Testing the independence of two random vectors where only one dimension is large

Weiming Li

School of Science, Beijing University of Posts and Telecommunications, Beijing, China

Jiaqi Chen

Department of Mathematics, Harbin Institute of Technology, Harbin, China

Jianfeng Yao

Department of Statistics and Actuarial Sciences, The University of Hong Kong, Hongkong, China

Abstract

For testing the independence of two vectors with respective dimensions $p_1$ and $p_2$, the existing literature in high-dimensional statistics all assume that both dimensions $p_1$ and $p_2$ grow to infinity with the sample size. However, as evidenced in RNA-sequencing data analysis, it happens frequently that one of the dimension is quite small and the other quite large compared to the sample size. In this paper, we address this new asymptotic framework for the independence test. A new test procedure is introduced and its asymptotic normality is established when the vectors are normally distributed. A Monte-Carlo study demonstrates the consistency of the procedure and ex-

Email addresses: liwm@bupt.edu.cn (Weiming Li), chenjq1016@gmail.com (Jiaqi Chen), jeffyao@hku.hk (Jianfeng Yao)

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hibits its superiority over some existing high-dimensional procedures. It is also shown that the procedure is robust against the normality assumption on the population vectors. Applied to a set of RNA-sequencing data, we obtain very convincing results on pairwise independence/dependence of gene isoform expressions as attested by prior knowledge established in that field. 

**Keywords:** Covariance matrix, Gene network, High-dimensional testing, Independence test

1. Introduction

Modern scientific researches increasingly encounter high dimensional data and then evoke corresponding statistical analyses. In genomics, next-generation sequencing techniques such as RNA-Sequencing (Feng et al., 2013) are designed to quantify gene expression, where typically a group of gene isoforms are analyzed and their expression data at exon levels are recorded into multidimensional vectors. The dimensions of these vectors vary in a wide range where the smallest dimension can be one or two and the largest one can be comparable to the sample size. A fundamental issue in such analyses is determining whether there is any interaction between two given gene isoforms. More formally, this problem involves testing the independence of two possibly correlated vectors in a situation where one dimension is small but the other is large compared to the sample size.

Generally, let $\mathbf{X} = (X_1, \ldots, X_{p_1})$, $\mathbf{Y} = (Y_1, \ldots, Y_{p_2})$ and $\mathbf{Z} = (\mathbf{X}, \mathbf{Y})$ be the joint vector of dimension $p := p_1 + p_2$. The covariance matrix of $\mathbf{Z}$ is partitioned as

$$
\Sigma = \begin{pmatrix}
\Sigma_{xx} & \Sigma_{xy} \\
\Sigma_{yx} & \Sigma_{yy}
\end{pmatrix}
$$
so that $\Sigma_{xx} = Var(X)$, $\Sigma_{yy} = Var(Y)$ and $\Sigma_{xy} = Cov(X, Y)$. Let $z_1, \ldots, z_N$ be a sample of size $N$ drawn from the population $Z$. The sample covariance matrix is

$$S_n = \frac{1}{n} \sum_{k=1}^{N} (z_k - \bar{z})(z_k - \bar{z})'$$

where $\bar{z} = \frac{1}{N} \sum_{k=1}^{N} z_k$ and $n = N - 1$ represents the degree of freedom. Accordingly, $S_n$ can be partitioned as

$$S_n = \begin{pmatrix} S_{xx} & S_{xy} \\ S_{yx} & S_{yy} \end{pmatrix}.$$ Assume that the joint vector $Z$ has a $p$-dimensional normal distribution with mean $\mu$ and covariance matrix $\Sigma$, the independence hypotheses of $X$ and $Y$ can be represented as

$$H_0 : \Sigma_{xy} = 0 \text{ v.s. } H_1 : \Sigma_{xy} \neq 0.$$ (1)

To test these hypotheses, the following three statistics are commonly used (Anderson, 2003), which are the likelihood ratio test (LRT) and two trace criteria:

$$\Lambda = \frac{\sup_{H_0} L(\mu, \Sigma)}{\sup L(\mu, \Sigma)} = \frac{|S_n|^{N/2}}{|S_{xx}|^{N/2}|S_{yy}|^{N/2}} = |I_p - S_{xy}S_{yy}^{-1}S_{yx}S_{xx}^{-1}|^N,$$

$$C_1 = \text{tr}(S_{xy}S_{yy}^{-1}S_{yx}S_{xx}^{-1}) \quad \text{and} \quad C_2 = \text{tr}(S_{xy}S_{yx}) - \frac{1}{n} \text{tr}(S_{xx})\text{tr}(S_{yy}).$$ (2)

The LRT statistic is the well-known Wilks’s $\Lambda$ (Wilks, 1935). Both statistics $C_1$ and $C_2$ are based on the idea that under the independence hypothesis, $\Sigma_{xy} = \Sigma_{yx}' = 0$ so that $S_{xy}$ as well as $S_{yx}$ should be small. A noticeable difference here is that the statistics $\Lambda$ and $C_1$ rely on the inverse matrices $S_{xx}^{-1}$ and $S_{yy}^{-1}$ so that essentially the conditions $p_i < n$ are required. Conversely,
the criterion $C_2$ can be applied when the dimensions $p_i$, $i = 1, 2$, are larger than the sample size $N$.

The test procedures for the classical situation where the dimensions $p_i$’s are reasonably small compared with the sample size are well studied (Anderson, 2003). It is however well understood today that these asymptotical approximations are no more valid when the dimensions are comparable to the sample size, see e.g. Ledoit & Wolf (2002), Bai et al. (2009), Chen & Qin (2010) and Wang & Yao (2013). New limiting distributions have to be found in the large-dimensional context.

Specifically for the independence test, the existing literature in the large-dimensional context includes

1. the large-dimensional limit of $\Lambda$ proposed in Jiang et al. (2013) under the asymptotic scheme $\min(p_1, p_2, n) \to \infty$, $p_1 + p_2 < n$ and $p_i/n \to c_i > 0$;

2. the large-dimensional limit of $C_1$ proposed in Jiang et al. (2013) under the asymptotic scheme $\min(p_1, p_2, n) \to \infty$, $\max(p_1, p_2) < n$ and $p_i/n \to c_i > 0$; and

3. the large-dimensional limit of $C_2$ proposed in Srivastava & Reid (2012) under the asymptotic scheme $\min(p_1, p_2, n) \to \infty$, $p_i/p \to d_i > 0$ and $n = O(p^\delta)$ for some constant $\delta > 0$ as $n \to \infty$.

Most recently, the trace criterion $C_1$ is generalized by Yang and Pan (2015) based on regularized canonical correlation coefficients to accommodate situations where $\max\{p_1, p_2\}$ can be larger than $n$. These existing asymptotic schemes are quite similar in that they all require that both dimensions $p_1$ and $p_2$ grow to infinity with the sample size $N$.

Motivated by RNA-sequencing analysis, our objective in this paper is
to test the hypotheses in (1) with the criterion $C_2$ assuming $p_1$ fixed and min($p_2, n$) → ∞. As far as we know, this scheme has not been addressed in the literature. It will be proved that the asymptotic distribution of the statistic exists under this asymptotic scenario and is consistent with the one in Srivastava & Reid (2012). Note that our proof is different from theirs and this new asymptotic scenario is not covered by their results.

The rest of this paper is organised as follows. In the next section, we present the new test procedure and examine its size and power through simulation experiments. Section 3 presents an analysis of a genomic data set and Section 4 presents some conclusions and remarks. The main theorem is proved in the last section.

2. Test for the independence in high dimensions

2.1. Test statistic and its asymptotic distribution

The null hypothesis in (1) is equivalent to $\text{tr}(\Sigma_{xy}\Sigma_{yx}) = 0$. Thus we may construct an unbiased estimator of this trace and reject the null hypothesis when this statistic is too large. Let

$$
\gamma_2 = \text{tr}(\Sigma^2), \quad \gamma_{xx} = \text{tr}(\Sigma_{xx}^2), \quad \gamma_{yy} = \text{tr}(\Sigma_{yy}^2), \quad \gamma_{xy} = \text{tr}(\Sigma_{xy}\Sigma_{yx}).
$$

We have by definition $2\gamma_{xy} = \gamma_2 - \gamma_{xx} - \gamma_{yy}$. From Srivastava (2005), an unbiased estimator of $\gamma_2$ is given as $k_n[\text{tr}(S_n^2) - \text{tr}^2(S_n)/n]$ with $k_n = n^2/(n-1)(n+2)$. Therefore an unbiased estimator of $\gamma_{xy}$ is constructed as

$$
\hat{\gamma}_{xy} = \frac{k_n}{2} \left\{ \text{tr}(S_n^2) - \text{tr}(S_{xx}^2) - \text{tr}(S_{yy}^2) - \frac{1}{n} \left[ \text{tr}^2(S_n) - \text{tr}^2(S_{xx}) - \text{tr}^2(S_{yy}) \right] \right\},
$$

$$
= k_n \left[ \text{tr}(S_{xy}S_{yx}) - \frac{1}{n} \text{tr}(S_{xx})\text{tr}(S_{yy}) \right].
$$
We thus get the trace criterion $C_2$ given in (2). Notice that the estimator $\hat{\gamma}_{xy}$ is a function of eigenvalues of the sample covariance matrices $S_{xx}, S_{yy},$ and $S_n$.

**Theorem 1.** Suppose that the dimension $p_1$ is fixed, $p_2$ and $n$ both tend to infinity, and

$$0 < \lim_{p \to \infty} \frac{1}{p} \text{tr}(\Sigma^k) < \infty, \ k = 1, 2, 4.$$  

Then under the null hypothesis in (1),

$$T_n := \frac{n}{\sqrt{2k_n}} \frac{\hat{\gamma}_{xy}}{\sqrt{\hat{\gamma}_{xx} \hat{\gamma}_{yy}}} \overset{d}{\to} N(0, 1),$$  

(3)

where $\hat{\gamma}_{xx} = k_n[\text{tr}(S_{xx}^2) - \text{tr}^2(S_{xx})/n]$ and $\hat{\gamma}_{yy} = k_n[\text{tr}(S_{yy}^2) - \text{tr}^2(S_{yy})/n]$ with $k_n = n^2/(n - 1)(n + 2)$.

This theorem establishes the asymptotic distribution of $T_n$ for the scenario that $p_1$ is fixed and $p_2$ approaches infinity. Notice that the convergence in (3) coincides with the result in Srivastava & Reid (2012), which assumes both $p_1$ and $p_2$ tend to infinity. This means that for practical applications, the proposed test is robust against different asymptotic scenarios of dimensions. Such robustness is especially welcomed since in a precise application (such as the gene isoform data analyzed in this paper) the explicit values of the dimensions $p_1$ and $p_2$ are known and it is somehow difficult to decide what is the most convenient asymptotic scenario to use. Synthesizing the two scenarios, we immediately get a more general one, i.e. $p = p_1 + p_2 \to \infty$, for the convergence in (3).

**Theorem 2.** Suppose that the dimensions $p = p_1 + p_2$ and $n$ both tend to infinity, and

(a) $0 < \lim_{p \to \infty} \frac{1}{p} \text{tr}(\Sigma^k) < \infty, \ k = 1, 2, 4;$
(b) for some $\delta \geq 0$, $\lim_{p,n \to \infty} \frac{p^{1-\delta}}{n} = 0$ and $0 < \lim_{p \to \infty} \frac{1}{p^\delta} \text{tr}(\Sigma_{xy} \Sigma_{yx})$.

Then the asymptotic power of the proposed test tends to 1 as $N \to \infty$.

Theorem 2 presents a set of sufficient conditions for the consistency of the test $T_n$. Condition (a) is the same as used in Theorem 1, which claims the existence of the fourth moments of the population spectral distribution. Condition (b) describes the range of the asymptotic frameworks that one can use. Condition (c) clarifies the requirement of the amount of $\text{tr}(\Sigma_{xy} \Sigma_{yx})$ for the consistency in a specified framework. For instance, in the most commonly used content $p/n \to c \in (0, \infty)$, we need $\text{tr}(\Sigma_{xy} \Sigma_{yx}) \to \infty$ to guarantee the consistency.

2.2. Monte-Carlo study

We numerically evaluate the finite-sample performance of the test $T_n$ and report the empirical size and power under different dimension settings. For the purpose of comparison, we also consider two tests discussed in Jiang et al. (2013): one is the corrected LRT, referred as $T_1$, and the other is based on the trace criterion $C_1$, referred as $T_2$. Since the test $T_1$ is limited to $p_1 + p_2 < n$ and $T_2$ is limited to $\max\{p_1, p_2\} < n$, we only consider the former case when comparing the three tests. The nominal significance level is fixed at $\alpha = 0.05$, and the number of independent replications is 100,000.

We first report the empirical sizes of the three tests. Samples are drawn from standard normal population, and thus $\Sigma$ is an identity matrix. The dimensions are $p_1 = 2, 6, 10$, $p_2 = 10, 30, 100, 200, 500$, and $n = 50$. The results are collected in Table 1, where the first six columns compare the sizes of the three tests when $p_1 + p_2 < n$ and the last three columns illustrate the
size of the proposed $T_n$ when $p_2 > n$. The results show that all the empirical sizes are close to the nominal significance level.

Table 1: Empirical sizes in percents for the three tests with the significant level $\alpha = 0.05$.

|        | $p_1$ | $p_2 = 10$ | $p_2 = 30$ | $T_n \& p_2$ |
|--------|-------|------------|------------|---------------|
| $T_n$  | 6.32  | 6.56       | 5.86       | 5.52          |
| $T_1$  | 6.17  | 5.90       | 4.70       | 5.30          |
| $T_2$  | 5.72  | 6.11       | 4.69       | 5.21          |

To examine the powers of the three tests, we employ a model studied in Jiang et al. (2013), where the populations $X$ and $Y$ are defined as

$$X = U_1 + \gamma U_2^{p_1}, \quad Y = U_2 + \gamma U_2, \quad U_i \sim N(0, I_{p_i}), \quad i = 1, 2,$$

respectively, where $U_1$ and $U_2$ are independent, $U_2^{p_1}$ is a subset of $U_2$ consisting of its first $p_1$ variables, and the factor $\gamma$ represents the degree of mixture. Therefore, the covariance matrices are respectively

$$\Sigma_{xx} = (1 + \gamma^2)I_{p_1}, \quad \Sigma_{yy} = (1 + \gamma)^2I_{p_2}, \quad \Sigma_{xy} = \gamma(1 + \gamma)(I_{p_1}, O_{p_1, p_2 - p_1}),$$

where $O_{m,n}$ represents an $m \times n$ zero matrix.

Figure 1 illustrates the powers of the three tests for this model. In the left panel, the parameters are $(p_1, p_2, n) = (4, 30, 50)$ and the factor $\gamma$ increases from 0 to 0.9; while on the right, $(p_1, n, \gamma) = (4, 50, 0.5)$ and $p_2$ increases from 5 to 45. The curves in the figure show that the powers of the tests $T_1$ and $T_2$ are similar, and are dominated by the proposed test $T_n$ in all the settings. Particularly, the curves in the right panel show that all the powers of the tests decrease as $p_2$ increases, which reflects the fact that in
Figure 1: Empirical powers of the three tests. The parameter settings are \((p_1, p_2, n) = (4, 30, 50), 0 \leq \gamma \leq 0.9\) in the left panel, and \((p_1, n, \gamma) = (4, 50, 0.5), 5 \leq p_2 \leq 45\) in the right panel.

this process the increasing number of zero entries of \(\Sigma_{xy}\) makes it closer to the zero matrix of the null hypothesis. However, the power of \(T_n\) declines much slower than \(T_1\) and \(T_2\), which demonstrates a greater robustness of \(T_n\) against the inflating \(p_2\).

Next we examine the robustness of the three test procedures when the assumed normal distribution of the vectors is contaminated by gamma-distributed errors. The studied model is the same as the previous except that the vector \(U_i\)'s are replaced by

\[
U_i + \theta V_i, \quad V_i = (v_{i1}, \ldots, v_{ip_i})', \quad i = 1, 2,
\]

where \(\{v_{ij}\}\), independent of \(\{U_i\}\), are \(i.i.d.\) standardized random variables derived from \(Gamma(a, b)\) distributed variables and the parameter \(\theta\) represents the level of contamination. The new parameters are set to be \(a = b = 3\) (positive skew, heavy-tailed) and \(\theta = 1/2, 2\) in this experiment. Thus the
Table 2: Empirical sizes in percents for the three tests with the significant level $\alpha = 0.05$.

|       | $p_1$ | $p_2 = 10$ | $p_2 = 30$ | $T_n \& p_2$ |
|-------|-------|------------|------------|---------------|
|       | $T_n$ | $T_1$ | $T_2$ | $T_n$ | $T_1$ | $T_2$ | 100 | 200 | 500 |
| $\theta = \frac{1}{2}$ | 6     | 5.91 | 6.17 | 5.44 | 5.47 | 5.84 | 4.60 | 5.34 | 5.24 | 5.16 |
|       | 10    | 5.71 | 5.94 | 5.18 | 5.55 | 5.80 | 4.80 | 5.33 | 5.12 | 5.22 |
| $\theta = 2$ | 6     | 6.02 | 6.15 | 5.49 | 5.65 | 5.79 | 4.67 | 5.42 | 5.42 | 5.22 |
|       | 10    | 5.85 | 6.03 | 5.27 | 5.68 | 5.82 | 4.84 | 5.35 | 5.33 | 5.06 |

covariance matrices become

$$
\Sigma_{xx} = (1 + \gamma^2)(1 + \theta^2)I_{p_1}, \quad \Sigma_{yy} = (1 + \gamma)^2(1 + \theta^2)I_{p_2},
$$

$$
\Sigma_{xy} = \gamma(1 + \gamma)(1 + \theta^2)(I_{p_1} - \rho_{p_1,p_2}).
$$

Results about the empirical sizes and powers of the tests are collected in Table 2 and Figure 2, respectively. It shows that all the sizes are close to the nominal one and the power curves are quite similar to those in Figure 1, which demonstrate that the additional gamma-distributed errors have little impact on the three tests. It is however worth noticing that the theoretic proof of Theorem 1 in this paper as well as the proofs for asymptotic normality of the test criteria $T_1$ and $T_2$ established in Jiang et al. (2013) all heavily rely on the assumed normality of the vectors, and to our best knowledge, it seems unclear how these proofs can be extended to cover non-normal data as the ones tested in the Monte-Carlo experiments reported here.
Figure 2: Empirical powers of the three tests for the non-normal distribution with $\theta = 1/2, 2$. The parameter settings are $(p_1, p_2, n) = (4, 30, 50)$, $0 \leq \gamma \leq 0.9$ in the left panel, and $(p_1, n, \gamma) = (4, 50, 0.5)$, $5 \leq p_2 \leq 45$ in the right panel.
3. Real data analysis

Genomes play a central role in the control of cellular processes (Barabasi & Oltvai, 2004). The dynamic interplay between various genes can be mapped as gene co-expression networks, which is an important and widely used method to understand the cause and prognosis of various diseases. To recover pairwise dependencies in a gene co-expression network, each co-expression edge has to be inferred by accepting or rejecting the independence hypothesis from the sample covariance matrix of respective isoform expressions.

We analyze a data set of liver cancer, which is downloaded from TCGA data portal: https://tcga-data.nci.nih.gov/tcga/dataAccessMatrix.htm, and filtered by data types RNASeqV2 and Level 3. The data set consists of 38 genes with their dimensions ranging from 1 to 31 (see Table 3) and their sample size is $N = 50$. Obviously the dimensions are not on the same order of magnitude as their sample size. For these genes, the relationship of dependency are totally known based on established knowledge from historical experiments: 29 pairs of them are dependent and the remaining 674 pairs are independent.

We test the pairwise gene dependencies using $T_n$ and compare the results with those from two other methods: one is from Hong et al. (2013), which is a variant of traditional canonical correlation analysis (CCA); the other is the large-dimensional trace criterion $T_2$, which is recently applied in Yalamanchili et al. (2014) and is demonstrated better than CCA. The corrected LRT $T_1$ is excluded from this comparison since its dimensional requirement is not met for the data set. The significance level is set to be $\alpha = 0.05$. To evaluate the accuracy of the test results, we employ the so called F-score
Table 3: Lung cancer data: 38 genes with different dimensions

| Name      | NM000222 | NM000321 | NM000636 | NM000791 | NM001126116 | NM001140 |
|-----------|----------|----------|----------|----------|--------------|----------|
| Dimension | 20       | 27       | 4        | 6        | 7            | 13       |
| Name      | NM001145102 | NM001204191 | NM001237 | NM001429 | NM001759      | NM001760 |
| Dimension | 9        | 7        | 8        | 31       | 5            | 4        |
| Name      | NM001786 | NM001880 | NM001950 | NM002198 | NM002228      | NM002421 |
| Dimension | 4        | 13       | 10       | 9        | 1            | 10       |
| Name      | NM002467 | NM002505 | NM002539 | NM002985 | NM003109      | NM003153 |
| Dimension | 3        | 10       | 11       | 3        | 6            | 22       |
| Name      | NM003221 | NM003998 | NM004379 | NM004417 | NM005194      | NM005238 |
| Dimension | 7        | 23       | 8        | 2        | 1            | 8        |
| Name      | NM005239 | NM005252 | NM005438 | NM007122 | NM022457      | NM033285 |
| Dimension | 10       | 2        | 4        | 11       | 20           | 4        |
| Name      | NM053056 | NM198253 |          |          |              |          |
| Dimension | 5        | 15       |          |          |              |          |

(Powers, 2007) which actually measures the trade-off between precision $P$ and recall $R$:

$$F = 2 \times \frac{P \times R}{P + R},$$  \hspace{1cm} (4)

where

$$P = \frac{\text{true positives}}{\text{true positives} + \text{false positives}}, \quad R = \frac{\text{true positives}}{\text{true positives} + \text{false negatives}}.$$

With the prior information of dependency, the true positives stands for the number of correctly identified correlated pairs of genes, the false positive is the number of misidentified correlated pairs of genes, and the false negatives is the number of misidentified uncorrelated pairs of genes.

The F-scores reported in Table 4 show that $T_n$ outperforms $T_2$ significantly. CCA fails to detect the relationship between gene NM002228 and
other genes due to the dimension of this gene is 1. The same phenomenon happens to gene NM005195. Therefore, we cannot get F-score for CCA.

| Method | $T_n$ | $T_2$ | CCA |
|--------|-------|-------|-----|
| F-score| 0.64  | 0.40  | NA  |

Next, we remove the 1-dimensional genes from the data set in order to incorporate CCA for comparison. The remaining 36 genes include 25 dependent pairs and 605 independent pairs. The F-scores collected in Table 5 demonstrate that $T_n$ again outperforms the others. Notice that such results on pairwise dependence of gene isoform expressions are further used to construct gene co-expression networks, see Yalamanchili et al. (2014).

| Method | $T_n$ | $T_2$ | CCA |
|--------|-------|-------|-----|
| F-score| 0.6465| 0.4238| 0.4187|

4. Concluding remarks

This paper investigates the independence test of two vectors in a high-dimensional situation where one of the dimensions $p_1$ is quite small while the other dimension $p_2$ is large compared to the sample size. The asymptotic scheme is novel and practically useful. A new procedure is introduced and the test statistic under the null is proved to be asymptotically normal distributed assuming that $p_1 + p_2 \to \infty$ and the vectors are normal distributed. The power of the proposed test is studied through Monte-Carlo simulations and a real data analysis, which demonstrates the superiority of the new test.
over the existing ones. Another interesting feature found in the Monte-Carlo study is that the proposed procedure is robust against deviations from the normality assumption on the vectors although a theoretic proof of this fact is still missing.

5. Proofs

5.1. Lemma

Lemma 1. Let \( u, v, \) and \( w \) be independent vectors of \( n \)-dimensional standard normal distribution \( N(0, I_n) \), and define

\[
\psi(x, y) = \frac{1}{n}(x'y)^2 - \frac{1}{n^2}(x'x)(y'y),
\]

then

\[
E[\psi(u, v)|u] = 0, \quad E[\psi(u, v)\psi(w, v)|u, w] = \frac{2}{n}\psi(u, w),
\]

\[
E[\psi(v, v)] = (n - 1)(n + 2)/n, \quad E[\psi^2(u, v)] = 2(n - 1)(n + 2)/n^2,
\]

\[
E[\psi^2(v, v)] = O(n^2), \quad \text{Var}[\psi^2(v, v)] = O(n), \quad E[\psi^4(u, v)] = O(1),
\]
as \( n \to \infty \).

Proof. The distribution of \( v'v \) is \( \chi^2(n) \) and the conditional distribution of \( u'v|u \) is \( N(0, u'u) \), thus \( E[\psi(u, v)|u] = 0 \). Write

\[
\psi(u, v)\psi(w, v) = \frac{1}{n^2}(u'v)^2(w'v)^2 - \frac{1}{n^3}(u'v)^2(w'w)(v'v)
\]

\[
- \frac{1}{n^3}(w'v)^2(u'u)(v'v) + \frac{1}{n^4}(u'u)(w'w)(v'v)^2
\]

\[= \frac{1}{n^2}S_1 - \frac{1}{n^3}S_2 - \frac{1}{n^3}S_3 + \frac{1}{n^4}S_4.\]
Then $E(S_4|\mathbf{u}, \mathbf{w}) = n(n + 2)(\mathbf{u}'\mathbf{u})(\mathbf{w}'\mathbf{w})$, and

$$E(S_1|\mathbf{u}, \mathbf{w}) = \sum_{i,j,k,l} u_i u_j w_k w_l E(v_i v_j v_k v_l)$$

$$= \sum_{i,j,k=l} u_i u_j w_k w_l + \sum_{i=k,j=l} u_i u_j w_k w_l + \sum_{i=l,j=k} u_i u_j w_k w_l$$

$$= (\mathbf{u}'\mathbf{u})(\mathbf{w}'\mathbf{w}) + 2(\mathbf{u}'\mathbf{w})^2,$$

$$E(S_2|\mathbf{u}, \mathbf{w}) = (\mathbf{w}'\mathbf{w}) \sum_{i,k} u_i^2 \cdot E(v_i^2 v_k^2) = (n + 2)(\mathbf{u}'\mathbf{u})(\mathbf{w}'\mathbf{w}),$$

and thus $E(S_3|\mathbf{u}, \mathbf{w}) = E(S_2|\mathbf{u}, \mathbf{w})$, where $(x_i)$ denote the elements of $\mathbf{x}$. Collecting these results, we get $E[\psi(\mathbf{u}, \mathbf{v})|\psi(\mathbf{w}, \mathbf{v})|\mathbf{u}, \mathbf{w}] = (2/n)\psi(\mathbf{u}, \mathbf{w})$.

Notice that $\psi(\mathbf{v}, \mathbf{v}) = (n - 1)(\mathbf{v}'\mathbf{v})^2/n^2$, and $E(\mathbf{v}'\mathbf{v})^k = n(n + 2) \cdots (n + 2k - 2), k \in \mathbb{N}^+$. We have then,

$$E[\psi(\mathbf{v}, \mathbf{v})] = (n - 1)(n + 2)/n,$$

$$E[\psi^2(\mathbf{u}, \mathbf{v})] = (2/n)E[\psi(\mathbf{u}, \mathbf{u})] = 2(n - 1)(n + 2)/n^2,$$

$$E[\psi^2(\mathbf{v}, \mathbf{v})] = E(\mathbf{v}'\mathbf{v})^4(n - 1)^2/n^4 = O(n^2),$$

$$Var[\psi^2(\mathbf{v}, \mathbf{v})] = [E(\mathbf{v}'\mathbf{v})^4 - E^2(\mathbf{v}'\mathbf{v})^2] (n - 1)^2/n^4 = O(n).$$

Finally, from Minkowski inequality,

$$E[\psi^4(\mathbf{u}, \mathbf{v})] = \frac{1}{n^4} E[(\mathbf{u}'\mathbf{v})^2 - (\mathbf{u}'\mathbf{u})(\mathbf{v}'\mathbf{v})]/n^4$$

$$\leq \frac{1}{n^4} \left\{ \left[ E(\mathbf{u}'\mathbf{v})^8 \right]^{\frac{1}{4}} + \left[ E(\mathbf{u}'\mathbf{u})^4 E(\mathbf{v}'\mathbf{v})^4 \right]^{\frac{1}{2}} / n \right\}^4$$

$$= \frac{1}{n^4} \left\{ \left[ E(\mathbf{v}'\mathbf{v})^4 \right]^{\frac{1}{4}} + \left[ E(\mathbf{v}'\mathbf{v})^4 \right]^{\frac{1}{2}} / n \right\}^4,$$

which is $O(1)$ as $n \to \infty$. \hfill \Box

5.2. Proof of Theorem 1

The sample covariance $S_n$ has the Wishart distribution $W_n(\Sigma)$ with $n$ degrees of freedom. It can be expressed as $\sum_{i=1}^n \tilde{z}_i \tilde{z}_k' / n$ where $(\tilde{z}_i)$ are i.i.d.
$N(0, \Sigma)$. Write $\tilde{z}_i = (\tilde{x}_i', \tilde{y}_i')' = (\tilde{x}_{i1}, \ldots, \tilde{x}_{ip_1}, \tilde{y}_{i1}, \ldots, \tilde{y}_{ip_2})'$, $i = 1, \ldots, n$, and denote $X = (\tilde{x}_1, \ldots, \tilde{x}_n)$ and $Y = (\tilde{y}_1, \ldots, \tilde{y}_n)$. Note that the matrices $X$ and $Y$ contain normal vectors which are independent under $H_0$. The matrices $X'X$ and $Y'Y$ can be standardized as

$$X'X = \sum_{i=1}^{p_1} \alpha_i u_i' u_i', \quad Y'Y = \sum_{j=1}^{p_2} \beta_j v_j' v_j,$$

where $(\alpha_i)$ and $(\beta_j)$ are the eigenvalues of $\Sigma_{xx}$ and $\Sigma_{yy}$, respectively, and $(u_i), (v_j)$ are i.i.d. $N(0, I_n)$. Therefore, we have

$$\frac{n}{k_n} \hat{\gamma}_{xy} = \frac{1}{n} \text{tr}(S_{xy}S_{yx}) - \frac{1}{n} \text{tr}(S_{xx})\text{tr}(S_{yy})$$

$$= \frac{1}{n} \text{tr}(X'XY') - \frac{1}{n^2} \text{tr}(X'X)\text{tr}(Y'Y)$$

$$= \sum_{i=1}^{p_1} \sum_{j=1}^{p_2} \alpha_i \beta_j \left[ \frac{1}{n} (u_i' v_j)^2 - \frac{1}{n^2} (u_i' u_i)(v_j' v_j) \right]$$

$$:= \sum_{i=1}^{p_1} \sum_{j=1}^{p_2} a_{ij} \psi(u_i, v_j),$$

where $a_{ij} = \alpha_i \beta_j$ and $\psi$ is defined in (5) with the dimension $n$, $i = 1, \ldots, p_1$, $j = 1, \ldots, p_2$.

We use the martingale CLT to establish the limiting distribution of $T_n$. Without loss of generality suppose that $p_1 \leq p_2$, and define $\phi_j^{(n)} = (1/\sqrt{p_1 p_2}) \sum_{i=1}^{p_1} a_{ij} \psi(u_i, v_j)$, $j = 1, \ldots, p_2$. Let $\mathcal{F}_j^{(n)}$ be the $\sigma$-algebra generated by the random variables $\{u_1, \ldots, u_{p_1}, v_1, \ldots, v_j\}$, then $\emptyset, \Omega = \mathcal{F}_0 \subset \mathcal{F}_1^{(n)} \subset \cdots \subset \mathcal{F}_{p_2}^{(n)} \subset \mathcal{F}$ with $(\Omega, \mathcal{F}, P)$ the probability space. From Lemma
1 and the law of iterated expectations,

\[
E \left[ \phi_j^{(n)} \bigg| \mathcal{F}_{j-1}^{(n)} \right] = \frac{1}{\sqrt{p_1p_2}} \sum_{i=1}^{p_1} a_{ij} E(\psi(u_i, v_j)|u_i) = 0,
\]

\[
E \left[ \phi_j^{(n)} \right]^2 = \frac{1}{p_1p_2} \sum_{i=1}^{p_1} \sum_{k=1}^{p_1} a_{ij} a_{kj} E(\psi(u_i, v_j)\psi(u_k, v_j))
\]

\[= \frac{2(n-1)(n+2)}{n^2p_1p_2} \sum_{i=1}^{p_1} a_{ij}^2,
\]

which is \(O(1/p_2)\) as \((p, n) \to \infty\). Thus \(\{\psi_j^{(n)}, \mathcal{F}_j^{(n)}\}\) forms a sequence of integrable martingale differences. On the other hand,

\[
\sum_{j=1}^{p_2} E \left[ \left( \phi_j^{(n)} \right)^2 \bigg| \mathcal{F}_{j-1}^{(n)} \right] = \frac{1}{p_1p_2} \sum_{j=1}^{p_2} \sum_{i=1}^{p_1} \sum_{k=1}^{p_1} a_{ij} a_{kj} E(\psi(u_i, v_j)\psi(u_k, v_j)|u_i, u_k)
\]

\[= 2 \sum_{j=1}^{p_2} b_{ij} \psi(u_i, u_j) + 2 \sum_{i \neq k} b_{ik} \psi(u_i, u_k)
\]

\[= A_{1n} + A_{2n},
\]

where \(b_{ik} = \sum_{j=1}^{p_2} a_{ij} a_{kj}, i, k = 1, \ldots, p_1\). Considering the variances of \(A_{1n}\) and \(A_{2n}\), \(\text{Var}(A_{1n}) = O(1/n)\) and

\[
\text{Var}(A_{2n}) = \frac{4}{n^2p_1p_2} \sum_{i \neq k} \sum_{l \neq s} b_{ik} b_{ls} E[\psi(u_i, u_k)\psi(u_l, u_s)]
\]

\[= \frac{8}{n^2p_1p_2} \sum_{i \neq k} b_{ik}^2 E[\psi^2(u_i, u_k)],
\]

which is \(O(1/n^2)\). Therefore, from the Chebyshev inequality,

\[
\sum_{j=1}^{p_2} E \left[ \left( \phi_j^{(n)} \right)^2 \bigg| \mathcal{F}_{j-1}^{(n)} \right] - \sum_{j=1}^{p_2} E \left( \phi_j^{(n)} \right)^2 \overset{\text{P}}{\to} 0, \quad \text{as} \quad (p, n) \to \infty,
\]

where the second expectation has expression \(s_n^2 := 2(1-1/n)(1+2/n) \sum_{i=1}^{p_1} \sum_{j=1}^{p_2} a_{ij}^2 / (p_1p_2)\).
Next we verify Lyapunov condition by showing that
\[ B_n = \sum_{j=1}^{p^2} E(\phi_j^{(n)})^4 \to 0. \]
From Lemma 1 and the law of iterated expectations,
\[
B_n = \frac{1}{p_1^2 p_2^2} \sum_{j=1}^{p^2} \sum_{i=1}^{p_1} a_{ij} a_{ij} a_{ij} a_{ij} E[\psi(u_i, v_j) \psi(u_i, v_j) \psi(u_s, v_j) \psi(u_t, v_j)]
\]
\[
= \frac{1}{p_1^2 p_2^2} \sum_{j=1}^{p^2} \sum_{i=1}^{p_1} a_{ij}^4 E[\psi^4(u_i, v_j)] + \frac{3}{p_1^2 p_2^2} \sum_{j=1}^{p^2} \sum_{i \neq s} a_{ij}^2 a_{sj}^2 E[\psi^2(u_i, v_j) \psi^2(u_s, v_j)]
\]
\[
= \frac{1}{p_1^2 p_2^2} \sum_{j=1}^{p^2} \sum_{i=1}^{p_1} a_{ij}^4 E[\psi^4(u_i, v_j)] + \frac{12}{n^2 p_1^2 p_2^2} \sum_{j=1}^{p^2} \sum_{i \neq s} a_{ij}^2 a_{sj}^2 E[\psi^2(v_j, v_j)],
\]
which is \( O(1/p_2) \) as \((p, n) \to \infty\).

Notice that \( \hat{\gamma}_{xx} \) and \( \hat{\gamma}_{yy} \) are unbiased and consistent estimators of \( \gamma_{xx} \)
and \( \gamma_{yy} \), respectively. The statistic \( s_n^2 := 2(1 - 1/n)(1 + 2/n)\hat{\gamma}_{xx} \hat{\gamma}_{yy}/(p_1 p_2) \)
is also an unbiased and consistent estimator of \( s_n^2 \) under the null hypothesis, therefore
\[
\frac{n}{\sqrt{2kn}} \sqrt{\frac{\hat{\gamma}_{xy}}{\hat{\gamma}_{xx} \hat{\gamma}_{yy}}} = \frac{1}{s_n} \sum_{j=1}^{p^2} \phi_j^{(n)} \xrightarrow{d} N(0, 1), \quad \text{as} \quad (p, n) \to \infty.
\]

5.3. Proof of Theorem 2

Denote \( \hat{\gamma}_2 = k_n \{ \text{tr}(S_n^2) - \frac{1}{n} \text{tr}^2(S_n) \} \). Under the assumptions of the theorem, from Lemma 6.4 in Srivastava (2005), we have
\[
E(\hat{\gamma}_2) = \text{tr}(\Sigma^2)
\]
and
\[
\text{Var}(\hat{\gamma}_2) = \frac{8(n + 3)}{n(n + 2)} \text{tr}(\Sigma^4) + \frac{4}{n + 2}(\text{tr}^2(\Sigma^2) - \text{tr}(\Sigma^4)).
\]

From Chebyshev inequality, \((\hat{\gamma}_2 - \text{tr}(\Sigma^2))/p\) converges in probability to 0, as \(n, p\) tend to infinity. Similarly, we can get
\[
\frac{1}{p_1} (\hat{\gamma}_{xx} - \gamma_{xx}) \overset{p}{\to} 0, \quad \frac{1}{p_2} (\hat{\gamma}_{yy} - \gamma_{yy}) \overset{p}{\to} 0.
\]
For the statistic \( \hat{\gamma}_{xy} \), it’s easy to see that 
\[
\hat{\gamma}_{xy} = \left( \hat{\gamma}_2 - \hat{\gamma}_{xx} - \hat{\gamma}_{yy} \right)/2,
\]
and thus we get 
\[
E(\hat{\gamma}_{xy}) = \gamma_{xy} \quad \text{and} \quad \text{Var}
\]
\[
(\hat{\gamma}_{xy}/p^\delta) = \frac{1}{4p^\delta^2} \text{Var}(\hat{\gamma}_2 - \hat{\gamma}_{xx} - \hat{\gamma}_{yy}) \leq \frac{3}{4p^\delta^2} \left( \text{Var}(\hat{\gamma}_2) + \text{Var}(\hat{\gamma}_{xx}) + \text{Var}(\hat{\gamma}_{yy}) \right) \to 0.
\]

Therefore, 
\[
(\hat{\gamma}_{xy} - \gamma_{xy})/p^\delta \xrightarrow{P} 0, \quad \text{as} \quad n, p \text{ tend to infinity.}
\]
From this, the results in (6), and again the assumptions of the theorem, we conclude that 
\[
T_n = \frac{n}{p^{1-\delta}} \cdot \frac{p}{\sqrt{2k_n p_1 p_2}} \cdot \frac{\hat{\gamma}_{xy}/p^\delta}{\sqrt{\hat{\gamma}_{xx}\hat{\gamma}_{yy}/p_1 p_2}} \xrightarrow{p} \infty, \quad n, p \to \infty,
\]
which implies the result of the theorem.

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