The effects of age, cigarette smoking, sex and race on the qualitative characteristics of lung transcriptome

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Qingzhou Guan
fujian medical university

Juan Zhang
fujian medical university

You Guo
fujian medical university

Jie Xia
fujian medical university

Jiahui Zhang
fujian medical university

Jiajing Xie
fujian medical unicersity

Hao Cai
fujian medical university

Haidan Yan
fujian medical university

Xianlong Wang
fujian medical university

Zheng Guo
guoz@ems.hrbmu.edu.cn
Harbin Medical University

Corresponding Author

ORCiD: 0000-0003-4466-6026

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Abstract

Background

Methods based on within-sample relative expression orderings (REOs) comparisons could be applied for various medical issues such as individualized diagnosis of cancer and subtype identification etc., it could also be used for identifying differentially expressed genes (DEGs) at the individual level and detecting disease-associated genes based on one-phenotype disease data by reusing data of normal samples from other sources. However, the common potential confounding factors, including age, cigarette smoking, sex and race whether could affect the REOs of gene pairs is still unclear. Here, we evaluated these confounding factors on the REOs of gene pairs within normal lung tissues transcriptome.

Results

For one confounding factor, based on the number of related gene pairs or DEGs, the effect of this confounding factor on REO was evaluated. Our results showed that age has little effect on REOs within lung tissues. We found that about 0.23% of the significantly stable REOs of gene pairs in non-smokers’ lung tissues are reversed in smokers’ lung tissues, introduced by 344 DEGs between the two groups of samples (RankCompV2, FDR < 0.05), which are enriched in metabolism of xenobiotics by cytochrome P450, glutathione metabolism and other pathways (hypergeometric test, FDR < 0.05). Comparison between the normal lung tissue samples of males and females revealed fewer reversal REOs introduced by 24 DEGs between the sex groups, among which 19 DEGs are located on sex chromosomes and 5 DEGs involving in spermatogenesis and regulation of oocyte are located on autosomes. Between the normal lung tissue samples of white and black people, we identified 22 DEGs (RankCompV2, FDR < 0.05) which introduced a few reversal REOs between the two races.
Conclusions

In summary, the REO-based study should consider the confounding factors of cigarette smoking, sex and race.

Background

Recently, we have revealed an important biological phenomenon that, despite high variations of gene expression levels among different individuals, the within-sample relative expression orderings (REOs) of genes, namely the qualitative transcriptional characteristics, are highly stable in a particular type of normal human tissue, which might be an intrinsic mechanism to keep genes functioning coordinately in the normal tissues. On the other hand, the stable REOs in the normal tissues are widely disturbed in diseased tissues[1, 2], providing abundant information for characterizing diseases[1, 3]. Compared with the quantitative transcriptional signature, the qualitative transcriptional characteristics are highly robust against measurement variations and experimental batch effects[4-6], which could be applied for the individualized analysis without the need of inter-sample data normalization or experimental batch adjustment. Besides, they are also insensitive to the specimens sampled from different tumor locations of the same patients[7], partial RNA degradation during the process of specimen preparation and storage[8] and amplification bias for minimum specimens[9], the above issues are the common causes of quantitative transcriptional signature failure in clinical practice. Actually, taking these unique advantages of the REOs, some REO-based methods such as TSP[10], K-TSP[11] and others[12, 13] have been developed for discriminating cancer subtypes. Especially, many REO-based prognostic signatures have been proposed for specific medical issues for various cancers such as non-small cell lung cancer[3, 14], colorectal cancer[4, 15] and other cancers[16-18]. Moreover, the REO-based algorithm could also identify the differentially expressed genes (DEGs) with changes in both mRNA
concentration and absolute abundances[19].

Besides, based on the REOs analysis, we have proposed an algorithm named RankComp[1] to detect DEGs for an individual disease sample compared with its previously normal state through analyzing which genes’ up- or down-regulation may lead to the reversal REOs in the disease sample, taking the stable REOs predetermined in a large collection of the normal tissue samples as the normal background[1, 20]. The individual-level analysis of DEGs allows us to identify subtype-specific genes, which can provide us novel perspectives for understanding the mechanisms of carcinogenesis[20]. In contrast, for a DEG detected at the population-level, we cannot know whether it is differentially expressed in a particular cancer sample because of the heterogeneity of cancer. The REOs analysis method could also be applied to the identification of disease-associated genes or pathways based on one-phenotype disease data when the normal tissues are unavailable or insufficient for some vital organs such like brain and heart[1, 21-23]. In this situation, it is of great value to reuse the normal control data accumulated in other studies. And we have proposed a REO-based algorithm, named DRFunc[23], to identify disease-associated pathways based on one-phenotype data through comparing the stable REO in the one-phenotype disease samples with the normal stable REOs background pre-determined in previously accumulated normal samples from other studies. Based on the REOs analysis, we have also proposed a method named “RankCompV2” for identifying DEGs at the population-level through comparing the stable REOs of two phenotypes[24].

In summary, the REO-based analysis methods could circumvent the critical limitation of quantitative transcriptional signatures, with giant potential value of clinical application[4]. However, some confounding factors such like age, cigarette smoking, sex and race may affect the gene expression levels in normal samples. Studies have shown that sex-biased gene expression is widespread across genomes on both sex chromosomes and
autosomes[25, 26]. Several studies have also reported that cigarette smoking[27] and race[28] could alert the gene expression levels, and the gene expression levels change with age in many organ tissues, including lung tissues[29]. However, whether those confounding factors could affect the REO of gene pairs is still unclear. Thus, in this paper, using the normal lung tissue samples from three different laboratories, we evaluated the effects of four confounding factors, including age, cigarette smoking, sex and race, on the REOs within normal lung tissues.

Methods

Data and preprocessing
The gene expression profiles analyzed in this study are described in Table 5. All the datasets were measured by the Affymetrix GPL570 platform and the processed data were directly downloaded from the Gene Expression Omnibus database. For the downloaded data, each probe ID was mapped to Entrez gene ID with the corresponding platform file. If a probe was mapped to multiple or zero genes, the data were discarded. If multiple probes were mapped to the same gene, the expression value of the gene was defined as the arithmetic mean of the values of these probes.

Evaluation of confounding factors on REO of gene pairs
Within a sample, the REO of two genes, A and B, is denoted as A > B (or B < A) if the expression level of gene A is higher (or lower) than that of gene B. For each of the three binary confounding factors, cigarette smoking, sex and race, we first divided the samples into two groups, and then identified the gene pairs with significantly stable REOs in each of the group. The significance of a gene pair with one REO pattern in a group of samples was determined by the binomial test as follows:
Due to technical limitations, Equation 1 has been placed in the Supplementary Files section.

where the REO pattern (A > B or A < B) is consistent among \( k \) samples out of \( n \) samples in total and \( p_0 \) (\( p_0 = 0.5 \)) is the probability of observing one of two possible REO outcomes in a sample by chance. The \( p \)-values were then adjusted using the Benjamini-Hochberg method[42].

A gene pair with stable REOs in both groups of samples but the REO directions are opposite is called a reversal gene pair. Otherwise, if the REO directions are consistent in both groups, it is called a concordant gene pair. If the two lists of stable gene pairs identified above have \( m \) common pairs, among which \( k \) have opposite REO directions, the reversal ratio is calculated as \( k/m \).

Between the two groups of samples classified by a binary confounding factor, the distribution of other confounding factors between the two groups was tested by the Fisher’s exact test to ensure there is no significant difference for the other confounding factors. For the age factor, the samples were divided into two groups based on the REO pattern of each gene pair, and then we test whether there is significant difference in age between the two groups of samples based on the Mann-Whitney U-test. The REO of the gene pair is significantly correlated with age if the age is significantly different between the two groups.

**Identification of Differentially Expressed Genes**

Focusing on the stable gene pairs commonly identified from two groups of samples, we identified the concordant and the reversal REOs between the two groups for a specific factor. RankCompV2[24] was applied to detect differentially expressed genes (DEGs) between the two groups of samples. The details of the RankCompV2 algorithm has been
described in ref. [24]. Briefly, Fisher’s exact test was applied to identify whether a gene may disrupt the gene correlation structure in one group compared to the other group based on the concordant and the reversal REOs between the two groups. For a particular gene, to minimize the potential effect of other genes’ expression changes on the Fisher’s exact test, an iterative filter process [43] was conducted.

Pathway enrichment analysis

Data of 238 pathways covering 6638 unique genes were extracted from the Kyoto Encyclopedia of Genes and Genomes (KEGG) on 3 May 2017. The hypergeometric distribution model was used to determine the significance of biological pathways enriched with up- and down-regulated DEGs, respectively.

Results

The influence of age on REOs within normal lung tissues

From three datasets (GSE31210, GSE19804 and GSE20257, as shown in Table 5), we selected 65 samples of non-smoking Asian females with age ranging from 37 to 80 years old for the analysis. Based on the REO pattern of each gene pair, the samples were divided into two groups, and then the Mann-Whitney U-test was used to test whether there is significant difference in age between the two groups of samples. We could not find any gene pair whose REO was significantly correlated with age with FDR < 0.05 or even with FDR < 0.2 (Methods). Similarly, using 34 samples for Caucasian males with age ranging from 27 to 80 years old, collected from the dataset GSE4115, no significant gene pair was found with either FDR < 0.05 or FDR < 0.2.

The above results indicated that the influence of age on REO of gene pair could be negligible. Accordingly, the age factor was not considered in the subsequent analyses.

The influence of cigarette smoking on REOs within normal lung tissues
We compared the gene expression profiles of normal lung tissue samples for 49 smokers and 44 non-smokers from the GSE20257 dataset. The detailed information on the sample composition was shown in Table 1. There is no significant difference in sex or race distribution between the smoker group and the non-smoker group (Fisher’s exact test, $P > 0.1$).

With FDR < 0.05, we identified the gene pairs with significantly stable REOs in the smoker group and non-smoker group, respectively. We found 187,875,560 gene pairs that have significantly stable REOs (binomial test, FDR < 0.05) in both groups, among which 0.227% showed reversal REO patterns. With RankCompV2, we identified 344 DEGs, including 210 up- and 134 down-regulated genes in the smoker group compared with the non-smoker group (FDR < 0.05). The 210 up-regulated genes and 134 down-regulated genes were enriched, respectively, in 7 pathways and 1 pathway (hypergeometric test, FDR < 0.05), as shown in Figure 1. For the pathway “metabolism of xenobiotics by cytochrome P450”, cytochrome P450 are known to be responsible for the metabolism of compounds present in cigarette smoke, including nicotine, benzene, polycyclic aromatic hydrocarbons (PAHs) and tobacco-specific nitrosamines (TSNAs)[30]. As for the “glutathione metabolism” pathway, it has been found that cigarette smoking could induce the deregulation of glutathione metabolism in bronchial epithelial cells[31]. It has also been reported that “metabolic pathways”[32], “steroid hormone biosynthesis”[33], “pentose phosphate pathway”[34], “arachidonic acid metabolism”[35] and “mineral absorption”[36] are affected by cigarette smoking.

The above results indicated that cigarette smoking can alter the REOs in normal lung tissues and disturb some important biological pathways.
The influence of sex on REOs within normal lung tissues

We compared the gene expression profiles of normal lung tissue samples for 64 males and 29 females from the dataset GSE20257. The detailed information of the sample composition was shown in Table 2. There is no significant difference in smoking rate or race distribution between the male group and the female group (Fisher’s exact test, \( P > 0.2 \)).

We identified the gene pairs with significantly stable REOs in the male and female groups, respectively, and found 187,481,246 gene pairs with significantly stable REOs (binomial test, FDR < 0.05) in both groups, among which 0.074% showed the reversal REO patterns. With RankCompV2, we identified 35 DEGs in the male group compared with the female group (FDR < 0.05). In another dataset GSE71181, including 201 male samples and 80 female samples which are all from smokers, 25 of the above 35 DEGs were also found (T-test, FDR < 0.05) and 96% (24 genes) have the same dysregulation directions in the male group compared with the female group. Among the 24 DEGs, 6 out of the 10 up-regulated genes in the male group are located on Y chromosome, 12 out of the 14 up-regulated genes in the female group are located on X chromosome, and the cytoband of these genes is shown in Table 3. In particular, DDX43, CRISP2 and PRDM7, which are up-regulated in the male group are located on autosome and involved in spermatogenesis and male fertility[37, 38]. For the other two genes, NLRP2 and C3orf79, located on autosome but up-regulated in the females, it is known that NLRP2 is a critical regulator of oocyte[39].

The influence of the race factor on REOs within normal lung tissues

Due to the limitation of the sample sizes for other races, we only compared the gene expression profiles of normal lung tissues for the white and black races. From the
GSE20257 dataset, we obtained 34 samples for white people and 59 samples for black people. The detailed information of the sample composition was shown in Table 4. There is no significant difference in cigarette smoking rate or sex distribution between the two groups (Fisher’s exact test, $P > 0.1$).

With FDR < 0.05, we found 187,973,147 gene pairs with significantly stable REOs in both groups, among which 0.0272% showed reversal REO patterns. With RankCompV2, we identified 22 DEGs, including 10 up- and 12 down-regulated genes in the white group compared with the black group (FDR < 0.05). Due to the small number of DEGs, we found no pathway significantly enriched with the up- or down-regulated DEGs with FDR < 0.05. With $P < 0.05$, the 10 up-regulated and 12 down-regulated genes were enriched in, respectively, 4 and 4 pathways, as shown in Figure 2. The result indicates that there are some differences in metabolism and immunity of the normal lung tissues between the white and black races[40, 41].

Discussion

Among the four confounding factors investigated in this paper, cigarette smoking alters the REOs within lung tissues most widely, and sex and race can also alter the REOs but only slightly, whereas there is no evidence that age could affect the REO of gene pairs. Therefore, the REO-based study should consider the confounding factors of cigarette smoking, sex and race. When building the normal stable REOs background based on previously accumulated normal samples from other studies, the normal samples should include enough samples with the same factors presenting in the one-phenotype disease samples analyzed in a study.

Our results showed that cigarette smoking disrupts “Metabolism of xenobiotics by cytochrome P450”, “Glutathione metabolism” and other pathways[30, 31], and there are
some differences in metabolism and immunity between different races. The sex factor affects some genes located on the sex chromosome and some genes located on the autosomes which are involved in spermatogenesis, male fertility[37] and are critical regulator of oocyte[39]. Because cigarette smoking, sex and race could affect the REO of gene pairs, the influence of these factors should be considered in the REO-based analysis for lung tissue.

This study exists some limitations. Due to the limitation of normal tissue samples and clinic information for many other organs, we only systematically analyze the influence of the four common confounding factors (age, cigarette smoking, sex and race) on REOs in the normal lung tissues. The effects of the confounding factors on the REOs might be tissue specific. We have preliminary analyzed the influence of sex on REOs of gene pairs in normal stomach tissues and esophagus tissues, respectively, and found that all the DEGs are located on sex chromosome, as described in Additional file 1. Studies on the effect of confounding factor on the REOs of gene pairs in tissues of other organs need to be further studied.

Conclusions

Our results show that the confounding factors, including cigarette smoking, sex and race could alter the REOs within lung tissues. Thus, the REO-based study should consider these confounding factors. Moreover, the effect of age on REO of gene pair could be negligible.

Abbreviations

REO: relative expression orderings

DEGs: differentially expressed genes

GEO: Gene Expression Omnibus KEGG: Kyoto Encyclopedia of Genes and Genomes

Declarations
Ethics approval and consent to participate

Not applicable

Consent for publication

Not applicable

Availability of data and materials

The public datasets generated and/or analysed during the current study are available in Gene Expression Omnibus repository (GEO, http://www.ncbi.nlm.nih.gov/geo/).

Competing interests

The authors declare that they have no competing interests

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Authors’ contributions

QZG and JZ conceived the study, analysed the data, made figures, performed the statistical analysis, and drafted the manuscript. YG, JX, JHZ, JJX and HC searched the data and participated in the statistical analysis. HDY participated in discussing and revising the manuscript. ZG and XLW conceived of the study, and participated in its design and coordination, helped to draft the manuscript and supervised the work. All authors read and approved the final manuscript.

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Tables

Table 1. The data extracted from GSE20257 for the analysis of the cigarette smoking factor

| Characteristics | Smoker  
| (n = 49) | Non-smoker  
| (n = 44) | Fisher’s Exact Test |
|---|---|---|---|
| Sex | | | \( P = 0.266 \) |
| Male | 31 | 33 | |
| Female | 18 | 11 | |
| Race | | | \( P = 0.131 \) |
| White | 14 | 20 | |
| Black | 35 | 24 | |

Table 2. The data extracted from GSE20257 for the analysis of the sex factor

| Characteristics | Male  
| (n = 64) | Female  
| (n = 29) | Fisher’s Exact Test |
|---|---|---|---|
| Race | | | \( P = 0.495 \) |
| White | 25 | 9 | |
| Black | 39 | 20 | |
| Smoking history | | | \( P = 0.266 \) |
| Smoker | 31 | 18 | |
| Non-smoker | 33 | 11 | |

Table 3. The cytoband of the 24 sex-biased genes
| Symbol | Cytoband | Symbol | Cytoband |
|--------|----------|--------|----------|
| TTTY10 | Yq11.221 | SMC1A  | Xp11.22-p11.21 |
| PRKY   | Yp11.2   | DDX3X  | Xp11.3-p11.23 |
| TBL1Y  | Yp11.2   | STS    | Xp22.32  |
| KDM5D  | Yq11     | RIBC1  | Xp11.22  |
| DDX3Y  | Yq11     | ZFX    | Xp21.3   |
| UTY    | Yq11     | EFHC2  | Xp11.3   |
| ARSE   | Xp22.3   | KDM6A  | Xp11.2   |
| PRDM7  | 16q24.3  | JPX    | Xq13.2   |
| DDX43  | 6q13     | ZRSR2  | Xp22.1   |
| CRISP2 | 6p12.3   | PNPLA4 | Xp22.3   |
|        |          | ARSD   | Xp22.3   |
|        |          | GEMIN8 | Xp22.2   |
|        |          | C3orf79| 3q25.2   |
|        |          | NLRP2  | 19q13.42 |

Table 4. The data extracted from GSE20257 for the analysis of the race factor

| Characteristics | White \((n = 34)\) | Black \((n = 59)\) | Fisher’s Exact Test | 
|-----------------|-------------------|-------------------|---------------------|
| Sex             |                   |                   | \(P = 0.495\)      |
| Male            | 25                | 39                |                     |
| Female          | 9                 | 20                |                     |
| Smoking history |                   |                   | \(P = 0.131\)      |
| Smokers         | 14                | 35                |                     |
| Non-smokers     | 20                | 24                |                     |

Table 5. Data used in this study
| Characteristics          | GEO Acc | GSE31210 | GSE19804 | GSE20257 |
|--------------------------|---------|----------|----------|----------|
| Sample Size              |         | 20       | 60       | 93       |
| Age                      | median  | 59 (30 - 89) | 61 (37 - 80) | 45 (21-73) |
| Smoking history           |         |          |          |          |
| smoker                   |         | 12       | /        | 49       |
| non-smoker               |         | 8        | 60       | 44       |
| Sex                      |         |          |          |          |
| male                     |         | 11       | /        | 64       |
| female                   |         | 9        | 60       | 29       |
| Race                     |         |          |          |          |
| Asian                    |         | 20       | 60       | /        |
| White                    |         | /        | /        | 34       |
| Black                    |         | /        | /        | 59       |

Notes: “/” cells indicate that there is no sample in the corresponding category

Figures

![KEGG Pathways Diagram](image)

**Figure 1**

The KEGG pathways separately enriched with up- and down-regulated genes in the smoker group compared with the non-smoker group.
Figure 2

The KEGG pathways separately enriched with up- and down-regulated genes in the white people compared with the black people.

Supplementary Files

This is a list of supplementary files associated with the primary manuscript. Click to download.

Equation 1.png
Additional file 1.docx