1	1.00	1.00	0.63	1.00	0.63	1.00	1.00	0.62	1.00	0.62	1.00	1.00	0.78	1.00	0.77
	d_2	1.00	1.00	0.71	1.00	0.72	1.00	1.00	0.70	1.00	0.69	1.00	1.00	0.84	1.00	0.84
	d_3	1.00	1.00	0.77	1.00	0.78	1.00	1.00	0.75	1.00	0.75	1.00	1.00	0.86	1.00	0.86
(1.b)	d_1	0.12	0.13	0.81	1.00	0.06	0.04	0.04	0.88	1.00	0.02	0.97	0.98	0.92	1.00	0.88
	d_2	0.19	0.19	0.86	1.00	0.12	0.07	0.07	0.91	1.00	0.03	0.99	0.99	0.95	1.00	0.94
	d_3	0.22	0.23	0.88	1.00	0.15	0.09	0.11	0.93	1.00	0.06	1.00	0.99	0.96	1.00	0.96
(1.c)	d_1	0.17	0.16	0.03	0.99	0.00	0.04	0.04	0.53	1.00	0.02	1.00	1.00	0.75	1.00	0.75
	d_2	0.22	0.22	0.06	1.00	0.01	0.08	0.08	0.71	1.00	0.03	1.00	1.00	0.85	1.00	0.86
	d_3	0.27	0.27	0.10	1.00	0.03	0.10	0.10	0.81	1.00	0.05	1.00	1.00	0.90	1.00	0.90
(1.d)	d_1	0.44	0.38	0.11	0.45	0.03	1.00	1.00	0.73	0.05	0.04	0.99	0.98	0.68	1.00	0.67
	d_2	0.51	0.46	0.18	0.53	0.05	1.00	1.00	0.81	0.09	0.08	1.00	0.98	0.76	1.00	0.75
	d_3	0.55	0.49	0.22	0.57	0.06	1.00	1.00	0.84	0.14	0.11	1.00	0.99	0.80	1.00	0.80
Case 3: $p = 5000$ and $\sigma_{ij} = 0.5^{i-j}$																
(1.a)	d_1	1.00	1.00	0.94	1.00	0.94	1.00	0.99	0.92	1.00	0.92	1.00	0.99	0.96	1.00	0.95
	d_2	1.00	1.00	0.95	1.00	0.95	1.00	1.00	0.95	1.00	0.95	1.00	1.00	0.97	1.00	0.97
	d_3	1.00	1.00	0.96	1.00	0.96	1.00	1.00	0.96	1.00	0.96	1.00	1.00	0.98	1.00	0.98
(1.b)	d_1	0.06	0.06	0.94	1.00	0.02	0.02	0.02	0.96	1.00	0.00	0.59	0.60	0.98	1.00	0.46
	d_2	0.09	0.09	0.96	1.00	0.03	0.03	0.03	0.97	1.00	0.01	0.72	0.72	0.99	1.00	0.61
	d_3	0.12	0.10	0.97	1.00	0.04	0.05	0.04	0.98	1.00	0.01	0.79	0.78	0.99	1.00	0.68
(1.c)	d_1	0.06	0.06	0.01	0.99	0.00	0.03	0.02	0.30	1.00	0.00	0.86	0.87	0.61	1.00	0.41
	d_2	0.10	0.10	0.02	1.00	0.00	0.04	0.03	0.45	1.00	0.00	0.92	0.93	0.69	1.00	0.57
	d_3	0.12	0.12	0.02	1.00	0.00	0.05	0.05	0.53	1.00	0.00	0.94	0.95	0.73	1.00	0.64
(1.d)	d_1	0.39	0.21	0.11	0.40	0.01	1.00	0.97	0.82	0.02	0.02	0.99	0.87	0.74	0.99	0.65
	d_2	0.44	0.24	0.14	0.45	0.01	1.00	0.98	0.88	0.04	0.03	0.99	0.90	0.81	0.99	0.75
	d_3	0.48	0.28	0.17	0.47	0.02	1.00	0.99	0.90	0.06	0.05	0.99	0.92	0.85	1.00	0.79
Case 4: $p = 5000$ and $\sigma_{ij} = 0.8^{i-j}$																
(1.a)	d_1	1.00	1.00	0.55	1.00	0.55	1.00	1.00	0.55	1.00	0.55	1.00	1.00	0.70	1.00	0.69
	d_2	1.00	1.00	0.61	1.00	0.62	1.00	1.00	0.61	1.00	0.61	1.00	1.00	0.76	1.00	0.76
	d_3	1.00	1.00	0.67	1.00	0.67	1.00	1.00	0.64	1.00	0.64	1.00	1.00	0.80	1.00	0.80
(1.b)	d_1	0.10	0.09	0.74	1.00	0.05	0.02	0.02	0.83	1.00	0.00	0.94	0.94	0.90	1.00	0.82
	d_2	0.12	0.13	0.81	1.00	0.07	0.03	0.04	0.87	1.00	0.01	0.97	0.97	0.93	1.00	0.89
	d_3	0.15	0.16	0.84	1.00	0.10	0.05	0.06	0.90	1.00	0.02	0.98	0.98	0.95	1.00	0.92
(1.c)	d_1	0.10	0.10	0.02	0.98	0.00	0.02	0.03	0.34	1.00	0.00	1.00	1.00	0.64	1.00	0.63
	d_2	0.13	0.14	0.04	0.99	0.01	0.04	0.04	0.50	1.00	0.01	1.00	1.00	0.74	1.00	0.74
	d_3	0.16	0.18	0.05	0.99	0.01	0.05	0.05	0.61	1.00	0.02	1.00	1.00	0.79	1.00	0.79
(1.d)	d_1	0.42	0.32	0.09	0.40	0.01	1.00	1.00	0.66	0.02	0.01	0.99	0.97	0.63	0.98	0.59
	d_2	0.48	0.39	0.12	0.44	0.02	1.00	1.00	0.74	0.04	0.03	0.99	0.97	0.70	1.00	0.68
	d_3	0.51	0.42	0.15	0.46	0.02	1.00	1.00	0.78	0.05	0.04	0.99	0.98	0.73	1.00	0.71

Table 3. The 5%, 25%, 50%, 75%, and 95% quantiles of the minimum model size S out of 500 replications in Example 2

S	$p = 2000$					$p = 5000$				
	5%	25%	50%	75%	95%	5%	25%	50%	75%	95%
$\sigma_{ij} = 0.5^{ i-j }$	4.0	4.0	4.0	5.0	12.0	4.0	4.0	4.0	6.0	16.1
$\sigma_{ij} = 0.8^{ i-j }$	4.0	5.0	7.0	9.0	15.2	4.0	5.0	7.0	9.0	21.0

Table 4. The 5%, 25%, 50%, 75%, and 95% quantiles of the minimum model size S out of 500 replications in Example 3

S	Model	$p = 2000$					$p = 5000$				
		5%	25%	50%	75%	95%	5%	25%	50%	75%	95%
$\sigma_{ij} = 0.5^{ i-j }$	(3.a)	4.0	9.0	18.0	39.3	112.3	6.0	22.0	48.0	95.3	296.4
	(3.b)	6.0	19.0	43.0	92.0	253.1	14.0	45.0	92.5	198.8	571.6
$\sigma_{ij} = 0.8^{ i-j }$	(3.a)	2.0	3.0	6.0	12.0	40.0	2.0	6.0	14.0	32.0	98.0
	(3.b)	4.0	4.0	4.0	6.0	10.0	4.0	4.0	5.0	8.0	18.1

Table 5. The proportions of \mathcal{P}_s and \mathcal{P}_a in Example 3. The user-specified model sizes are $d_1 = \lceil n / \log n \rceil$, $d_2 = 2\lceil n / \log n \rceil$, and $d_3 = 3\lceil n / \log n \rceil$

Size		$p = 2000$								$p = 5000$							
		(3.a)			(3.b)					(3.a)			(3.b)				
		\mathcal{P}_s		\mathcal{P}_a	\mathcal{P}_s		\mathcal{P}_a			\mathcal{P}_s		\mathcal{P}_a	\mathcal{P}_s			\mathcal{P}_a	
		X_1	X_2	All	X_1	X_2	X_3	X_4	All	X_1	X_2	All	X_1	X_2	X_3	X_4	All
$\sigma_{ij} = 0.5^{ i-j }$	d_1	0.95	0.76	0.74	0.71	0.98	0.98	0.72	0.47	0.79	0.49	0.42	0.48	0.91	0.90	0.53	0.20
	d_2	0.98	0.90	0.90	0.85	0.99	0.99	0.85	0.71	0.93	0.70	0.67	0.67	0.97	0.97	0.71	0.45
	d_3	1.00	0.95	0.95	0.91	0.99	1.00	0.90	0.81	0.97	0.81	0.80	0.75	0.98	0.99	0.78	0.55
$\sigma_{ij} = 0.8^{ i-j }$	d_1	0.98	0.95	0.94	1.00	1.00	1.00	1.00	1.00	0.92	0.84	0.81	1.00	1.00	1.00	0.99	0.99
	d_2	1.00	0.98	0.99	1.00	1.00	1.00	1.00	1.00	0.98	0.95	0.93	1.00	1.00	1.00	1.00	1.00
	d_3	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	0.99	0.96	0.96	1.00	1.00	1.00	1.00	1.00

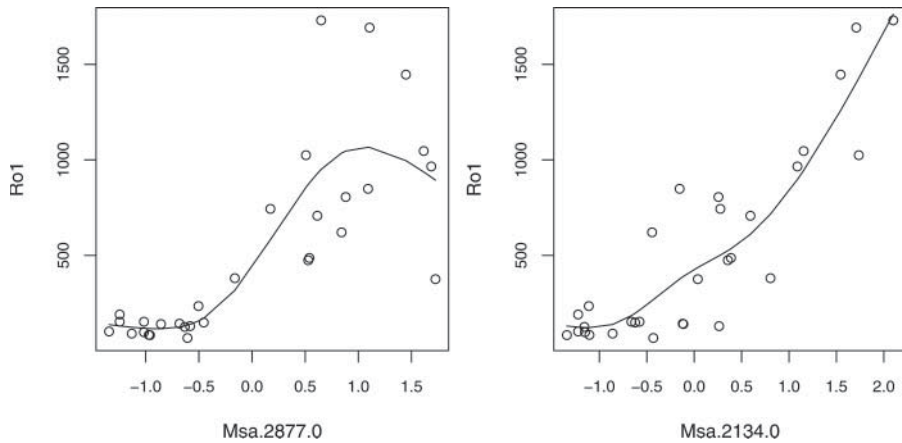


Figure 1. The scatterplot of Y versus two gene expression levels identified by the DC-SIS.

Y versus these two gene expression levels with cubic spline fit curves in Figure 1 indicate clearly the existence of nonlinear patterns. Yet, our finding is different from Hall and Miller (2009) in that they ranked Msa.2877.0 and Msa.1166.0 at the top, with their proposed generalized correlation ranking. A natural question arises: which screening procedure performs better in terms of ranking? To compare the performance of these two procedures, we fit an additive model as follows:

$$Y = \ell_{k1}(X_{k1}) + \ell_{k2}(X_{k2}) + \varepsilon_k, \text{ for } k = 1, 2.$$

The DC-SIS, corresponding to $k = 1$, regards Msa.2134.0 and Msa.2877.0 as the two predictors, while the generalized correlation ranking proposed by Hall and Miller (2009), corresponding to $k = 2$, regards Msa.2877.0 and Msa.1166.0 as predictors in the above model. We fit the unknown link functions ℓ_{ki} using the R “mgcv” package. The DC-SIS method clearly achieves better performance with the adjusted R^2 of 96.8% and the deviance explained of 98.3%, in contrast to the adjusted R^2 of 84.5% and the deviance explained of 86.6% for the generalized correlation ranking method. We remark here that deviance explained means the proportion of the null deviance explained by the proposed model, with a larger value indicating better performance. Because both the adjusted R^2 values and the explained deviance are very large, it seems unnecessary to extract any additional genes.

4. DISCUSSION

In this article, we propose an SIS procedure using DC, that is, DC-SIS. We establish the sure screening property for this procedure when the number of predictors diverges with an exponential rate of the sample size. We examine the finite sample performance of the proposed procedure via Monte Carlo studies and illustrate the proposed methodology through a real-data example. We follow Fan and Lv (2008) to set the cutoff d in this article and examine the effect of different values of d . As pointed out by a referee, the choice of d is very important at the screening stage. Zhao and Li (2012) proposed an approach to selecting d for Cox models based on controlling false positive rate. Their approach is merely for model-based feature screening methods. Zhu et al. (2011) proposed an alternate method to determine d for the SIRS. One may adopt their procedure for the DC-SIS. We opt not to pursue this further. Certainly, the selection of d is similar to the selection of the tuning parameter in regularization methods, and plays an important role in practical implementation. This is a good topic for future research.

Similar to the SIS, the DC-SIS may fail to identify some important predictors that are marginally independent of the response. Thus, it is of interest to develop an iterative procedure to fix such an issue. In the earlier version of this article, we proposed an iterative version of DC-SIS. Our empirical studies including Monte Carlo simulation and real-data analysis imply that the proposed iterative DC-SIS may be used to fix the problem in a similar spirit of iterative SIS (ISIS; Fan and Lv 2008). A theoretical analysis of the iterative DC-SIS needs further study. New methods to deal with the identification of important predictors that are marginally independent of the response is an important topic for future research.

APPENDIX A: SOME LEMMAS

Lemmas 1 and 2 will be used repeatedly in the proof of Theorem 1. These two lemmas provide us two exponential inequalities, and are extracted from lemma 5.6.1.A and theorem 5.6.1.A of Serfling (1980, pp. 200–201).

Lemma 1. Let $\mu = E(Y)$. If $\Pr(a \leq Y \leq b) = 1$, then

$$E[\exp\{s(Y - \mu)\}] \leq \exp\{s^2(b - a)^2/8\}, \text{ for any } s > 0.$$

Lemma 2. Let $h(Y_1, \dots, Y_m)$ be a kernel of the U statistics U_n , and $\theta = E\{h(Y_1, \dots, Y_m)\}$. If $a \leq h(Y_1, \dots, Y_m) \leq b$, then, for any $t > 0$ and $n \geq m$,

$$\Pr(U_n - \theta \geq t) \leq \exp\{-2[n/m]t^2/(b - a)^2\},$$

where $[n/m]$ denotes the integer part of n/m .

Due to the symmetry of U statistics, Lemma 2 entails that

$$\Pr(|U_n - \theta| \geq t) \leq 2 \exp\{-2[n/m]t^2/(b - a)^2\}.$$

Let us introduce some notations before giving the proof of Theorem 1. Let $\{\tilde{X}_k, \tilde{\mathbf{y}}\}$ be an independent copy of $\{X_k, \mathbf{y}\}$, and define $S_{k1} = E\|X_k - \tilde{X}_k\|_1 \|\mathbf{y} - \tilde{\mathbf{y}}\|_q$, $S_{k2} = E\|X_k - \tilde{X}_k\|_1 E\|\mathbf{y} - \tilde{\mathbf{y}}\|_q$, and $S_{k3} = E\{E(\|X_k - \tilde{X}_k\|_1 | X_k) E(\|\mathbf{y} - \tilde{\mathbf{y}}\|_q | \mathbf{y})\}$, and their sample counterparts

$$\begin{aligned} \hat{S}_{k1} &= \frac{1}{n^2} \sum_{i,j=1}^n \|X_{ik} - X_{jk}\|_1 \|\mathbf{y}_i - \mathbf{y}_j\|_q, \\ \hat{S}_{k2} &= \frac{1}{n^2} \sum_{i,j=1}^n \|X_{ik} - X_{jk}\|_1 \frac{1}{n^2} \sum_{i,j=1}^n \|\mathbf{y}_i - \mathbf{y}_j\|_q, \\ \hat{S}_{k3} &= \frac{1}{n^3} \sum_{i,j,l=1}^n \|X_{ik} - X_{lk}\|_1 \|\mathbf{y}_j - \mathbf{y}_l\|_q. \end{aligned}$$

By definitions of distance covariance and sample distance covariance, it follows that

$$\begin{aligned} \text{dcov}^2(X_k, \mathbf{y}) &= S_{k1} + S_{k2} - 2S_{k3} \quad \text{and} \\ \widehat{\text{dcov}}^2(X_k, \mathbf{y}) &= \hat{S}_{k1} + \hat{S}_{k2} - 2\hat{S}_{k3}. \end{aligned}$$

APPENDIX B: PROOF OF THEOREM 1

We aim to show the uniform consistency of the denominator and the numerator of $\hat{\omega}_k$ under regularity conditions respectively. Because the denominator of $\hat{\omega}_k$ has a similar form as the numerator, we deal with its numerator only below. Throughout the proof, the notations C and c are generic constants, which may take different values at each appearance.

We first deal with \hat{S}_{k1} . Define $\hat{S}_{k1}^* = \{n(n-1)\}^{-1} \sum_{i \neq j} \|X_{ik} - X_{jk}\|_1 \|\mathbf{y}_i - \mathbf{y}_j\|_q$, which is a usual U statistics. We shall establish the uniform consistency of \hat{S}_{k1}^* by using the theory of U statistics (Serfling 1980, sec. 5). By using the Cauchy–Schwartz inequality,

$$\begin{aligned} S_{k1} &= E(\|X_{ik} - X_{jk}\|_1 \|\mathbf{y}_i - \mathbf{y}_j\|_q) \\ &\leq \{E(\|X_{ik} - X_{jk}\|_1^2) E(\|\mathbf{y}_i - \mathbf{y}_j\|_q^2)\}^{1/2} \\ &\leq 4\{E(X_k^2) E\|\mathbf{y}\|_q^2\}^{1/2}. \end{aligned}$$

This, together with Condition (C1), implies that S_{k1} is uniformly bounded in p , that is, $\sup_{1 \leq k \leq p} S_{k1} < \infty$. For any given $\varepsilon > 0$, take n large enough such that $S_{k1}/n < \varepsilon$. Then, it can be easily shown that

$$\begin{aligned} &\Pr(|\hat{S}_{k1} - S_{k1}| \geq 2\varepsilon) \\ &= \Pr\{|\hat{S}_{k1}^* (n-1)/n - S_{k1} (n-1)/n - S_{k1}/n| \geq 2\varepsilon\} \\ &\leq \Pr\{|\hat{S}_{k1}^* - S_{k1}| (n-1)/n \geq 2\varepsilon - S_{k1}/n\} \\ &\leq \Pr(|\hat{S}_{k1}^* - S_{k1}| \geq \varepsilon). \end{aligned} \tag{B.1}$$

To establish the uniform consistency of \widehat{S}_{k1} , it thus suffices to show the uniform consistency of \widehat{S}_{k1}^* . Let $h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) = \|X_{ik} - X_{jk}\|_1 \|\mathbf{y}_i - \mathbf{y}_j\|_q$ be the kernel of the U statistics \widehat{S}_{k1}^* . We decompose the kernel function h_1 into two parts: $h_1 = h_1 \mathbf{1}(h_1 > M) + h_1 \mathbf{1}(h_1 \leq M)$, where M will be specified later. The U statistics can now be written as follows:

$$\begin{aligned} \widehat{S}_{k1}^* &= \{n(n-1)\}^{-1} \sum_{i \neq j} h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) \\ &\quad \times \mathbf{1}\{h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) \leq M\} \\ &\quad + \{n(n-1)\}^{-1} \sum_{i \neq j} h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) \\ &\quad \times \mathbf{1}\{h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) > M\} \\ &= \widehat{S}_{k1,1}^* + \widehat{S}_{k1,2}^*. \end{aligned}$$

Accordingly, we decompose S_{k1} into two parts:

$$\begin{aligned} S_{k1} &= E[h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) \mathbf{1}\{h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) \leq M\}] \\ &\quad + E[h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) \mathbf{1}\{h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) > M\}] \\ &= S_{k1,1} + S_{k1,2}. \end{aligned}$$

Clearly, $\widehat{S}_{k1,1}^*$ and $\widehat{S}_{k1,2}^*$ are unbiased estimators of $S_{k1,1}$ and $S_{k1,2}$, respectively.

We deal with the consistency of $\widehat{S}_{k1,1}^*$ first. With the Markov's inequality, for any $t > 0$, we can obtain that

$$\Pr(\widehat{S}_{k1,1}^* - S_{k1,1} \geq \varepsilon) \leq \exp(-t\varepsilon) \exp(-tS_{k1,1}) E\{\exp(t\widehat{S}_{k1,1}^*)\}.$$

Serfling (1980, sec. 5.1.6) showed that any U statistics can be represented as an average of averages of iid random variables. That is, $\widehat{S}_{k1,1}^* = (n!)^{-1} \sum_{n!} \Omega_1(X_{1k}, \mathbf{y}_1; \dots; X_{nk}, \mathbf{y}_n)$, where $\sum_{n!}$ denotes the summation over all possible permutations of $(1, \dots, n)$, and each $\Omega_1(X_{1k}, \mathbf{y}_1; \dots; X_{nk}, \mathbf{y}_n)$ is an average of $m = [n/2]$ iid random variables (i.e., $\Omega_1 = m^{-1} \sum_r h_1^{(r)} \mathbf{1}\{h_1^{(r)} \leq M\}$). Since the exponential function is convex, it follows from Jensen's inequality that, for $0 < t \leq 2s_0$,

$$\begin{aligned} E\{\exp(t\widehat{S}_{k1,1}^*)\} &= E\left[\exp\left\{t(n!)^{-1} \sum_{n!} \Omega_1(X_{1k}, \mathbf{y}_1; \dots; X_{nk}, \mathbf{y}_n)\right\}\right] \\ &\leq (n!)^{-1} \sum_{n!} E[\exp\{t\Omega_1(X_{1k}, \mathbf{y}_1; \dots; X_{nk}, \mathbf{y}_n)\}] \\ &= E^m\{\exp(m^{-1}t h_1^{(r)} \mathbf{1}\{h_1^{(r)} \leq M\})\}, \end{aligned}$$

which, together with Lemma 1, entails immediately that

$$\begin{aligned} \Pr(\widehat{S}_{k1,1}^* - S_{k1,1} \geq \varepsilon) &\leq \exp(-t\varepsilon) E^m\left\{\exp\left(m^{-1}t[h_1^{(r)} \mathbf{1}\{h_1^{(r)} \leq M\} - S_{k1,1}]\right)\right\} \\ &\leq \exp\{-t\varepsilon + M^2 t^2 / (8m)\}. \end{aligned}$$

By choosing $t = 4\varepsilon m / M^2$, we have $\Pr(\widehat{S}_{k1,1}^* - S_{k1,1} \geq \varepsilon) \leq \exp(-2\varepsilon^2 m / M^2)$. Therefore, by the symmetry of U statistics, we can obtain easily that

$$\Pr(|\widehat{S}_{k1,1}^* - S_{k1,1}| \geq \varepsilon) \leq 2 \exp(-2\varepsilon^2 m / M^2). \quad (\text{B.2})$$

Next we show the consistency of $\widehat{S}_{k1,2}^*$. With Cauchy-Schwartz and Markov's inequalities,

$$\begin{aligned} S_{k1,2}^2 &\leq E\{h_1^2(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j)\} \Pr\{h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) > M\} \\ &\leq E\{h_1^2(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j)\} E[\exp\{s'h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j)\}] / \exp(s'M), \end{aligned}$$

for any $s' > 0$. Using the fact $(a^2 + b^2)/2 \geq (a + b)^2/4 \geq |ab|$, we have

$$\begin{aligned} h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) &= \{(X_{ik} - X_{jk})^2 (\mathbf{y}_i - \mathbf{y}_j)^T (\mathbf{y}_i - \mathbf{y}_j)\}^{1/2} \\ &\leq 2\{(X_{ik}^2 + X_{jk}^2) (\|\mathbf{y}_i\|_q^2 + \|\mathbf{y}_j\|_q^2)\}^{1/2} \\ &\leq \{(X_{ik}^2 + X_{jk}^2 + \|\mathbf{y}_i\|_q^2 + \|\mathbf{y}_j\|_q^2)\}^{1/2} \\ &= X_{ik}^2 + X_{jk}^2 + \|\mathbf{y}_i\|_q^2 + \|\mathbf{y}_j\|_q^2, \end{aligned}$$

which yields that

$$\begin{aligned} E[\exp\{s'h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j)\}] &\leq E\{\exp\{s'(X_{ik}^2 + X_{jk}^2 + \|\mathbf{y}_i\|_q^2 + \|\mathbf{y}_j\|_q^2)\}\} \\ &\leq E\{\exp(2s'X_{ik}^2)\} E\{\exp(2s'\|\mathbf{y}_i\|_q^2)\}. \end{aligned}$$

The last inequality follows from the Cauchy-Schwartz inequality. If we choose $M = cn^\gamma$ for $0 < \gamma < 1/2 - \kappa$, then $S_{k1,2} \leq \varepsilon/2$ when n is sufficiently large. Consequently,

$$\Pr(|\widehat{S}_{k1,2}^* - S_{k1,2}| > \varepsilon) \leq \Pr(|\widehat{S}_{k1,2}^*| > \varepsilon/2). \quad (\text{B.3})$$

It remains to bound the probability $\Pr(|\widehat{S}_{k1,2}^*| > \varepsilon/2)$. We observe that the events satisfy

$$\{|\widehat{S}_{k1,2}^*| > \varepsilon/2\} \subseteq \{X_{ik}^2 + \|\mathbf{y}_i\|_q^2 > M/2, \text{ for some } 1 \leq i \leq p\}. \quad (\text{B.4})$$

To see this, we assume that $X_{ik}^2 + \|\mathbf{y}_i\|_q^2 \leq M/2$, for all $1 \leq i \leq p$. This assumption will lead to a contradiction. To be precise, under this assumption, $h_1(X_{ik}, \mathbf{y}_i; X_{jk}, \mathbf{y}_j) \leq X_{ik}^2 + X_{jk}^2 + \|\mathbf{y}_i\|_q^2 + \|\mathbf{y}_j\|_q^2 \leq M$. Consequently, $|\widehat{S}_{k1,2}^*| = 0$, which is a contrary to the event $|\widehat{S}_{k1,2}^*| > \varepsilon/2$. This verifies that the relation (B.4) is true.

By invoking Condition (C1), there must exist a constant C such that

$$\begin{aligned} \Pr(\|X_k\|_1^2 + \|\mathbf{y}_q\|_q^2 \geq M/2) &\leq \Pr(\|X_k\|_1 \geq \sqrt{M}/2) + \Pr(\|\mathbf{y}_q\|_q \geq \sqrt{M}/2) \leq 2C \exp(-sM/4). \end{aligned}$$

The last inequality follows from Markov's inequality for $s > 0$. Consequently,

$$\begin{aligned} \max_{1 \leq k \leq p} \Pr(|\widehat{S}_{k1,2}^*| > \varepsilon/2) &\leq n \max_{1 \leq k \leq p} \Pr(\|X_k\|_1^2 + \|\mathbf{y}_q\|_q^2 \geq M/2) \\ &\leq 2nC \exp(-sM/4). \end{aligned} \quad (\text{B.5})$$

Recall that $M = cn^\gamma$. Combining the results (B.2), (B.3), and (B.5), we have

$$\Pr(|\widehat{S}_{k1} - S_{k1}| \geq 4\varepsilon) \leq 2 \exp(-\varepsilon^2 n^{1-2\gamma}) + 2nC \exp(-sn^\gamma/4). \quad (\text{B.6})$$

In the sequel, we turn to \widehat{S}_{k2} . We write $\widehat{S}_{k2} = \widehat{S}_{k2,1} \widehat{S}_{k2,2}$, where $\widehat{S}_{k2,1} = n^{-2} \sum_{i \neq j} \|X_{ik} - X_{jk}\|_1$, and $\widehat{S}_{k2,2} = n^{-2} \sum_{i \neq j} \|\mathbf{y}_i - \mathbf{y}_j\|_q$. Similarly, we write $S_{k2} = S_{k2,1} S_{k2,2}$, where $S_{k2,1} = E\{\|X_{ik} - X_{jk}\|_1\}$ and $S_{k2,2} = E\{\|\mathbf{y}_i - \mathbf{y}_j\|_q\}$. Following arguments for proving (B.6), we can show that

$$\begin{aligned} \Pr(|\widehat{S}_{k2,1} - S_{k2,1}| \geq 4\varepsilon) &\leq 2 \exp(-\varepsilon^2 n^{1-2\gamma}) + 2nC \exp(-sn^{2\gamma}/4), \text{ and} \\ \Pr(|\widehat{S}_{k2,2} - S_{k2,2}| \geq 4\varepsilon) &\leq 2 \exp(-\varepsilon^2 n^{1-2\gamma}) + 2nC \exp(-sn^{2\gamma}/4). \end{aligned} \quad (\text{B.7})$$

Condition (C1) ensures that $S_{k2,1} \leq \{E(\|X_{ik} - X_{jk}\|_1^2)\}^{1/2} \leq \{4E(X_k^2)\}^{1/2}$ and $S_{k2,2} \leq \{E(\|\mathbf{y}_i - \mathbf{y}_j\|_q^2)\}^{1/2} \leq \{4E(\|\mathbf{y}_q\|_q^2)\}^{1/2}$ are uniformly bounded. That is,

$$\max_{1 \leq k \leq p} \{S_{k2,1}, S_{k2,2}\} \leq C,$$

for some constant C . Using (B.7) repetitively, we can easily prove that

$$\begin{aligned} \Pr(|(\widehat{S}_{k2,1} - S_{k2,1}) S_{k2,2}| \geq \varepsilon) &\leq \Pr(|\widehat{S}_{k2,1} - S_{k2,1}| \geq \varepsilon/C) \\ &\leq 2 \exp\{-\varepsilon^2 n^{1-2\gamma} / (16C^2)\} \\ &\quad + 2nC \exp(-sn^{2\gamma}/4), \\ \Pr(|S_{k2,1} (\widehat{S}_{k2,2} - S_{k2,2})| \geq \varepsilon) &\leq \Pr(|\widehat{S}_{k2,2} - S_{k2,2}| \geq \varepsilon/C) \\ &\leq 2 \exp\{-\varepsilon^2 n^{1-2\gamma} / (16C^2)\} \\ &\quad + 2nC \exp(-sn^{2\gamma}/4), \end{aligned} \quad (\text{B.8})$$

and

$$\begin{aligned} \Pr\{|\widehat{S}_{k2,1} - S_{k2,1}| |\widehat{S}_{k2,2} - S_{k2,2}| \geq \varepsilon\} &\leq \Pr(|\widehat{S}_{k2,1} - S_{k2,1}| \geq \sqrt{\varepsilon}) + \Pr(|\widehat{S}_{k2,2} - S_{k2,2}| \geq \sqrt{\varepsilon}) \\ &\leq 4 \exp(-\varepsilon n^{1-2\gamma} / 16) + 4nC \exp(-sn^{2\gamma}/4). \end{aligned} \quad (\text{B.9})$$

It follows from Bonferroni's inequality and inequalities (B.8) and (B.9) that

$$\begin{aligned} \Pr(|\widehat{S}_{k2} - S_{k2}| \geq 3\varepsilon) &= \Pr(|\widehat{S}_{k2,1}\widehat{S}_{k2,2} - S_{k2,1}S_{k2,2}| \geq 3\varepsilon) \\ &\leq \Pr\{(|\widehat{S}_{k2,1} - S_{k2,1}|)S_{k2,2} \geq \varepsilon\} \\ &\quad + \Pr\{|S_{k2,1}(\widehat{S}_{k2,2} - S_{k2,2})| \geq \varepsilon\} \\ &\quad + \Pr\{(|\widehat{S}_{k2,1} - S_{k2,1}|)(\widehat{S}_{k2,2} - S_{k2,2}) \geq \varepsilon\} \\ &\leq 8 \exp\{-\varepsilon^2 n^{1-2\gamma}/(16C^2)\} + 8nC \exp(-sn^{2\gamma}/4), \end{aligned} \tag{B.10}$$

where the last inequality holds when ε is sufficiently small and C is sufficiently large.

It remains to the uniform consistency of \widehat{S}_{k3} . We first study the following U statistics:

$$\begin{aligned} \widehat{S}_{k3}^* &= \frac{1}{n(n-1)(n-2)} \\ &\quad \times \sum_{i < j < l} \{\|X_{ik} - X_{jk}\|_1 \|y_j - y_l\|_q + \|X_{ik} - X_{lk}\|_1 \|y_j - y_l\|_q \\ &\quad + \|X_{ik} - X_{jk}\|_1 \|y_i - y_l\|_q \\ &\quad + \|X_{lk} - X_{jk}\|_1 \|y_i - y_l\|_q \\ &\quad + \|X_{ik} - X_{jk}\|_1 \|y_i - y_j\|_q \\ &\quad + \|X_{lk} - X_{ik}\|_1 \|y_i - y_j\|_q\} \\ &=: \frac{6}{n(n-1)(n-2)} \sum_{i < j < l} h_3(X_{ik}, y_i; X_{jk}, y_j; X_{lk}, y_l). \end{aligned} \tag{B.11}$$

Here, $h_3(X_{ik}, y_i; X_{jk}, y_j; X_{lk}, y_l)$ is the kernel of U statistics \widehat{S}_{k3}^* . Following the arguments to deal with \widehat{S}_{k1}^* , we decompose h_3 into two parts: $h_3 = h_3\mathbf{1}(h_3 > M) + h_3\mathbf{1}(h_3 \leq M)$. Accordingly,

$$\begin{aligned} \widehat{S}_{k3}^* &= \frac{6}{n(n-1)(n-2)} \sum_{i < j < l} h_3\mathbf{1}(h_3 \leq M) \\ &\quad + \frac{6}{n(n-1)(n-2)} \sum_{i < j < l} h_3\mathbf{1}(h_3 > M) \\ &= \widehat{S}_{k3,1}^* + \widehat{S}_{k3,2}^*, \\ S_{k3} &= E\{h_3\mathbf{1}(h_3 \leq M)\} + E\{h_3\mathbf{1}(h_3 > M)\} = S_{k3,1} + S_{k3,2}. \end{aligned}$$

Following similar arguments for proving (B.2), we can show that

$$\Pr(|\widehat{S}_{k3,1}^* - S_{k3,1}| \geq \varepsilon) \leq 2 \exp(-2\varepsilon^2 m'/M^2), \tag{B.12}$$

where $m' = \lfloor n/3 \rfloor$ because $\widehat{S}_{k3,1}^*$ is a third-order U statistics.

Then, we deal with $\widehat{S}_{k3,2}^*$. We observe that $h_3(X_{ik}, y_i; X_{jk}, y_j; X_{lk}, y_l) \leq 4(X_{ik}^2 + X_{jk}^2 + X_{lk}^2 + \|y_i\|_q^2 + \|y_j\|_q^2 + \|y_l\|_q^2)/6$, which will be smaller than M if $X_{ik}^2 + \|y_i\|_q^2 \leq M/2$, for all $1 \leq i \leq p$. Thus, for any $\varepsilon > 0$, the events satisfy

$$\{|\widehat{S}_{k3,2}^*| > \varepsilon/2\} \subseteq \{X_{ik}^2 + \|y_i\|_q^2 > M/2, \text{ for some } 1 \leq i \leq p\}.$$

By using the similar arguments to prove (B.5), it follows that

$$\begin{aligned} \Pr(|\widehat{S}_{k3,2}^* - S_{k3,2}| > \varepsilon) &\leq \Pr(|\widehat{S}_{k3,2}^*| > \varepsilon/2) \\ &\leq 2nC \exp(-sM/4). \end{aligned} \tag{B.13}$$

Then, we combine the results (B.12) and (B.13) with $M = cn^\gamma$ for some $0 < \gamma < 1/2 - \kappa$ to obtain that

$$\begin{aligned} \Pr(|\widehat{S}_{k3}^* - S_{k3}| \geq 2\varepsilon) &\leq 2 \exp(-2\varepsilon^2 n^{1-2\gamma}/3) \\ &\quad + 2nC \exp(-sn^\gamma/4). \end{aligned} \tag{B.14}$$

By the definition of \widehat{S}_{k3} ,

$$\widehat{S}_{k3} = \frac{(n-1)(n-2)}{n^2} \left\{ \widehat{S}_{k3}^* + \frac{1}{(n-2)} \widehat{S}_{k1}^* \right\}.$$

Thus, using similar techniques to deal with \widehat{S}_{k1} , we can obtain that

$$\begin{aligned} \Pr(|\widehat{S}_{k3} - S_{k3}| \geq 4\varepsilon) &= \Pr\left\{ \left| \frac{(n-1)(n-2)}{n^2} (\widehat{S}_{k3}^* - S_{k3}) \right. \right. \\ &\quad \left. \left. - \frac{3n-2}{n^2} S_{k3} + \frac{n-1}{n^2} (\widehat{S}_{k1}^* - S_{k1}) \right. \right. \\ &\quad \left. \left. + \frac{n-1}{n^2} S_{k1} \right| \geq 4\varepsilon \right\}. \end{aligned}$$

Using similar arguments for dealing with S_{k1} , we can show that S_{k3} is uniformly bounded in p . Taking n large enough such that $\{(3n-2)/n^2\}S_{k3} \leq \varepsilon$ and $\{(n-1)/n^2\}S_{k1} \leq \varepsilon$, then

$$\begin{aligned} \Pr(|\widehat{S}_{k3} - S_{k3}| \geq 4\varepsilon) &\leq \Pr(|\widehat{S}_{k3}^* - S_{k3}| \geq \varepsilon) \\ &\quad + \Pr\{|\widehat{S}_{k1}^* - S_{k1}| \geq \varepsilon\} \\ &\leq 4 \exp(-\varepsilon^2 n^{1-2\gamma}/6) \\ &\quad + 4nC \exp(-sn^\gamma/4). \end{aligned} \tag{A.15}$$

The last inequality follows from (B.6) and (B.14). This, together with (B.6), (B.10), and the Bonferroni's inequality, implies

$$\begin{aligned} \Pr\{(|\widehat{S}_{k1} + \widehat{S}_{k2} - 2\widehat{S}_{k3}) - (S_{k1} + S_{k2} - 2S_{k3})| \geq \varepsilon\} \\ \leq \Pr(|\widehat{S}_{k1} - S_{k1}| \geq \varepsilon/4) \\ + \Pr(|\widehat{S}_{k2} - S_{k2}| \geq \varepsilon/4) + \Pr(|\widehat{S}_{k3} - S_{k3}| \geq \varepsilon/4) \\ = O\{\exp(-c_1\varepsilon^2 n^{1-2\gamma}) + n \exp(-c_2 n^\gamma)\}, \end{aligned} \tag{A.16}$$

for some positive constants c_1 and c_2 . The convergence rate of the numerator of $\widehat{\omega}_k$ is now achieved. Following similar arguments, we can obtain the convergence rate of the denominator. In effect, the convergence rate of $\widehat{\omega}_k$ has the same form of (B.16). We omit the details here. Let $\varepsilon = cn^{-\kappa}$, where κ satisfies $0 < \kappa + \gamma < 1/2$. We thus have

$$\begin{aligned} \Pr\left\{ \max_{1 \leq k \leq p} |\widehat{\omega}_k - \omega_k| \geq cn^{-\kappa} \right\} &\leq p \max_{1 \leq k \leq p} \Pr\{|\widehat{\omega}_k - \omega_k| \geq cn^{-\kappa}\} \\ &\leq O\left\{ p \left[\exp\{-c_1 n^{1-2(\kappa+\gamma)}\} \right. \right. \\ &\quad \left. \left. + n \exp(-c_2 n^\gamma) \right] \right\}. \end{aligned}$$

The first part of Theorem 1 is proven.

Now, we deal with the second part of Theorem 1. If $\mathcal{D} \not\subseteq \widehat{\mathcal{D}}^*$, then there must exist some $k \in \mathcal{D}$ such that $|\widehat{\omega}_k - \omega_k| > cn^{-\kappa}$. It follows from Condition (C2) that $|\widehat{\omega}_k - \omega_k| > cn^{-\kappa}$ for some $k \in \mathcal{D}$, indicating that the events satisfy $\{\mathcal{D} \not\subseteq \widehat{\mathcal{D}}^*\} \subseteq \{|\widehat{\omega}_k - \omega_k| > cn^{-\kappa}, \text{ for some } k \in \mathcal{D}\}$, and hence $\mathcal{E}_n = \{\max_{k \in \mathcal{D}} |\widehat{\omega}_k - \omega_k| \leq cn^{-\kappa}\} \subseteq \{\mathcal{D} \subseteq \widehat{\mathcal{D}}^*\}$. Consequently,

$$\begin{aligned} \Pr(\mathcal{D} \subseteq \widehat{\mathcal{D}}^*) &\geq \Pr(\mathcal{E}_n) = 1 - \Pr(\mathcal{E}_n^c) \\ &= 1 - \Pr\left(\min_{k \in \mathcal{D}} |\widehat{\omega}_k - \omega_k| \geq cn^{-\kappa}\right) \\ &= 1 - s_n \Pr\{|\widehat{\omega}_k - \omega_k| \geq cn^{-\kappa}\} \\ &\geq 1 - O\left\{s_n \left[\exp\{-c_1 n^{1-2(\kappa+\gamma)}\} \right. \right. \\ &\quad \left. \left. + n \exp(-c_2 n^\gamma) \right] \right\}, \end{aligned}$$

where s_n is the cardinality of \mathcal{D} . This completes the proof of the second part.

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REFERENCES

Ashburner, M., Ball, C. A., Blake, J. A., Botstein, D., Butler, H., Cherry, J. M., Davis, A. P., Dolinski, K., Dwight, S. S., Eppig, J. T., Harris, M. A., Hill, D. P., Issel-Tarver, L., Kasarskis, A., Lewis, S., Matese, J. C., Richardson, J. E., Ringwald, M., Rubin, G. M., and Sherlock, G. (2000), "Gene Ontology: Tool for the Unification of Biology. The Gene Ontology Consortium," *Nature Genetics*, 25, 25-29. [1130]
 Bild, A., Yao, G., Chang, J. T., Wang, Q., Potti, A., Chasse, D., Joshi, M.-B., Harpole, D., Lancaster, J. M., Berchuck, A., Olson, J. A., Jr, Marks, J. R., Dressman, H. K., West, M., and Nevins, J. R. (2006), "Oncogenic Pathway

- Signatures in Human Cancers as a Guide to Targeted Therapies," *Nature*, 439, 353–357. [1130]
- Candes, E., and Tao, T. (2007), "The Dantzig Selector: Statistical Estimation When p is Much Larger Than n " (with discussion), *The Annals of Statistics*, 35, 2313–2404. [1129]
- Chen, L. S., Paul, D., Prentice, R. L., and Wang, P. (2011), "A Regularized Hotelling's T^2 Test for Pathway Analysis in Proteomic Studies," *Journal of the American Statistical Association*, 106, 1345–1360. [1130]
- Efron, B., Hastie, T., Johnstone, I., and Tibshirani, R. (2004), "Least Angle Regression" (with discussion), *The Annals of Statistics*, 32, 409–499. [1129]
- Efron, B., and Tibshirani, R. (2007), "On Testing the Significance of Sets of Genes," *The Annals of Applied Statistics*, 1, 107–129. [1130]
- Fan, J., Feng, Y., and Song, R. (2011), "Nonparametric Independence Screening in Sparse Ultra-High Dimensional Additive Models," *Journal of the American Statistical Association*, 106, 544–557. [1129,1130]
- Fan, J., and Li, R. (2001), "Variable Selection via Nonconcave Penalized Likelihood and Its Oracle Properties," *Journal of the American Statistical Association*, 96, 1348–1360. [1129]
- Fan, J., and Lv, J. (2008), "Sure Independence Screening for Ultrahigh Dimensional Feature Space" (with discussion), *Journal of the Royal Statistical Society, Series B*, 70, 849–911. [1129,1130,1131,1132,1133,1136]
- Fan, J., Samworth, R., and Wu, Y. (2009), "Ultrahigh Dimensional Feature Selection: Beyond the Linear Model," *Journal of Machine Learning Research*, 10, 1829–1853. [1129,1130]
- Fan, J., and Song, R. (2010), "Sure Independence Screening in Generalized Linear Models With NP-Dimensionality," *The Annals of Statistics*, 38, 3567–3604. [1129,1130]
- Hall, P., and Miller, H. (2009), "Using Generalized Correlation to Effect Variable Selection in Very High Dimensional Problems," *Journal of Computational and Graphical Statistics*, 18, 533–550. [1129,1133,1136]
- Ji, P., and Jin, J. (2012), "UPS Delivers Optimal Phase Diagram in High Dimensional Variable Selection," *The Annals of Statistics*, 40, 73–103. [1129]
- Jones, S., Zhang, X., Parsons, D. W., Lin, J. C.-H., Leary, R. J., Angenendt, P., Mankoo, P., Carter, H., Kamiyama, H., Jimeno, A., Hong, S. M., Fu, B., Lin, M. T., Calhoun, E. S., Kamiyama, M., Walter, K., Nikolskaya, T., Nikolsky, Y., Hartigan, J., Smith, D. R., Hidalgo, M., Leach, S. D., Klein, A. P., Jaffee, E. M., Goggins, M., Maitra, A., Iacobuzio-Donahue, C., Eshleman, J. R., Kern, S. E., Hruban, R. H., Karchin, R., Papadopoulos, N., Parmigiani, G., Vogelstein, B., Velculescu, V. E., and Kinzler, K. W. (2008), "Core Signaling Pathways in Human Pancreatic Cancers Revealed by Global Genomic Analyses," *Science*, 321, 1801–1806. [1130]
- Kim, Y., Choi, H., and Oh, H. S. (2008), "Smoothly Clipped Absolute Deviation on High Dimensions," *Journal of the American Statistical Association*, 103, 1665–1673. [1129]
- Mootha, V. K., Lindgren, C. M., Eriksson, K. F., Subramanian, A., Sihag, S., Lehar, J., Puigserver, P., Carlsson, E., Ridderstråle, M., Laurila, E., Houstis, N., Daly, M. J., Patterson, N., Mesirov, J. P., Golub, T. R., Tamayo, P., Spiegelman, B., Lander, E. S., Hirschhorn, J. N., Altshuler, D., and Groop, L. C. (2003), "PGC-1 α -Responsive Genes Involved in Oxidative Phosphorylation Are Coordinately Downregulated in Human Diabetes," *Nature Genetics*, 34, 267–273. [1130]
- Segal, M. R., Dahlquist, K. D., and Conklin, B. R. (2003), "Regression Approach for Microarray Data Analysis," *Journal of Computational Biology*, 10, 961–980. [1133]
- Serfling, R. J. (1980), *Approximation Theorems of Mathematical Statistics*, New York: Wiley. [1136,1137]
- Subramanian, A., Tamayo, P., Mootha, V. K., Mukherjee, S., Ebert, B. L., Gillette, M. A., Paulovich, A., Pomeroy, S. L., Golub, T. R., Lander, E. S., and Mesirov, J. P. (2005), "Gene Set Enrichment Analysis: A Knowledge-Based Approach for Interpreting Genome-Wide Expression Profiles," *Proceedings of the National Academy of Sciences of the USA*, 102, 15545–15550. [1130]
- Székely, G. J., and Rizzo, M. L. (2009), "Brownian Distance Covariance," *The Annals of Applied Statistics*, 3, 1233–1303. [1129,1130]
- Székely, G. J., Rizzo, M. L., and Bakirov, N. K. (2007), "Measuring and Testing Dependence by Correlation of Distances," *The Annals of Statistics*, 35, 2769–2794. [1129,1130,1131]
- Tian, L., Greenberg, S. A., Kong, S. W., Altschuler, J., Kohane, I. S., and Park, P. J. (2005), "Discovering Statistically Significant Pathways in Expression Profiling Studies," *Proceedings of the National Academy of Sciences of the USA*, 102, 13544–13549. [1130]
- Tibshirani, R. (1996), "Regression Shrinkage and Selection via LASSO," *Journal of the Royal Statistical Society, Series B*, 58, 267–288. [1129]
- Wang, H. (2009), "Forward Regression for Ultra-High Dimensional Variable Screening," *Journal of the American Statistical Association*, 104, 1512–1524. [1130,1131]
- Zhao, S. D., and Li, Y. (2012), "Principled Sure Independence Screening for Cox Models With Ultra-High-Dimensional Covariates," *Journal of Multivariate Analysis*, 105, 397–411. [1136]
- Zhu, L. P., Li, L., Li, R., and Zhu, L. X. (2011), "Model-Free Feature Screening for Ultrahigh Dimensional Data," *Journal of the American Statistical Association*, 106, 1464–1475. [1129,1132,1136]
- Zou, H. (2006), "The Adaptive Lasso and Its Oracle Properties," *Journal of the American Statistical Association*, 101, 1418–1429. [1129]
- Zou, H., and Hastie, T. (2005), "Regularization and Variable Selection via the Elastic Net," *Journal of the Royal Statistical Society, Series B*, 67, 301–320. [1129]
- Zou, H., and Li, R. (2008), "One-Step Sparse Estimates in Nonconcave Penalized Likelihood Models," *The Annals of Statistics*, 36, 1509–1533. [1129]
- Zou, H., and Zhang, H. H. (2009), "On the Adaptive Elastic-Net With a Diverging Number of Parameters," *The Annals of Statistics*, 37, 1733–1751. [1129]