# Row and Column Correlations (Are a set of microarrays independent of each other?) 

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#### Abstract

Having observed an $m \times n$ matrix $X$ whose rows are possibly correlated, we wish to test the hypothesis that the columns are independent of each other. Our motivation comes from microarray studies, where the rows of $X$ record expression levels for $m$ different genes, often highly correlated, while the columns represent $n$ individual microarrays, presumably obtained independently. The presumption of independence underlies all the familiar permutation, cross-validation, and bootstrap methods for microarray analysis, so it is important to know when independence fails. We develop nonparametric and normal-theory testing methods. The row and column correlations of $X$ interact with each other in a way that complicates test procedures, essentially by reducing the accuracy of the relevant estimators.


Keywords: total correlation, effective sample size, permutation tests

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## 1 Introduction

The formal statistical problem considered here can be stated simply: having observed an $m \times n$ data matrix $X$ with possibly correlated rows, test the hypothesis that the columns are independent of each other. Relationships between the row correlations and column correlations of $X$, discussed in Section 2, complicate the problem's solution.

Why are we interested in column-wise independence? the motivation in this paper comes from microarray studies, where $X$ is a matrix of expression levels for $m$ genes on $n$ microarrays. In the "Cardio" study I will use for illustration there are $m=20426$ genes each measured on $n=63$ arrays, with the microarrays corresponding to 63 subjects, 44 healthy controls and 19 cardiovascular patients. We expect the gene expressions to be correlated, inducing substantial correlations between the rows (Owen, 2005; Efron, 2007a; Qiu, Brooks, Klebanov and Yakovlev, 2005a), but most of the standard analysis techniques begin with an assumption of independence across microarrays, that is, across the columns of $X$. This can be a risky assumption, as discussed next.

An immediate purpose of the Cardio study is to identify genes involved in the disease process. For gene $i$ we compute the two-sample $t$-statistic " $t_{i}$ " comparing sick versus healthy subjects. It will be convenient for discussion to convert these to $z$-scores,

$$
\begin{equation*}
z_{i}=\Phi^{-1}\left(F_{61}\left(t_{i}\right)\right) \quad i=1,2, \ldots, m \tag{1.1}
\end{equation*}
$$

with $\Phi$ and $F_{61}$ the cumulative distribution functions (cdf) of standard normal and $t_{61}$ distributions; under the usual assumptions, $z_{i}$ will have a standard $N(0,1)$ null distribution, called here the "theoretical null." Unusually large values of $z_{i}$ or $-z_{i}$ are used to identify non-null genes, with the meaning of "unusual" depending heavily on column-wise independence.

The left panel of Figure 1 shows the histogram of all $20426 z_{i}$ values, which is seen to be much wider than $N(0,1)$ near its center. An "empirical null" fit to the center as in Efron (2007b) was estimated to be $N\left(.03,1.57^{2}\right)$. Null overdispersion has many possible causes (Efron, 2004, 2007a,b), one of which is positive correlation across the columns of $X$. Such correlations reduce the effective degrees of freedom for the $t$ statistic, causing (1.1) to yield overdispersed null $z_{i} \mathrm{~s}$, and of course changing our assessment of significance for outlying values.

The right panel of Figure 1 seems to offer a "smoking gun" for correlation: the scattergram of expression levels for microarrays 31 and 32 is strikingly correlated, with a sample correlation coefficient of .805. Here $X$ has been standardized by subtraction of its row means, so the effect is not due to so-called ecological correlations. ( $X$ is actually "doubly standardized," as discussed in Section 2.) Nevertheless the question of whether or not correlation .805 is significantly positive turns out to be surprisingly close, as discussed in Section 4, because the row-wise correlations in $X$ drastically reduce the degrees of freedom for the scatterplot.

The permutation and normal-theory tests introduced in Sections 3-5 will make it clear that there is in fact correlation across microarrays in the Cardio matrix, enough to explain the null overdispersion seen in Figure 1 (though other causes are possible, as well). All the familiar permutation, cross-validation, and bootstrap methods for microarray analysis, such as the popular SAM program of Tusher, Tibshirani and Chu (2001), depend on column-wise independence of $X$. The diagnostic tests developed here can help warn the statistician when caution is required in their use.

The paper is organized as follows: Section 2 describes the relationship between the row and column correlations of $X$, showing how row-wise correlation alone can give a misleading impression of columnwise correlation. A class of simple but useful permutation tests is discussed in Section 3. The Kronecker matrix normal distribution is introduced in Section 4 as a parametric model allowing both row and column correlations on $X$, and then used to calculate the loss of information for testing column-wise independence due to row-wise correlation. Various normal-theory tests are discussed in Section 5, with difficulties seen for all of them. The paper ends in Section 6 with a collection of remarks and details.

## 2 Row and Column Correlations

There is a close correspondence between row and column correlations, discussed next, that complicates the question of column-wise independence. For convenient presentation we assume that the data matrix $X$ has


Figure 1 Left panel: histogram of $m=20426 z$-values (1.1) for Cardio study; center of histogram is much wider than $N(0,1)$ theoretical null. Right panel: scatterplot of microarrays 31 and $32,\left(x_{i 31}, x_{i 32}\right)$ for $i=1,2, \ldots, m$, after removal of row-wise gene means; the scattergram seems to indicate substantial correlation between the two arrays.
been "demeaned" by the subtraction of row and column means, giving

$$
\begin{equation*}
\sum_{i} x_{i j}=\sum_{j} x_{i j}=0 \quad \text { for } i=1,2, \ldots, m \text { and } j=1,2, \ldots, n \tag{2.1}
\end{equation*}
$$

Our numerical results go further and assume "double standardization": that in addition to (2.1),

$$
\begin{equation*}
\sum_{j} x_{i j}^{2}=n \quad \text { and } \quad \sum_{i} x_{i j}^{2}=m \quad \text { for } i=1, \ldots, m \text { and } j=1, \ldots, n \tag{2.2}
\end{equation*}
$$

i.e., that each row and column of $X$ has mean 0 and variance 1.

The singular value decomposition of $X$ (SVD) is

$$
\begin{equation*}
\underset{m \times n}{X}=\underset{m \times K}{U} \underset{K \times K}{\underset{K}{d}} \underset{K \times n}{V^{\prime}}, \tag{2.3}
\end{equation*}
$$

where $K$ is the rank, $d$ the diagonal matrix of ordered singular values, and $U$ and $V$ orthonormal matrices of sizes $m \times K$ and $n \times K$,

$$
\begin{equation*}
U^{\prime} U=V^{\prime} V=I_{K} \tag{2.4}
\end{equation*}
$$

$I_{K}$ the $K \times K$ identity. The squares of the diagonal elements, say

$$
\begin{equation*}
e_{1} \geq e_{2} \geq \cdots \geq e_{K}>0, \quad\left(e_{k}=d_{k}^{2}\right) \tag{2.5}
\end{equation*}
$$

are the eigenvalues of $X^{\prime} X=V^{\prime} d^{2} V$.
The $n \times n$ matrix $\widehat{\boldsymbol{C o v}}$ of sample covariances between the columns of $X$ is

$$
\begin{equation*}
\widehat{\boldsymbol{C o v}}=X^{\prime} X / m \tag{2.6}
\end{equation*}
$$

and likewise

$$
\begin{equation*}
\widehat{\boldsymbol{c o v}}=X X^{\prime} / n, \tag{2.7}
\end{equation*}
$$

for the $m \times m$ matrix of row-wise sample covariances (having more than $400,000,000$ entries in the Cardio example!).

Theorem 1. If $X$ has row and column means $0,(2.1)$, then the $n^{2}$ entries of $\widehat{\boldsymbol{C o v}}$ have empirical mean 0 and variance $c_{2}$,

$$
\begin{equation*}
c_{2}=\sum_{k=1}^{K} e_{k}^{2} /(m n)^{2} \tag{2.8}
\end{equation*}
$$

and so do the $m^{2}$ entries of $\widehat{\boldsymbol{c o v}}$.
Proof. The sum of $\widehat{\boldsymbol{C o v}}$ 's entries is

$$
\begin{equation*}
1_{n}^{\prime} X^{\prime} X 1_{n} / m=0 \tag{2.9}
\end{equation*}
$$

according to (2.1), while the mean of squared entries is

$$
\begin{equation*}
\frac{\sum_{j=1}^{n} \sum_{j^{\prime}=1}^{n} \widehat{\operatorname{Cov}}_{j j^{\prime}}^{2}}{n^{2}}=\frac{\operatorname{tr}\left(\left(\mathrm{X}^{\prime} \mathrm{X}\right)^{2}\right)}{m^{2} n^{2}}=\frac{\operatorname{tr}\left(\mathrm{V}^{\prime} \mathrm{d}^{4} \mathrm{~V}\right)}{m^{2} n^{2}}=c_{2} \tag{2.10}
\end{equation*}
$$

Replacing $X^{\prime} X$ with $X X^{\prime}$ yields the same results for the row covariances $\widehat{\boldsymbol{c o v}}$.
Under double standardization (2.1)-(2.2), the covariances become sample correlations, say $\widehat{\boldsymbol{C o r} \boldsymbol{r}}$ and $\widehat{\boldsymbol{c o r}}$ for the columns and rows. Theorem 1 has a surprising consequence: whether or not the columns of $X$ are independent, the column sample correlations will have the same mean and variance as the row correlations. In other words, substantial row-wise correlation can induce the appearance of column-wise correlation.

Figure 2 concerns the 44 healthy subjects in the Cardio study, with $X$ now an $(m, n)=(20426,44)$ doubly standardized matrix. All $44^{2}$ column correlations are shown by the solid histogram, while the line histogram is a random sample of 10,000 row correlations. Here $c_{2}=.283^{2}$, so according to the Theorem both histograms have mean 0 and standard deviation .283.

The 44 diagonal elements of $\widehat{\boldsymbol{C o r}}$ protrude as a prominent spike at 1 . (We can't see the spike of 20426 diagonal elements for the row correlation matrix $\widehat{\boldsymbol{c o r}}$ because they form such a small fraction of all $20426^{2}$.) It is easy to remove the diagonal 1's from consideration.


Figure 2 Left panel: solid histogram the $44^{2}$ column sample correlations for $X$ the doubly standardized matrix of healthy Cardio subjects; line histogram is sample of 10000 of the $20426^{2}$ row correlations. Right panel: solid histogram the column correlations excluding diagonal 1s; line histogram the row correlations corrected for sampling overdispersion.

Corollary. In the doubly standardized situation, the off-diagonal elements of the column correlation matrix $\widehat{\text { Cor }}$ have empirical mean and variance

$$
\begin{equation*}
\hat{\mu}=-\frac{1}{n-1} \quad \text { and } \quad \hat{\alpha}^{2}=\frac{n}{n-1}\left(c_{2}-\frac{1}{n-1}\right) \tag{2.11}
\end{equation*}
$$

For $n=44$ and $c_{2}=.283$ this gives

$$
\begin{equation*}
\left(\hat{\mu}, \hat{\alpha}^{2}\right)=\left(-.023, .241^{2}\right) \tag{2.12}
\end{equation*}
$$

The corresponding diagonal-removing corrections for the row correlations (replacing $n$ by $m$ in (2.11)) are neglible for $m=20426$. However $c_{2}$ overestimates the variance of the row correlations for another reason: with only 44 points available to estimate each correlation, estimation error adds a considerable component of variance to the $\widehat{\boldsymbol{c o r}}$ histogram in the left panel, as discussed next.

Suppose now that the columns of $X$ are in fact independent, in which case the substantial column correlations seen in Figure 2 must actually be induced by row correlations, via Theorem 1. Let cor $\mathrm{ii}^{\prime}$ indicate the true correlation between rows $i$ and $i^{\prime}$ (that is, between $X_{i j}$ and $X_{i^{\prime} j}$ ), and define $\alpha$ the total correlation to be the root mean square of the $\operatorname{cor}_{i i^{\prime}}$ values,

$$
\begin{equation*}
\alpha^{2}=\sum_{i<i^{\prime}} \operatorname{cor}_{i i^{\prime}}^{2} /\binom{m}{2} \tag{2.13}
\end{equation*}
$$

where Remark 6.5 of Section 6 shows that $\hat{\alpha}^{2}$ in (2.11) is an approximately unbiased estimate of $\alpha^{2}$, assuming column-wise independence. For the Cardio example $\hat{\alpha}=.241$, similar to the size of the microarray correlation estimates in Efron (2007a), Owen (2005), and Qiu et al. (2005a). Section 4 discusses the crucial role of $\alpha$ in determining the accuracy of estimates based on $X$.

The right panel of Figure 2 compares the histogram of the column correlation $\widehat{C o r}_{j j^{\prime}}$, now excluding cases $j=j^{\prime}$, with the row correlation histogram corrected for sampling overdispersion via the shrinkage factor $.241 / .283$. As predicted by Theorem 1, the similarity is striking. A possible difference lies in the long right tail of the $\widehat{\boldsymbol{C O r}}$ distribution (including $\widehat{\boldsymbol{C O r}_{31,32}}$, the case illustrated in Figure 1), whose significance is examined in Section 4.

Standardization A matrix $X$ is "column standardized" by individually subtracting the mean and dividing by the standard deviation of each column, and similarly for row standardization. Table 1 shows the effect of successive row and column standardizations on the $20426 \times 44$ demeaned matrix of healthy Cardio subjects. Here "Col" is the empirical standard deviation of the 946 column-wise correlations $\widehat{C o r}_{j j^{\prime}}$, $j<j^{\prime}$; "Eig" is $\hat{\alpha}$ in (2.11); and "Row" is the empirical standard deviation " $\hat{\beta}$ " of a $1 \%$ sample of the row correlations $\widehat{c o r}_{i i^{\prime}}$, but adjusted for overdispersion,

$$
\begin{equation*}
\operatorname{Row}^{2}=\frac{n}{n-1}\left(\hat{\beta}^{2}-\frac{1}{n-1}\right) \tag{2.14}
\end{equation*}
$$

Sampling error of the Row entries is about $\pm .0034$.
The doubly standardized matrix $X$ used for Figure 2 was obtained after five successive column-row standardizations. This was excessive; the Figure looked almost the same after two iterations. Other microarray examples converged equally rapidly, though small counterexamples can be constructed where double standardization isn't possible.

Microarray analyses usually begin with some form of column-wise standardization (Bolstad, Irizarry, Åstrand and Speed, 2003; Qiu, Klebanov and Yakovlev, 2005b), designed to negate "brightness" differences between the $n$ microarrays. In the same spirit, row standardization helps prevent incidental gene differences (for example, very great or very small expression level variabilities) from obscuring the actual effects of interest. Standardization tends to reduce the apparent correlations; see Remark 6.4. Without standardization, the scatterplot in Figure 1 stretches out along the main diagonal, correlation .917 , driven by genes with unusually large or small inherent expression levels.

|  | Col | Row | Eig |  | Col | Row | Eig |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| demeaned | 0.252 | 0.286 | 0.000 | demeaned | 0.252 | 0.286 | 0.000 |
| col | 0.252 | 0.249 | 0.251 | row | 0.241 | 0.283 | 0.279 |
| row | 0.242 | 0.255 | 0.246 | col | 0.241 | 0.251 | 0.240 |
| col | 0.242 | 0.241 | 0.242 | row | 0.240 | 0.247 | 0.241 |
| row | 0.241 | 0.246 | 0.235 | col | 0.240 | 0.247 | 0.240 |
| col | 0.241 | 0.244 | 0.241 | row | 0.241 | 0.240 | 0.235 |
| row | 0.241 | 0.245 | 0.234 | col | 0.241 | 0.237 | 0.240 |
| col | 0.241 | 0.238 | 0.241 | row | 0.241 | 0.233 | 0.233 |

Table 1 Successive row and column standardizations of the $20426 \times 44$ matrix of healthy Cardio subjects. "Col" empirical standard deviation of $\widehat{C o r}_{j j^{\prime}}, j<j^{\prime}$; "Eig" $\hat{\alpha}$ from (2.11); "Row" from $1 \%$ sample of $\widehat{\operatorname{cor}}_{i i^{\prime}}$ values, adjusted for overdispersion (2.14), sampling standard error .0034.

## 3 Permutation Tests of Column-wise Independence

Simple permutation tests can provide strong evidence against column-wise independence, as we will see for the Cardio data. Let $\widehat{\Delta}$ be the familiar estimate of the $n \times n$ covariance matrix $\Delta$ for the rows of $X$,

$$
\begin{equation*}
\widehat{\Delta}=X^{\prime} X / m \tag{3.1}
\end{equation*}
$$

also called $\widehat{\boldsymbol{C o v}}$ at (2.6); notice that it is the sample covariances between columns that estimate the entries of the row covariance matrix $\Delta$, as described more carefully in Section 4. Also let $v_{1}$ denote $\widehat{\Delta}$ 's first eigenvector, the first column of $V$ in the SVD (2.3). The left panel of Figure 3 plots the components of $v_{1}$ versus array number $1,2, \ldots, 44$ for $\widehat{\Delta}$ obtained from the healthy Cardio subjects' data matrix $X$ - doubly standardized so $\widehat{\Delta}$ estimates the row correlation matrix.

Suppose that the columns of the original expression matrix, before standardization, are independent and identically distributed $m$-vectors ("i.i.d."). Then it is easy to see, Remark 6.3 of Section 6 , that all orderings of the components of $v_{1}$ are equally likely. This is not what Figure 3 shows, however, where the components seem to increase from left to right, with a noticeable block of large values for arrays $27-32$.

Let $S\left(v_{1}\right)$ be a statistic that measures structure, for instance a linear regression of $v_{1}$ versus array index. Comparing $S\left(v_{1}\right)$ with a set of permuted values

$$
\begin{equation*}
\left\{S^{* l}=S\left(v^{* l}\right), \quad l=1,2, \ldots, L\right\} \tag{3.2}
\end{equation*}
$$

$v^{* l}$ a random permutation of the components of $v_{1}$, provides a quick test of the i.i.d. null hypothesis.
Permutation testing was applied to $v_{1}$ for the Cardio data, using the "block" statistic

$$
\begin{equation*}
S\left(v_{1}\right)=v_{1}^{\prime} B v_{1} \tag{3.3}
\end{equation*}
$$

where $B$ is the $n \times n$ matrix

$$
\begin{equation*}
B=\sum_{h} \beta_{h} \beta_{h}^{\prime} . \tag{3.4}
\end{equation*}
$$

The sum in (3.4) is over all vectors $\beta_{h}$ of the form

$$
\begin{equation*}
\beta_{h}=(0,0, \cdots, 0,1,1, \cdots, 1,0,0, \cdots, 0), \tag{3.5}
\end{equation*}
$$

with the 1 s forming blocks of length between 2 and 10 inclusive. A heuristic rationale for block testing appears below; intuitively, microarray experiments are prone to block disturbances because of the way they are developed and read; see Callow et al. (2000). After $L=5000$ permutations, only three $S^{*}$ values exceeded the actual value $S\left(v_{1}\right)$, $p$-value .0006 , yielding strong evidence against the i.i.d. null hypothesis.

The right panel of Figure 3 pertains to a microarray prostate cancer study (Singh et al., 2002) discussed in Efron (2008): $m=6033$ genes were measured on each of $n=102$ men, 50 healthy controls and 52 prostate cancer patients. The right panel plots first eigenvectors for $\widehat{\Delta}$, (3.1), computed separately for the healthy


Figure 3 Left panel: Components of first eigenvector of row sample correlation matrix for the 44 healthy Cardio subjects, plotted versus array number $1,2, \ldots, 44$; dashes emphasize the block of large components for arrays 27-32. Right panel: First eigenvectors for healthy (solid line) and cancer (dashed) subjects, prostate cancer study, Singh, Febbo, et al. (2002); there was a systematic drift in expression levels as the study progressed.
controls and the cancer patients (the two matrices being individually doubly standardized). Both vectors increase almost linearly from left to right. Taking $S\left(v_{1}\right)$ as the linear regression of $v_{1}$ versus array number, permutation testing overwhelmingly rejected the i.i.d. null hypothesis, as it also did using the block test (3.3)-(3.5).

The prostate study appears as a favorable example of microarray technology in Efron (2008): the total correlation estimate $\hat{\alpha},(2.11)$, is quite small, .02 compared to .24 in (2.12); the $z$-value histogram, from twosample $t$-tests, comes close to following the theoretical $N(0,1)$ null near its center, $N\left(0,1.06^{2}\right)$ as opposed to $N\left(.03,1.57^{2}\right)$ in Figure 1; and False Discovery Rate analysis yields some 50 "significant" genes. Nevertheless, Figure 3 indicates a systematic drift in the expression level readings as the study progressed. Some genes drift up, others down (the average drift equaling 0 because of standardization), inducing a small amount of column-wise correlation.

Section 5 discusses models for $X$ where the $n \times n$ row covariance matrix $\Delta$ is of the "single degree of freedom" form

$$
\begin{equation*}
\Delta=I+\lambda \beta \beta^{\prime} \tag{3.6}
\end{equation*}
$$

for some known fixed vector $\beta$, the null hypothesis of column-wise independence being $H_{0}: \lambda=0$. An obvious choice of test statistic in this situation is

$$
\begin{equation*}
S_{\beta}=\beta^{\prime}(\widehat{\Delta}-I) \beta, \tag{3.7}
\end{equation*}
$$

a monotone increasing function of $\beta^{\prime} \widehat{\Delta} \beta$.
If $\beta$ is unknown we can replace $S_{\beta}$ with

$$
\begin{equation*}
S_{B}=\sum_{h=1}^{H} \beta_{h}^{\prime} \widehat{\Delta} \beta_{h}=\operatorname{tr}\left(\widehat{\Delta} \sum_{\mathrm{h}} \beta_{\mathrm{h}} \beta_{\mathrm{h}}^{\prime}\right) \equiv \operatorname{tr}(\widehat{\Delta} \mathrm{B}), \tag{3.8}
\end{equation*}
$$

where $\left\{\beta_{1}, \beta_{2}, \ldots, \beta_{H}\right\}$ is a catalog of "likely prospects" as in (3.5). From the SVD (2.3), $\widehat{\Delta}=\sum_{j}\left(d_{j}^{2} / m\right) v_{j} v_{j}^{\prime}$,
giving

$$
\begin{equation*}
S_{B}=\sum_{j} \frac{d_{j}^{2}}{m}\left(v_{j}^{\prime} B v_{j}\right) \tag{3.9}
\end{equation*}
$$

Finally, the model leading to (3.6) suggests that most of the information against $H_{0}$ lies in the first eigenvector $v_{1}$, getting us back to test statistic $S\left(v_{1}\right)=v_{1}^{\prime} B v_{1}$ as in (3.3). (Notice that permutations of $X$ 's columns do not affect the eigenvalues $d_{j}^{2}$.)

Two objections can be raised to our permutation tests: (1) they are really testing i.i.d., not independence; (2) non-independence might not manifest itself in the order of $v_{1}$ (particularly if the order of the microarrays has been shuffled in some unknown way).

Column-wise standardization makes the column distributions more similar, mitigating objection (1). Going further, "quantile standardization" - say replacing each column's entries by normal scores (Bolstad et al., 2003) - makes the marginals exactly the same. The Cardio data was reanalyzed using normal scores, with almost identical results.

Objection (2) is more worrisome from the point of view of statistical power. The order in which the arrays were obtained should be available to the statistician, and should be analyzed to expose possible trends like those in Figure 3. It would be desirable, nevertheless, to have independence tests that do not depend on order - that is, test statistics invariant under column-wise permutations. The discussion in Section 5 concerns the difficulties in constructing such "non-permutation" tests.

## 4 Normal Theory

Multivariate normal theory offers an alternative to the nonparametric methods of Section 3. We consider the matrix normal distribution for $X$,

$$
\begin{equation*}
\underset{m \times n}{X} \sim \mathcal{N}_{m, n}(0, \underset{m \times m}{\notin} \otimes \underset{n \times n}{\Delta}) \tag{4.1}
\end{equation*}
$$

where the Kronecker notation indicates covariance structure

$$
\begin{equation*}
\operatorname{cov}\left(X_{i j}, X_{i^{\prime} j^{\prime}}\right)=\Sigma_{i i^{\prime}} \Delta_{j j^{\prime}} \tag{4.2}
\end{equation*}
$$

Row $x_{i}$ of $X$ has covariance matrix proportional to $\Delta$,

$$
\begin{equation*}
x_{i} \sim \mathcal{N}_{n}\left(0, \nvdash_{i i} \Delta\right) \tag{4.3}
\end{equation*}
$$

( not independently unless $\ngtr$ is diagonal), and likewise for column $\boldsymbol{x}_{j}, \boldsymbol{x}_{j} \sim \mathcal{N}_{m}\left(0, \Delta_{j j} \not \subset\right)$. As in (2.1), we take all means equal 0 .

Much of classical multivariate analysis focuses on the situation $\xi=I$, where the rows $x_{i}$ are independent replicates,

$$
\begin{equation*}
\mathscr{L}=I: x_{i} \stackrel{\mathrm{iid}}{\sim} \mathcal{N}_{n}(0, \Delta) \quad i=1,2, \ldots, m \tag{4.4}
\end{equation*}
$$

in which case the sample covariance matrix $\widehat{\Delta}=X^{\prime} X / m$ has a scaled Wishart distribution,

$$
\begin{equation*}
\widehat{\Delta} \sim \operatorname{Wishart}(m, \Delta) / m \tag{4.5}
\end{equation*}
$$

Distribution (4.5) has first and second moments

$$
\begin{equation*}
\underset{n \times n}{\widehat{\Delta}} \sim\left(\underset{n \times n}{\Delta}, \Delta_{n^{2} \times n^{2}}^{(2)} / m\right) \quad \text { with } \Delta_{j k, l h}^{(2)}=\Delta_{j l} \Delta_{k h}+\Delta_{j h} \Delta_{k l} \tag{4.6}
\end{equation*}
$$

for $j, k, l, h=1,2, \ldots, n$; see Mardia, Kent and Bibby (1979, p. 92).
Relation (4.6) says that when $\not \subset=I$, that is when the rows of $X$ are independent, $\widehat{\Delta}$ unbiasedly estimates the row covariance matrix $\Delta$ with accuracy proportional to $m^{-1 / 2}$. Correlation between rows reduces the accuracy of $\widehat{\Delta}$, as shown next.

Returning to the general situation (4.1)-(4.3), define

$$
\begin{equation*}
\widetilde{\Delta}=X^{\prime} \boldsymbol{\sigma}^{-2} X / m \tag{4.7}
\end{equation*}
$$

where $\boldsymbol{\sigma}$ is the diagonal matrix with diagonal entries $\sigma_{i}=\mathcal{F}_{i i}^{1 / 2}$.

Theorem 2. Under model (4.1), $\widetilde{\Delta}$ has first and second moments

$$
\begin{equation*}
\widetilde{\Delta} \sim\left(\Delta, \Delta^{(2)} / \tilde{m}\right), \quad \tilde{m}=m /\left[1+(m-1) \alpha^{2}\right] \tag{4.8}
\end{equation*}
$$

where $\alpha$ is the total correlation as in (2.13),

$$
\begin{equation*}
\alpha^{2}=\sum_{i<i^{\prime}}\left(\not_{i i^{\prime}}^{2} / \Sigma_{i i} \not_{i^{\prime} i^{\prime}}\right) /\binom{m}{2} \tag{4.9}
\end{equation*}
$$

and $\Delta^{(2)}$ is the Wishart covariance (4.6).
Comparing (4.8) with (4.6), we see that correlation between the rows reduces "effective sample size" from $m$ to $\tilde{m}$ : for $\alpha=.241$ as in (2.12), the reduction is from $m=20426$ to $\tilde{m}=17.2$ ! (Notice that row standardization effectively makes $\sigma_{i} \doteq 1$ in (4.7), so $\widetilde{\Delta} \doteq \widehat{\Delta}$, (3.1), justifying the comparison.) The total correlation $\alpha$ shows up in other efficiency calculations; see Remark 6.7.

Proof. The row-standardized matrix $\widetilde{X}=\sigma^{-1} X$ has matrix normal distribution

$$
\begin{equation*}
\widetilde{X} \sim \mathcal{N}_{m, n}(0, \widetilde{\Sigma} \otimes \Delta) \tag{4.10}
\end{equation*}
$$

where $\widetilde{\sharp}=\boldsymbol{\sigma}^{-1} \not \mathbb{Z} \boldsymbol{\sigma}^{-1}$ has diagonal elements $\widetilde{\mathscr{Z}}_{i i}=1$. From (4.2) we see that $\widetilde{\mathscr{F}}_{i i^{\prime}}=\mathbb{Z}_{i i^{\prime}} /\left(\mathbb{Z}_{i i} \mathbb{Z}_{i^{\prime} i^{\prime}}\right)^{1 / 2}$ is the correlation between elements $X_{i j}$ and $X_{i^{\prime} j}$ in the same column of $X ; \widetilde{\Delta}=\widetilde{X}^{\prime} \widetilde{X} / m$ has entries $\widetilde{\Delta}_{j k}=$ $\sum_{i} \widetilde{X}_{i j} \widetilde{X}_{i k} / m$, and is unbiased for $\Delta$,

$$
\begin{equation*}
E\left\{\widetilde{\Delta}_{j k}\right\}=\Delta_{j k} \tag{4.11}
\end{equation*}
$$

using (4.2).
The covariance calculation for $\widetilde{\Delta}$ involves expansion

$$
\begin{align*}
\widetilde{\Delta}_{j k} \widetilde{\Delta}_{l h} & =\left(\sum_{i} \widetilde{X}_{i j} \widetilde{X}_{i k} / m\right)\left(\sum_{i^{\prime}} \widetilde{X}_{i^{\prime} l} \widetilde{X}_{i^{\prime} h} / m\right)  \tag{4.12}\\
& =\frac{1}{m^{2}}\left(\sum_{i} \widetilde{X}_{i j} \widetilde{X}_{i k} \widetilde{X}_{i l} \widetilde{X}_{i h}+\sum_{i \neq i^{\prime}} \widetilde{X}_{i j} \widetilde{X}_{i k} \widetilde{X}_{i^{\prime} l} \widetilde{X}_{i^{\prime} h}\right) \tag{4.13}
\end{align*}
$$

Using the formula

$$
\begin{equation*}
E\left\{Z_{1} Z_{2} Z_{3} Z_{4}\right\}=\gamma_{12} \gamma_{34}+\gamma_{13} \gamma_{24}+\gamma_{14} \gamma_{23} \tag{4.14}
\end{equation*}
$$

for a normal vector $\left(Z_{1} Z_{2} Z_{3} Z_{4}\right)^{\prime}$ with 0 means and covariances $\gamma_{i j}$, (4.2) gives

$$
\begin{equation*}
E\left\{\sum_{i} \tilde{X}_{i j} \tilde{X}_{i k} \tilde{X}_{i l} \tilde{X}_{i h}\right\}=m\left[\Delta_{j k} \Delta_{l h}+\Delta_{j l} \Delta_{k h}+\Delta_{j h} \Delta_{k l}\right] \tag{4.15}
\end{equation*}
$$

and

$$
\begin{equation*}
E\left\{\sum_{i \neq i^{\prime}} \widetilde{X}_{i j} \widetilde{X}_{i k} \widetilde{X}_{i^{\prime} l} \widetilde{X}_{i^{\prime} h}\right\}=m(m-1) \Delta_{j k} \Delta_{l k}+\left(\Delta_{j l} \Delta_{k h}+\Delta_{j h} \Delta_{k l}\right) \sum_{i \neq i^{\prime}} \widetilde{\mathscr{F}}_{i i^{\prime}}^{2} \tag{4.16}
\end{equation*}
$$

Then (4.13) yields

$$
\begin{equation*}
E\left\{\widetilde{\Delta}_{j k} \widetilde{\Delta}_{l h}\right\}=\Delta_{j k} \Delta_{l h}+\left(\Delta_{j l} \Delta_{k h}+\Delta_{j h} \Delta_{k l}\right)\left(\frac{1+(m-1) \alpha^{2}}{m}\right) \tag{4.17}
\end{equation*}
$$

giving

$$
\begin{equation*}
\operatorname{cov}\left(\widetilde{\Delta}_{j k}, \widetilde{\Delta}_{l h}\right)=\left(\Delta_{j l} \Delta_{k h}+\Delta_{j h} \Delta_{k l}\right) / \tilde{m} \tag{4.18}
\end{equation*}
$$

as in (4.8).
A corollary of Theorem 2 , used in Section 5 , concerns bilinear functions of $\Delta$ and $\widetilde{\Delta}$,

$$
\begin{equation*}
\tau^{2}=w^{\prime} \Delta w \quad \text { and } \quad \tilde{\tau}^{2}=w^{\prime} \widetilde{\Delta} w \tag{4.19}
\end{equation*}
$$

where $w$ is a given $n$-vector.

Corollary. Under model (4.1), $\tilde{\tau}^{2}$ has mean and variance

$$
\begin{equation*}
\tilde{\tau}^{2} \sim\left(\tau^{2}, 2 \tau^{4} / \tilde{m}\right) \tag{4.20}
\end{equation*}
$$

The proof follows that for Theorem 2; see Remark 6.9.
If $\mathbb{Z}=I$ in (4.1), then $\widetilde{\Delta}=\widehat{\Delta}$ and $\tilde{\tau}^{2}$ has a scaled chi-squared distribution,

$$
\begin{equation*}
\tilde{\tau}^{2} \sim \tau^{2} \cdot \chi_{m}^{2} / m \tag{4.21}
\end{equation*}
$$

with mean and variance $\tilde{\tau}^{2} \sim\left(\tau^{2}, 2 \tau^{4} / m\right)$, so again the effect of correlation within $\not \subset$ is to reduce the effective sample size from $m$ to $\tilde{m}$, (4.8).

We can approximate $\widetilde{\Delta},(4.7)$, with

$$
\begin{equation*}
\widehat{\Delta}=X^{\prime} \hat{\boldsymbol{\sigma}}^{-2} X / m \tag{4.22}
\end{equation*}
$$

where $\hat{\sigma}_{i i}^{2}$ is an estimate of $\mathbb{F}_{i i}$ based on the observed variability in row $i$. If the rows of $X$ have been standardized then $\hat{\sigma}_{i i}^{2}=1$, and $\widehat{\Delta}$ returns to its original definition $X^{\prime} X / m$.

Both Theorem 2 and the Corollary encourage us to think of $\widehat{\Delta}$ as, approximately, a scaled Wishart distribution based on an independent sample of size $\tilde{m}$,

$$
\begin{equation*}
\widehat{\Delta} \dot{\sim} \operatorname{Wishart}(\tilde{m}, \Delta) / \tilde{m} \tag{4.23}
\end{equation*}
$$

The dangers of this approximation are discussed in Section 5, but it is, nevertheless, an evocative heuristic, as shown below.

Figure 4 returns to the question of the seemingly overwhelming correlation .805 between arrays 31 and 32 seen in Figure 1. A one-sided $p$-value was calculated for each of the 946 column correlations, using as a null hypothesis the normal theory correlation coefficient distribution based on a sample size of $\tilde{m}=17.2$ pairs of points (the correct null if $\Delta=I$ in (4.23)). Benjamini and Hochberg's (1995) False Discovery Rate test, level $q=.1$, was applied to the $946 p$-values. This yielded 7 significant cases, those with sample correlation $\geq .723$; all 7 were from the block of arrays 27 to 32 indicated in Figure 3. Correlation .805 does turn out to be significant, but by a much closer margin than Figure 1's scattergram suggests.

The Fdr procedure was also applied using the simpler null distribution $\mathcal{N}\left(-.023, .241^{2}\right)$, (2.12). This raised the significance threshold from .723 to .780 , removing two of the previously significant correlations.

Theorem 1 showed that the variance of the observed column correlations is useless for testing columnwise independence, since any value at all can be induced by row correlations. The test in Figure 4 avoids this trap by looking for unusual outliers among the column correlations. It does not depend on the order of the columns, objection (2) in Section 3 for permutation tests, but pays the price of increased modeling assumptions.

## 5 Other Test Statistics

Theorem 2 offers a normal-theory strategy for testing column-wise independence. We begin with $X \sim$ $\mathcal{N}_{m, n}(0, \not \subset \otimes \Delta),(4.1)$, taking

$$
\begin{equation*}
\mathscr{E}_{i i}=1 \quad \text { and } \quad \Delta_{j j}=1 \quad \text { for all } i \text { and } j \tag{5.1}
\end{equation*}
$$

as suggested by double standardization. The null hypothesis of column-wise independence is equivalent to the row covariance matrix $\Delta$ equaling the identity,

$$
\begin{equation*}
H_{0}: \Delta=I \tag{5.2}
\end{equation*}
$$

since then (4.2) says that all pairs in different columns are independent.
To test (5.2), we estimate $\Delta$ with $\widehat{\Delta}$, (4.22) or more simply $\widehat{\Delta}=X^{\prime} X / m$ after standardization, and compute a test statistic

$$
\begin{equation*}
S=s(\widehat{\Delta}) \tag{5.3}
\end{equation*}
$$



Figure 4 Dashed curve is normal theory density for correlation coefficient from $\tilde{m}=17.2$ pairs of points; see Remark 6.6. Histogram is the 946 column correlations, right panel Figure 2. FDR test, $q=.1$, yielded 7 significant correlations, $\widehat{C o r} \geq .723$, including .805 between arrays 31 and 32, Figure 1.
where $s(\cdot)$ is some measure of distance between $\widehat{\Delta}$ and $I$. The accuracy approximation $\widehat{\Delta} \dot{\sim}\left(\Delta, \Delta^{(2)} / \tilde{m}\right)$ from (4.8), with $\Delta=I$, is used to assess the significance level of the observed $S$, maybe even employing the more daring approximation $\widehat{\Delta} \dot{\sim} \operatorname{Wishart}(\tilde{m}, I) / \tilde{m}$. Strategy (5.3) looks promising but, as the examples of this section will show, it suffers from serious difficulties.

One of the difficulties stems from Theorem 1. An obvious test statistic for $H_{0}: \Delta=I$ is

$$
\begin{equation*}
S=\sum_{j<j^{\prime}} \widehat{\Delta}_{j, j^{\prime}}^{2} /\binom{n}{2} \tag{5.4}
\end{equation*}
$$

the average squared off-diagonal element of $\widehat{\Delta}$. But $\widehat{\Delta}=\widehat{\boldsymbol{C o v}},(2.6)$, so in the doubly standardized situation of (2.11), $S$ is an increasing monotone function of $\hat{\alpha}$, the estimated total correlation. This disqualifies $S$ as a test statistic for (5.2), since large values of $\hat{\alpha}$ can always be attributed to row-wise correlation alone.

Similarly, the variance of the eigenvalues (2.5),

$$
\begin{equation*}
S=\sum_{k=1}^{K}\left(e_{k}-e .\right)^{2} / k \quad\left(e .=\sum e_{k} / K\right) \tag{5.5}
\end{equation*}
$$

looks appealing since the true eigenvalues all equal 1 when $\Delta=I$. However (5.5) is also a monotonic function of $\hat{\alpha}$; see Remark 6.2.

The general difficulty here is "leakage," the fact that row-wise correlations affect the observed pattern of column-wise correlations. This becomes clearer by comparison with classical multivariate methods, where row-wise correlations are assumed away by taking $\mathbb{Z}=I$ in (4.1). Johnson and Graybill (1972) consider a two-way ANOVA problem where, after subtraction of main effects, $X$ has the form

$$
\begin{equation*}
X_{i j}=a_{i} \beta_{j}+\epsilon_{i j} \quad \text { for } i=1,2, \ldots, m \text { and } j=1,2, \ldots, n \tag{5.6}
\end{equation*}
$$

$a_{i} \sim \mathcal{N}(0, \lambda)$ and $\epsilon_{i j} \sim \mathcal{N}(0,1)$, all independently, with $\beta=\left(\beta_{1}, \beta_{2}, \ldots, \beta_{n}\right)$ a fixed but unknown vector (representing "one degree of freedom for nonadditivity" in the two-way table $X$, Johnson and Graybill's extension of Tukey's procedure).

In the Kronecker notation (4.1), $X \sim \mathcal{N}_{m, n}(0, I \otimes \Delta)$ with

$$
\begin{equation*}
\Delta=I+\lambda \beta \beta^{\prime} \tag{5.7}
\end{equation*}
$$

Now (5.2) becomes $H_{0}: \lambda=0$. Johnson and Graybill show that, with $\beta$ unknown, the likelihood ratio test rejects $H_{0}$ for large values of the eigenvalue ratio

$$
\begin{equation*}
S=e_{1} / \sum_{k=1}^{K} e_{k} \tag{5.8}
\end{equation*}
$$

Since the $m$ rows of $X$ are assumed independent, they can test $H_{0}$ by comparison of $S$ with values $S^{*}=$ $e_{1}^{*} / \sum_{k=1}^{K} e_{k}^{*}$ obtained from

$$
\begin{equation*}
\widehat{\Delta}^{*} \sim \operatorname{Wishart}(m, I) / m \tag{5.9}
\end{equation*}
$$

as in (4.5).
Getting back to the correlated rows situation, Theorem 2 suggests comparing $S$ with values $S^{*}$ from

$$
\begin{equation*}
\widehat{\Delta}^{*} \sim \mathrm{Wishart}(\tilde{m}, I) / \tilde{m} \tag{5.10}
\end{equation*}
$$

$\tilde{m}$ as in (4.8). The solid histogram in Figure 5 compares $100 S^{*}$ values from (5.10), $\tilde{m}=17.2$ for the Cardio data, with the observed value $S=.207$ from the doubly standardized Cardio matrix for the healthy subjects used in Figure 2. All $100 S^{*}$ values are much smaller than $S$, providing strong evidence against $H_{0}: \Delta=I$.


Figure 5 Eigenratio statistic (5.8) equals . 207 for $20426 \times 44$ Cardio matrix $X$; solid histogram 100 simulations $S^{*}$ from Wishart (5.10), $\tilde{m}=17.2$; line histogram 100 simulations from correlated-row $X^{*}$ matrices (5.11), $\alpha=$ .241, $\Delta=I$.

The evidence looks somewhat weaker, though, if we simulate $S^{*}$ values with $\widehat{\Delta}^{*}$ obtained from random matrices

$$
\begin{equation*}
X^{*} \sim \mathcal{N}_{20426,44}(0, \not \subset \otimes I) \tag{5.11}
\end{equation*}
$$

doubly standardized, where $\not \approx$ has total correlation $\alpha=.241$, the estimated value for $X,(4.9)$. The line histogram in Figure 5 shows 100 such $S^{*}$ values, all still smaller than $S$, but substantially less so. (Remark 6.8 describes the construction of $X^{*}$.)

Why does (5.11) produce larger "null" $S^{*}$ values than (5.10)? The answer is simple: even though the first and second moments of $\widehat{\Delta}^{*}=\boldsymbol{X}^{* \prime} \boldsymbol{X}^{*} / m$ match $\widehat{\Delta}^{*}$ from (5.10), its eigenvalues do not. The non-zero eigenvalues of $X^{* \prime} X^{*} / m$ equal those of $\widehat{\mathscr{y}}^{*}=X^{*} X^{* \prime} / n$. This is another example of leakage, where the fact that $\mathbb{Z}$ in (5.11) is not the identity $I_{m}$ distorts the estimated eigenvalue of $\widehat{\Delta}^{*}$, even if $\Delta=I_{n}$.

The eigenratio statistic $S=e_{1} / \sum e_{k}$ is invariant under permutations of the columns of $X$, answering objection (2) to permutation testing of Section 3. Because of invariance, the eigenratio and permutation tests provide independent $p$-values for testing the null hypothesis of i.i.d. columns, and so can be employed together. Figure 5 is disturbing nonetheless, in suggesting that an appropriate null distribution for $S$ depends considerably on the choice of the nuisance parameter $\mathbb{\Sigma}$ in (5.11).

The bilinear form (4.19)-(4.20) yields another class of test statistics,

$$
\begin{equation*}
\hat{\tau}^{2}=w^{\prime} \widehat{\Delta} w \dot{\sim}\left(\tau^{2}, 2 \tau^{4} / \tilde{m}\right) \tag{5.12}
\end{equation*}
$$

where $w$ is a pre-chosen $n$-vector and $\tau^{2}=w^{\prime} \Delta w$. Delta-method arguments give $C V(\hat{\tau}) \doteq(2 \tilde{m})^{-1 / 2}$ for the coefficient of variation of $\hat{\tau}$. Defining

$$
\begin{equation*}
Z_{i}=x_{i}^{\prime} w \quad\left(x_{i}^{\prime} \text { the } i \text { th row of } X\right) \tag{5.13}
\end{equation*}
$$

yields the alternative form

$$
\begin{equation*}
\hat{\tau}^{2}=\sum_{i=1}^{m} Z_{i}^{2} / m \tag{5.14}
\end{equation*}
$$

In a two-sample situation like that for the Cardio study, sample sizes $n_{1}$ and $n_{2}$, we can choose

$$
\begin{equation*}
w^{\prime}=\left(\frac{n_{1} n_{2}}{n_{1}+n_{2}}\right)^{1 / 2}\left(-1_{n_{1}} / n_{1}, 1_{n_{2}} / n_{2}\right) \tag{5.15}
\end{equation*}
$$

" $1_{n}$ " indicating a vector of $n$ 1's. This choice makes

$$
\begin{equation*}
Z_{i}=\left(\frac{n_{1} n_{2}}{n_{1}+n_{2}}\right)^{1 / 2}\left(\bar{x}_{2 i}-\bar{x}_{1 i}\right) \tag{5.16}
\end{equation*}
$$

the multiple of the mean response difference between the two samples that has variance 1 if $\Delta=I$. In terms of (5.12), $\|w\|^{2}=1$ so $\tau^{2}=1$.

For the Cardio study, with $n_{1}=44, n_{2}=19$, and $\tilde{m}=17.2$, we obtain $\hat{\tau}=1.48$, coefficient of variation 0.17. This puts $\hat{\tau}$ more than 2.8 standard errors above the null hypothesis value $\tau=1$, again providing evidence against column-wise independence. The $Z_{i}$ values from (5.16) are nearly indistinguishable from the $z_{i}$ values in Figure 1 - not surprisingly since with the rows of $X$ standardized, $Z_{i}$ is an equivalent form of the two-sample $t$-statistic $t_{i}$ in (1.1).

Once again, however, there are difficulties with this as a test for column-wise independence. There is no question that the $Z_{i}$ 's are overdispersed compared to the theoretical value $\tau=1$. But problems other than column dependence can cause overdispersion, in particular unobserved covariate differences between subjects in the two samples (Efron, 2004, 2008).

The statistic $S=w^{\prime} \widehat{\Delta} w$ in (5.15) does not depend upon the order of the columns of $X$ within each of the two samples, answering objection (2) against permutation tests, but it is the only such choice for a two-sample situation. Other $w$ 's might yield interesting results. The version of (5.15) comparing the first 22 healthy Cardio subjects with the second 22 provided the spectacular value $\hat{\tau}=1.87$, and here the "unobserved covariate" objection has less force.

Now, however, the test statistic depends on the order of the columns within the healthy subjects' matrix, reviving objection (2). We might again want to check a catalog of possible $w$ vectors $w_{1}, w_{2}, \ldots, w_{H}$, leading back to test statistic

$$
\begin{equation*}
S_{B}=\sum_{h} w_{h}^{\prime} \widehat{\Delta} w_{h}=\operatorname{tr}(\widehat{\Delta} \mathrm{B}) \quad\left(\mathrm{B}=\sum_{\mathrm{h}} \mathrm{w}_{\mathrm{h}} \mathrm{w}_{\mathrm{h}}^{\prime}\right) \tag{5.17}
\end{equation*}
$$

as in (3.8), the only difference being that the null distribution of $\widehat{\Delta}$ now involves normal theory rather than permutations. Remark 6.9 shows that the null first and second moments of $S_{B}$ are similar to (5.12),

$$
\begin{equation*}
S_{B} \underset{H_{0}}{\sim}\left(\operatorname{tr}(\mathrm{~B}), \frac{2}{\tilde{\mathrm{~m}}} \operatorname{tr}\left(\mathrm{~B}^{2}\right)\right) \tag{5.18}
\end{equation*}
$$

In summary, normal-theory methods are interesting and promising, but are not yet proven competitors for the permutation tests of Section 3.

## 6 Remarks

This section presents some brief remarks and details supplementing the previous material.

Remark 6.1. Normal theory independence testing Our problem differs from classic multivariate normal independence testing, as described in Chapter 9 of Anderson (2003). There the columns of $X$ are taken always to be independent, while the independence hypotheses concern the rows - for example, whether the top half of $X$ is independent of the bottom half.

Remark 6.2. The constant $c_{2}$ The variance constant $c_{2}$ in Theorem 1, (2.8), can be expressed as

$$
\begin{equation*}
c_{2}=\frac{K}{(m n)^{2}}\left[\bar{e}^{2}+\sum_{k=1}^{K}\left(e_{k}-\bar{e}\right)^{2}\right] \quad\left(\bar{e} \equiv \sum_{1}^{K} e_{k} / K\right) \tag{6.1}
\end{equation*}
$$

so that $c_{2} \geq K(\bar{e} / m n)^{2}$, with equality only if the eigenvalues $e_{k}$ are equal. In the doubly standardized case $\bar{e}=m n / K$, giving

$$
\begin{equation*}
c_{2} \geq 1 / K \tag{6.2}
\end{equation*}
$$

where $K$ is the rank of $X$.

Remark 6.3. Permutation invariance If the columns of $X$ are i.i.d. observations from a distribution on $\mathbb{R}^{m}$, then the distribution of $X$ is invariant under permutations: $X \boldsymbol{\pi} \sim X$ for any $n \times n$ permutation matrix $\boldsymbol{\pi}$. Now suppose $\widetilde{X}=L(X)$ where $L$ performs the same operation on each column of $X$, for example replacing each column by its normal scores vector. Then

$$
\begin{equation*}
\widetilde{X} \boldsymbol{\pi}=L(X) \boldsymbol{\pi}=L(X \boldsymbol{\pi}) \sim L(X)=\widetilde{X} \tag{6.3}
\end{equation*}
$$

showing that $\tilde{X}$ is permutation invariant.
Similarly, suppose $\widetilde{X}=R(X), R$ performing the same operation $\widetilde{X}_{i}=r\left(X_{i}\right)$ on each row of $X$, where now we require $r(x) \boldsymbol{\pi}=r(x \boldsymbol{\pi})$ for all $n$-vectors $x$. The same argument as (6.3) demonstrates that $\widetilde{X}$ is still permutation invariant. Iterating row and column standardizations as in Table 1 then shows that if the original data matrix $X$ is permutation invariant, so is its doubly standardized version.

Remark 6.4. Covariances after demeaning Suppose that $X$ is normally distributed, with covariances $\not \subset \otimes \Delta$, (4.2), all columns having the same expectation vector $\mu$. Let $\widetilde{X}$ be the demeaned matrix obtained by subtracting all the row and column means of $X$. Then

$$
\begin{equation*}
\widetilde{X} \sim \mathcal{N}_{m, n}(0, \tilde{\Sigma} \otimes \widetilde{\Delta}) \tag{6.4}
\end{equation*}
$$

where

$$
\begin{equation*}
\widetilde{\Delta}_{j j^{\prime}}=\Delta_{j j^{\prime}}-\Delta_{\cdot j^{\prime}}-\Delta_{j}+\Delta_{. .} \tag{6.5}
\end{equation*}
$$

dots indicating averaging over the missing subscripts, and similarly for $\widetilde{\Sigma}$. This shows that demeaning tends to reduce covariances by recentering them around 0 .

Remark 6.5. Corrected estimates of the total correlation Suppose that the true row correlations cor ii $^{\prime}$ have mean 0 and variance $\alpha^{2}$, as in (2.13) with $\overline{c o r}=0$, and that given $c o r_{i i^{\prime}}$, the usual estimate $\widehat{c o r}_{i i^{\prime}}$ has mean and variance

$$
\begin{equation*}
\widehat{\operatorname{cor}}_{i, i^{\prime}} \doteq\left[\operatorname{cor}_{i i^{\prime}},\left(1-\operatorname{cor}_{i i^{\prime}}^{2}\right)^{2} /(n-3)\right] \tag{6.6}
\end{equation*}
$$

(6.6) being a good normal-theory approximation (Johnson and Kotz, 1970, Chap. 32). Letting $\bar{\alpha}^{2}$ be the empirical variance of the $\widehat{\operatorname{cor}}_{i i^{\prime}}$ values, a standard empirical Bayes derivation yields

$$
\begin{equation*}
\hat{\alpha}^{2}=A^{2}-\frac{3}{n-5} A^{4} \quad\left[A^{2}=\frac{(n-3) \bar{\alpha}^{2}-1}{n-5}\right] \tag{6.7}
\end{equation*}
$$

as an approxiately unbiased estimate of $\alpha^{2}$. (If $\overline{c o r}$ is not assumed to equal 0 , a slightly more complicated formula applies.) Of course $\hat{\alpha}^{2}=0$ if the right side of (6.7) is negative.

Theorem 1 implies that $\bar{\alpha}^{2}$ nearly equals $c_{2}$, (2.8), in the doubly standardized situation. Formula (2.11), with say

$$
\begin{equation*}
\tilde{\alpha}^{2}=\frac{n}{n-1}\left(\bar{\alpha}^{2}-\frac{1}{n-1}\right) \tag{6.8}
\end{equation*}
$$

is not identical to (6.7), but provides an excellent approximation for values of $\bar{\alpha} \leq 0.5$ : with $n=44$ and $\bar{\alpha}=.283$ as in (2.11), $\hat{\alpha}=.2415$ while $\tilde{\alpha}=.2412$.

Remark 6.6. Column and row centerings The column correlation mean $\hat{\mu}=-1 /(n-1)$ in (2.11) is forced by the row-wise demeaning $\sum_{j} x_{i j}=0,(2.1)$, centering the solid histogram in the right panel of Figure 2 at -.023 . With $m=20426$, the corresponding center for the line histogram is nearly 0 , and the difference in the two centerings is noticeable. The dashed density curve in Figure 4, and the corresponding $p$-values for the FDR analysis, were shifted .023 units leftwards.

Remark 6.7. The total correlation $\alpha$ The total correlation $\alpha$, which plays a key role in Theorem 2, (4.9), also is the central parameter of the theory developed in Efron (2007a). Equations (3.15)-(3.16) there are equivalent to (5.12) here. In both papers, $\alpha$ has the very convenient feature of summarizing the effects of an enormous $m \times m$ correlation matrix $\not \subset$ in a single number.

Remark 6.8. Z for simulation (5.11) The $X^{*}$ simulation used in Figure 5 began with $m \times n$ matrix $Y=\left(y_{i j}\right)$,

$$
y_{i j}=c_{I j}+e_{i j} \quad\left\{\begin{array}{l}
e_{i j} \sim \mathcal{N}(0,1)  \tag{6.9}\\
c_{I j} \sim \mathcal{N}\left(0, \alpha^{2}\right)
\end{array} \quad\right. \text { (all independent) }
$$

where $I=1,2,3,4,5$ as $i$ is in the first, second, ..., last fifth of 1 through $m$; $Y$ was then column standardized to give $X^{*}$, so that $\not \approx$ had a block form, with large positive correlations (about 0.61 ) in the $(m / 5) \times(m / 5)$ diagonal blocks. The choice $\gamma=1.23$ was required to yield $\alpha=.241$.

Remark 6.9. Bilinear statistics Since $\widetilde{\Delta} \sim\left(\Delta, \Delta^{(2)} / \tilde{m}\right)$, (4.8), it is clear that $E\left\{\tilde{\tau}^{2}\right\}=\tau^{2}$ in Corollary (4.20). The variance calculation proceeds as in Theorem 2:

$$
\begin{align*}
\operatorname{var}\left\{\tilde{\tau}^{2}\right\} & =\sum_{j k} \sum_{l h} \Delta_{j k, l h}^{(2)} w_{j} w_{k} w_{l} w_{h} / \tilde{m} \\
& =\sum_{j k} \sum_{l h}\left[\Delta_{j l} \Delta_{k h}+\Delta_{j h} \Delta_{k l}\right] w_{j} w_{k} w_{l} w_{h} / \tilde{m} \\
& =\left[\sum_{j l} \sum_{k h}\left(\Delta_{j l} w_{j} w_{l}\right)\left(\Delta_{k h} w_{k} w_{h}\right)+\sum_{j h} \sum_{k l}\left(\Delta_{j h} w_{j} w_{h}\right)\left(\Delta_{k l} w_{k} w_{l}\right)\right] / \tilde{m}  \tag{6.10}\\
& =2\left(\sum_{j k} \Delta_{j k} w_{j} w_{l}\right)^{2} / \tilde{m}=2 \tau^{4} / \tilde{m}
\end{align*}
$$

The verification of (5.18) is the same, except with element $b_{j k}$ of $B$ replacing $w_{j} w_{k}$ above, $b_{l h}$ replacing $w_{l} w_{h}$, etc.

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