Genome-wide mRNA profiling identifies the NRF2-regulated lymphocyte oxidative stress status in patients with silicosis

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Abstract

Background: The immunomodulatory abnormalities of silicosis are related to the lymphocyte oxidative stress state. The potential effect of antioxidant therapy on silicosis may depend on the variation in nuclear factor erythroid 2-related factor 2 (NRF2)-regulated antioxidant genes in peripheral blood mononuclear cells (PBMCs). As NRF2 is a redox-sensitive transcription factor, its possible roles and underlying mechanism in the treatment of silicosis need to be clarified.

Methods: Ninety-two male patients with silicosis and 87 male healthy volunteers were randomly selected. PBMCs were isolated from fresh blood from patients with silicosis and healthy controls. The lymphocyte oxidative stress state was investigated by evaluating NRF2 expression and NRF2-dependent antioxidative genes in PBMCs from patients with silicosis. Key differentially expressed genes (DEGs) and signaling pathways were identified utilizing RNA sequencing (RNA-Seq) and bioinformatics technology. Gene set enrichment analysis was used to identify the differences in NRF2 signaling networks between patients with silicosis and healthy controls.

Results: The number of monocytes was significantly higher in patients with silicosis than that of healthy controls. Furthermore, RNA-Seq findings were confirmed using quantitative polymerase chain reaction and revealed that NRF2-regulated DEGs were associated with glutathione metabolism, transforming growth factor-β, and the extracellular matrix receptor interaction signaling pathway in PBMCs from patients with silicosis. The top 10 hub genes were identified by PPI analysis: SMAD2, MAPK3, THBS1, SMAD3, ITGB3, integrin alpha-V (ITGAV), von Willebrand factor (VWF), BMP4, CD44, and SMAD7.

Conclusions: These findings suggest that NRF2 signaling regulates the lymphocyte oxidative stress state and may contribute to fibrogenic responses in human PBMCs. Therefore, NRF2 might serve as a novel preventive and therapeutic candidate for silicosis.

Keywords: Silicosis, Nuclear factor-erythroid 2-related factor 2, Fibrosis, GSEA-analysis, Peripheral blood mononuclear cells

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Introduction

Silicosis is a pulmonary interstitial fibrosis disease caused by the exposure to crystalline silica dust. This disease is a progressive process characterized by lung inflammation during the early phase and triggering pulmonary fibrosis with subsequent impairment of lung function [1, 2]. Unfortunately, no effective treatments are available, except the possibility of a lung transplant in a small minority of patients [3]. At present, a large number of employees are exposed to high concentrations of dust in a booming mining industry in China. The number of new pneumoconiosis reported per year reached more than 20,000 in China, almost half of which was silicosis [4].

Silicosis is characterized not only by its direct fibrotic effect on lung tissue but by immunomodulatory abnormalities, such as the appearance of complications of autoimmune diseases and autoantibodies in silica-exposed populations [5–7]. Peripheral blood mononuclear cells (PBMCs) are collected from the peripheral or circulating blood and possess a single, round nucleus. PBMCs are the major immune cells in the human body and provide selective responses to the immune system. The roles of these blood cells are to adapt to intruders and fight infection. PBMCs mainly comprise T-lymphocytes, B-lymphocytes, and natural killer (NK) cells and are also a rich source of monocytes, such as macrophages and dendritic cells [8]. It was reported that lymphocytes account for 70–95% in PBMCs [9, 10]. PBMCs play a crucial role in maintaining and controlling the immunomodulatory process in silicosis pathogenesis [11]. It has been reported that the gene associated with apoptosis was significantly overexpressed in PBMCs derived from patients with silicosis and the dysregulation of the genes in silicosis cases could cause the appearance of autoantibodies and acquisition of autoimmune diseases sequentially [12, 13]. Animal experiments demonstrated that silica exposure induces migration of dendritic cells from the peripheral blood into the alveoli in rats [14]. Taken together, PBMCs are associated with the pathogenesis or development of silicosis.

Oxidative stress occurs due to a persistent imbalance of redox homeostasis induced by overproduction of reactive oxygen species (ROS) in silica-exposed populations. When ROS production exceeds antioxidant capacity, oxidative stress may have harmful effects on the structure and function of biological tissues. It is essential for oxidative damage to occur in the development of silicosis [15–17]. It was showed that the antioxidant N-acetylcysteine alleviates lung fibrosis induced by silica in rats by downregulating ROS and interfering with apoptosis signaling in our previous study [18]. Antioxidant treatments have been shown to be effective for silicosis in animal experiments [19–21]. However, antioxidant therapy has produced conflicting results in clinical studies of the treatment of pulmonary fibrosis and other chronic diseases [22, 23]. The failure of clinical trials may be due to the lack of understanding of the role of ROS in the development of silicosis. Indeed, clinical effects of antioxidant therapy on silicosis depend on the redox microenvironment, which changes spatially and temporally in different cell types and in different subcellular compartments [24]. Thus, the individual’s difference in redox microenvironment potentially depends on variation of their antioxidant genes. We hypothesize that the endogenous antioxidant levels in the subjects might have greatly affected the results of the clinical trials.

Antioxidant defense systems to combat ROS from tissues oxidative damage include enzymatic antioxidants and nonenzymatic antioxidants. Enzymatic antioxidants comprise superoxide dismutases (SODs), catalase (CAT), glutathione peroxidases (GPx), glutathione reductase (GR), and aldehyde dehydrogenases (ALDH). Glutathione (GSH), coenzyme Q10, ascorbic acid (vitamin C) and α-tocopherol (vitamin E) represent nonenzymatic antioxidants, which influence each other and have their unique roles [25]. The GPx/GR antioxidant system is related to other antioxidant systems, including the SOD/CAT system. GSH modulates the neutralization of free radicals by vitamin C (ascorbic acid) and vitamin E [26]. Occupational silica exposure leads to the alteration of SOD and CAT activities in the antioxidant defense system [27]. Oxidative stress status as demonstrated by the level of antioxidant biomolecules could provide an early warning signal in diseases related to oxidative damage such as cancer, cataract, and heart disease [28].

Nuclear factor erythroid 2-related factor 2 (NRF2) is a redox-sensitive transcription factor and plays a critical role in maintaining redox homeostasis by regulating the expression of antioxidant defense enzymes [29]. Levels of NRF2 in PBMCs increase immediately after ozone/oxygen exposure [30]. In a human intervention study, antioxidant modulated the NRF2-related gene expression in PBMCs [31]. Our previous research shows that NRF2 is involved in mediating the development of silicosis in animal experiments [32]. Although studies in animals have preliminarily revealed that the pathogenic process of silicosis associated with oxidative stress is regulated by NRF2, the integrated mechanism requires further exploration. Advances in molecular biology and genomics have enabled further exploration as high-throughput sequencing technology has been widely used in disease diagnosis and prognosis. However, gene expression profile changes have rarely been discussed in PBMCs of patients with silicosis.

In the present study, genome-wide mRNA profiling was used to identify the lymphocyte oxidative stress state
regulated by NRF2 in patients with silicosis, and to investigate whether there was an association between lymphocyte oxidative stress state and the pathogenesis of silicosis. RNA sequencing (RNA-Seq) technology and bioinformatics technology were comprehensively integrated and subsequently identified key differentially expressed genes (DEGs) and signaling pathways. The role of DEGs in silicosis was explored from three aspects, including cellular components, molecular function, and biological process, which were also used to mine potential pathogenic genes and biomarkers. Furthermore, the clinical significance of identified genes was verified using clinical samples. These data provide novel information and further understanding of the mechanism underlying oxidative stress in PBMCs from patients with silicosis.

Materials and methods
Study design and participants
A total of 92 male patients with silicosis (the range of age: 40–65 years) and 87 male healthy volunteers (the range of age: 40–65 years) were randomly selected from an institute of occupational disease prevention and control from Henan Province in the study. The healthy controls group had no dust exposure in the past. The group of patients with silicosis were strictly selected according to the standard diagnoses of pneumoconiosis (GBZ70–2015), and the patients were diagnosed as silicosis stage I. All subjects had no clinical symptoms of autoimmune disease, including Raynaud’s phenomenon, sclerotic skin, arthralgia or facial erythema, or malignant tumors. Additionally, all subjects were negative for pneumonia, active pulmonary tuberculosis, pulmonary heart disease, infectious diseases and other lung related diseases. All subjects confirmed that they understood the experimental procedure and provided written informed consent. The study was approved by the Ethics Committee of the Xinxiang Medical University (protocol number XYLL-2017086, approved 3 March 2017).

In this study, information was collected using selected survey questionnaires by trained medical personnel. The survey mainly included information related to age, smoking status, basic health status, and family history. Physical examinations included blood pressure, weight, and height measurements to determine body mass and body mass index (BMI). Levels of PBMCs in every subject were detected by an automated hematology analyzer (Cell-Dyn Sapphire, Abbott, USA). Pulmonary function evaluation was performed using a portable handheld spirometer (Drägerwerk AG, Lübeck, Germany). The general specifications for the performance, as well as interpretation, of the pulmonary function test were followed [33, 34]. The pulmonary function indices documented were the forced vital capacity (FVC) and forced expiratory volume in 1 second (FEV1). All readings were in units of liters and percentage predicted values. The transforming growth factor (TGF)-β1 content was detected in accordance with manufacturer instructions by ELISA. In addition, three patients with silicosis and three healthy donors were randomly selected for the detection of genome-wide mRNA profiling. Thirty-six participants were respectively selected from the remaining healthy and patient samples with random and double-blindness method for mRNA analysis, immunofluorescence and western blot analysis.

PBMC isolation
PBMCs were isolated from 10 ml of fresh blood from patients with silicosis and healthy controls within 30 min after collection using the Ficoll-Paque (GE Healthcare Bio-Sciences, Pittsburgh, PA, USA) density centrifugation method. Briefly, blood was diluted 1:1 with RPMI-1640 medium at room temperature (Solarbio, Beijing, China), underlaid with 10 ml Ficoll-Paque, and then centrifuged (1000×g, 10 min, 20 °C) with acceleration set to six and deceleration set to zero in a Heraeus Multifuge X3R (Thermo Fisher Scientific, Langenselbold, Germany). Separated PBMCs were carefully collected and transferred to a 50 ml conical tube, which was brought up to 50 ml with RPMI-1640 medium, and then washed three times with RPMI-1640 medium (100×g, 10 min, 4 °C) with acceleration set to nine and deceleration set to six. The supernatant was removed and the pellet was resuspended in 1.0 ml RPMI-1640 medium for total RNA extraction, immunofluorescence assays or western blotting. PBMCs were enumerated and assessed for viability using trypan blue. Samples were stored at −80 °C before use.

Expression of NRF2 in PBMCs of patients with silicosis using immunofluorescence
After PBMCs were isolated from peripheral blood, the expression of NRF2 in PBMCs of patients with silicosis was determined using a fluorescent immunocytochemistry assay. PBMCs were diluted to 2×10⁶/ml with phosphate-buffered saline (PBS). A total of 300 μl of diluted suspension was spread evenly over the slide and allowed to air-dry. Then, PBMCs were fixed with 4% paraformaldehyde for 20 min at room temperature and washed three times with PBS. Sections were incubated with primary antibodies to NRF2 (Abcam, 1:100) overnight at 4 °C and then incubated with Cy3-conjugated secondary antibodies (1:200) at room temperature for 50 min in the dark. Nuclei were labeled using 4′, 6-diamidine-2′-phenylindole dihydrochloride and the results observed under a confocal laser scanning microscope (TCS STED-CW, Leica Microsystems, Mannheim, Germany). Sixteen areas per sample covering the whole
spectrum of blood smear, were evaluated in a blinded fashion. Immunofluorescence images were analyzed using Image J Software. Finally, fluorescent intensity of NRF2 staining was assessed by a pathologist blinded to study groups.

Expression of NRF2 and NRF2 regulated proteins in PBMCs of patients with silicosis using western blotting

Total proteins were extracted from PBMCs isolated from peripheral blood. Protein concentration was quantified using a BCA assay kit (Thermo, USA). Equal amounts of total protein were separated by 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis and then transferred onto a polyvinylidene fluoride membrane (Millipore). After blocked for 2 h in 5% nonfat milk, they were incubated with the primary antibodies: NRF2 (1:1000) (Affinity Biosciences, USA), GCLM (1:2000) (Abcam, USA), KEAP-1(1:2000, Cell Signaling, Boston, MA, USA), anti-GAPDH antibodies (1:5000, Proteintech, USA) overnight at 4°C. The membranes were washed for 3 times in 1% Tris-buffered saline with Tween (TBST) containing 0.01% Tween 20 and incubated with horseradish peroxidase conjugated secondary antibodies (1:3000, Affinity Biosciences, USA) for 1.0 h at room temperature. After washing membranes, immunoreactive bands were visualized using a Fluor Chem R imaging analysis system (Protein Simple, California, USA). Finally, the density of proteins in each band were analyzed using Image J Software. All western blot analyses were performed at least three times.

Total RNA extraction, library construction, and sequencing

Total cellular RNA was extracted from PBMCs using the TRIzol reagent (Gibco BRL Life Technologies, Gaithersburg, MD, USA), according to the manufacturer’s instructions. Extracted total RNA was analyzed using the Agilent 2100 RNA 6000 Nano kit (Agilent Technologies, Mississauga, Ontario, Canada) to confirm its quality and quantity. An mRNA library was constructed using the NEN Next Ultra RNA library prep kit (New England Biolabs, Ipswich, MA, USA). An Agilent 2100 (Agilent, Santa Clara, CA, USA) was used for library quality inspection and an Illumina HiSeq TM 4000 (Illumina, Santa Clara, CA, USA) was used for sequencing. mRNA sequencing was performed by Gene Denovo Biotechnology Co. (Guangzhou, China). To ensure data quality, the original data was filtered to reduce data noise before analysis.

Genome-wide mRNA profiling

DEG analysis

Bioinformatics was used to reanalyzed the original data from the sequencing results. Principal component analysis was performed using the R statistical software package with normalized counts to investigate if samples from the same group clustered together. The R-based software package edgeR was used to determine whether there was a significant difference in gene expression levels between the two groups. Genes with a fold change ≥2 and P < 0.05 in a comparison were classified as significant DEGs. The gene ontology (GO) project provides a uniform way to describe the functions of gene products and enable analysis of genomic data with respect to three aspects including molecular function, cellular component, biological process. Functional analysis of DEGs was performed using the GO project (http://www.geneontology.org) [35]. KEGG is a database resource for understanding biological functions of the genes obtained by genome sequencing through intuitional graphics. Pathway enrichment analysis was performed to identify significant pathways involving DEGs using the KEGG database [36].

Gene set enrichment analysis (GSEA)

GSEA is a method to evaluate genomic data at the level of gene sets. It can detect subtle enrichment signals and avoid ignoring genes with insignificant differences. The biological functions of the key NRF2-driven genes were clarified using GSEA (http://software.broadinstitute.org/gsea/index.jsp) [37]. A nominal P < 0.001 and a normalized enrichment score ≥1.00 were chosen as the cutoff criteria.

Co-expression network analysis

To reveal functional associations between proteins, the STRING database was used to construct a PPI network [38]. In the PPI network, each node represents a protein and each edge (between two nodes) represents an interaction between these two proteins. Hub proteins were the nodes with a relatively large number of edges. Within the network analysis, the degree of association is an important factor to determine the relative importance of a gene. Different colors and sizes of nodes were employed to discriminate the degree of associations for one gene with the surrounding nodes. The co-expression networks were constructed using Cytoscape [39].

Quantitative real-time polymerase chain reaction (RT-qPCR) validation of responses in patients with silicosis

RT-qPCR analysis was performed as previously described [18]. Total RNA was extracted from PBMCs of healthy controls and patients with silicosis using the TRIzol reagent (Invitrogen, San Diego, CA, USA) and reverse-transcribed by the Maxima First Strand cDNA Synthesis Kit for RT-qPCR (Thermo Scientific, USA). The relative expression levels of mRNAs encoding
NRF2, Kelch-like ECH-associated protein 1 (KEAP1), glutamate-cysteine ligase modifier subunit (GCLM), TGF-β, mothers against decapentaplegic homolog 3 (SMAD3), SMAD2, mitogen-activated protein kinase 3 (MAPK3), thrombospondin 1 (THBS1), integrin beta-3 (ITGB3), bone morphogenetic protein 4 (BMP4), CD44 molecule (Indian blood group) (CD44), and glyceraldehyde 3-phosphate dehydrogenase (GAPDH) were measured using a LightCycler® 96 (Roche Diagnostics, Mannheim, Germany). The genes were amplified using SYBR Green PCR SuperMix (SYBR High-Sensitivity qPCR SuperMix, Novoprotein, Shanghai, China) and 300 nM of each primer pair. Primers were designed using the Primer Design online software (Sangon Biotech, Shanghai, China) and synthesized by Sangon Biotech (Shanghai, China). Normalized gene expression levels are provided as the ratio between the mean value for the target gene and that for the reference gene (GAPDH) in each sample. All reactions were repeated three times. All RNA samples exhibited a 260/280 ratio > 1.8. The primer sequences used are shown in Table 1.

### Table 1 Gene name and GenBank accession number used in quantitative reverse transcriptase PCR

| Gene          | Accession No. | Gene name                                      | Primer sequence                                      |
|---------------|---------------|-----------------------------------------------|-----------------------------------------------------|
| NRF2          | NC_000002.12  | nuclear factor, erythroid 2 like 2           | 5′-ACGGTATGCAACAGGACATTGAGC-3′ 5′-TTGGCTCTGGACTGGAAACATG-3′ |
| KEAP1         | NC_000019.10  | kelch like ECH associated protein 1          | 5′-ACATCGTCTGGTTTACTACCC-3′ 5′-CAGCATAGATACTAGGTGTG-3′ |
| GCLM          | NC_000009.6   | glutamate-cysteine ligase, modifier subunit | 5′-GGCCACAGTTAAAACCAAATATG-3′ 5′-TCTTTCAACAAAGCAGATTCC-3′ |
| SMAD3         | NC_000015.10  | SMAD family member 3                          | 5′-AGAGAGTGGAGACACGAGTTCTC-3′ 5′-GAAGGTAGGTGGTTTCGGAGATG-3′ |
| SMAD2         | NC_000018.10  | SMAD family member 2                          | 5′-CTTCTCTGGCTGCTGTTGTTA-3′ 5′-AGAGAGTGGAGACACGAGTTCTC-3′ |
| MAPK3         | NC_000016.10  | Mitogen-activated protein kinase 3            | 5′-TCTGCTACTTCCATCAGGAT-3′ 5′-CAGGCGGAAATCACAATCTTA-3′ |
| THBS1         | NC_000015.10  | thrombospondin 1                              | 5′-TTGGACATTTTGGAACCTTCAAGC-3′ 5′-AGAGAGGAAACTCCCTGTG-3′ |
| ITGB3         | NC_000017.11  | Integrin β-3                                  | 5′-TCATCGTGGACCTTCTTATACA-3′ 5′-GTAGACTGTCGGCTTCTTTTATACA-3′ |
| LTPB1         | NC_000002.12  | latent transforming growth factor beta binding protein 1 | 5′-GCCATGAGTGTTAACAAGCAGATCG-3′ 5′-ACAAGGCCCTATACGAGATTG-3′ |
| TGFβ          | NC_000019.10  | transforming growth factor beta 1             | 5′-TGAACAGCCGTCCCTCCTTCTCAGG-3′ 5′-CCGGAAAGCATGAGTGACGCGGCC-3′ |
| BMP4          | NC_000014.9   | Bone morphogenetic protein 4                  | 5′-GTTGGGAAAACCTTTGGGTGAG-3′ 5′-TGGGAGTAGCTGCAGTTCATC-3′ |
| CD44          | NC_000011.10  | CD44 molecule                                  | 5′-TCTGATCTAGGAGACACTCAG-3′ 5′-CAGTCCTGGGTTTCTCATTACG-3′ |
| ALOX12        | NC_000017.11  | arachidonate 12-lipoxygenase, 12S type         | 5′-GATCCGGAGAGAACGAAATAC-3′ 5′-TAGTGTTCAGCAAGGATGT-3′ |
| GAPDH         | NC_000012.12  | glyceraldehyde-3-phosphate dehydrogenase      | 5′-AAGAAGGTGGTTAAAGCAGGCC-3′ 5′-TCCACACCCCTGTGCTGA-3′ |

### Statistics
Routine test data were analyzed using SPSS 19.0 analysis software. Data are reported as the mean ± SD values. Group differences were calculated using a t-test and a Chi-squared test. The Pearson’s correlation coefficients were calculated to test the relationship between the variables. RNA sequencing data between two groups were analyzed using the R statistical software package. P < 0.05 was considered to be significantly different.

### Results
#### Clinical and cytology characteristics of patients with silicosis and healthy volunteers
A total of 92 male patients with silicosis and 87 male healthy volunteers participated in this study; among them, only 77 patients with silicosis and 62 healthy controls fully satisfied the enrollment criteria. Clinical and cytology characteristics of patients with silicosis and healthy volunteers are described in Table 2. There were no statistical differences in age, BMI, blood pressure, and smoking status between the two groups. FVC and FEV1 for respiratory function were significantly lower in the group of patients with silicosis than that of healthy controls. The number of monocytes was significantly higher in patients with silicosis.
silicosis than that of healthy controls through cytology detection, indicating that monocytes may play an important role in cellular responses induced in patients with silicosis (Table 2). The clinical and cytology characteristics in the samples used for the biomolecular detection and Genome-wide mRNA profiling are similar to that of all silicosis patients and healthy controls (Table 2).

**Expression of NRF2 and NRF2-dependent antioxidative genes in PBMCs from patients with silicosis**

To investigate the potential role of NRF2 in PBMCs from patients with silicosis, the expression of NRF2 was examined using a confocal laser scanning microscope. Notably, the expression of NRF2 was significantly higher in patients with silicosis than in healthy controls, and it is showing a similar tendency in the lymphocytes and the monocytes (Fig. 1A). Semi-quantitative analysis revealed a significantly higher NRF2 fluorescence level in patients with silicosis compared to that in healthy controls (Fig. 1B). Moreover, RT-qPCR analysis showed that the expression of NRF2, KEAP1 and GCLM mRNA in PBMCs from patients with silicosis was higher than that in healthy controls (Fig. 1C). Accordingly, the protein expression in PBMCs from patients with silicosis were measured by western blotting. The results of western bolt showed the protein expression of NRF2, KEAP1 and GCLM in PBMCs was higher in patients with silicosis than that of healthy controls (Fig. 1D and E). These results showed that the expression levels of NRF2 and NRF2-dependent antioxidative genes increased in PBMCs from patients with silicosis.

**Network analysis of the DEGs from transcriptome sequencing** found that the genes in the NRF2 module related to NRF2 involved in the silicosis, which suggests that DEGs associated with NRF2 might affect the pathogenesis of silicosis (Fig. 1F).

**Identification of DEGs in the PBMCs of patients with silicosis**

In total, six samples were included: three samples in the healthy control group and three samples in the group of patients with silicosis. Using commercial RNA preparation kits, total RNA was prepared from PBMC samples and the integrity number (RIN) value of each sample is greater than 8.9.

To identify key pathways and genes in the silicosis, the differentially expressed mRNAs among healthy controls and patients with silicosis were analyzed using the Gene-e software. Three conditions were set for DEG screening: a false discovery rate < 20%, unfolded change > 2 and \( P < 0.05 \). A total of 1158 genes were identified after the analysis, of which, 475 were upregulated and 683 were downregulated (Fig. 2A and B).

### Table 2 Clinical and cytology characteristics of silicosis patients and healthy volunteers

| Characteristics | Overall | Samples for detection of Nrf2-related molecular | Samples for RNA sequencing |
|-----------------|---------|-----------------------------------------------|---------------------------|
|                 | Healthy Controls (n = 62) | Silicosis Patients (n = 77) | Healthy Controls (n = 36) | Silicosis Patients (n = 36) | Healthy Controls (n = 3) | Silicosis Patients (n = 3) |
| Age (years)     | 52.56 ± 6.18 | 51.55 ± 8.38 | 52.19 ± 6.49 | 54.31 ± 7.50 | 52.67 ± 9.07 | 53.67 ± 9.29 |
| BMI (kg/m²)     | 25.01 ± 3.36 | 25.47 ± 3.15 | 25.43 ± 3.34 | 25.62 ± 2.78 | 22.79 ± 2.66 | 27.25 ± 2.93 |
| Smokers/non-Smokers | 32/45 | 28/38 | 16/20 | 15/21 | 0/3 | 0/3 |
| SBP (mmHg)      | 126.13 ± 15.38 | 130.00 ± 14.60 | 123.47 ± 14.63 | 128.19 ± 9.19 | 130.00 ± 10.00 | 125.00 ± 5.00 |
| DBP (mmHg)      | 84.84 ± 9.19 | 85.00 ± 9.42 | 84.03 ± 8.77 | 82.64 ± 6.38 | 85.00 ± 5.00 | 76.67 ± 5.77 |
| Percentage of FVC (%) | 91.05 ± 9.24 | 80.81 ± 23.54# | 93.19 ± 6.68 | 79.97 ± 18.97# | 89.33 ± 4.04 | 66.63 ± 7.51# |
| Percentage of FEV1.0 (%) | 94.05 ± 7.67 | 84.35 ± 19.68g | 90.72 ± 8.70 | 76.18 ± 23.59# | 91.33 ± 5.51 | 67.07 ± 9.64* |
| WBC (×10⁶/L)    | 5.90 ± 1.35 | 6.04 ± 1.28 | 5.88 ± 1.37 | 5.85 ± 1.27 | 5.70 ± 1.05 | 6.59 ± 0.92 |
| Neutrophils (× 10⁷/L) | 3.70 ± 1.02 | 3.40 ± 1.16 | 3.51 ± 0.72 | 3.26 ± 0.87 | 3.77 ± 0.93 | 3.67 ± 0.79 |
| Monocytes (×10⁶/ L) | 0.23 ± 0.09 | 0.47 ± 0.15a | 0.23 ± 0.08 | 0.40 ± 0.10a | 0.20 ± 0.10 | 0.38 ± 0.16a |
| Lymphocytes (×10⁶/L) | 2.07 ± 0.65 | 2.03 ± 0.65 | 1.97 ± 0.63 | 1.99 ± 0.75 | 1.73 ± 0.32 | 2.06 ± 0.67 |
| TGF-β (pg/mL)   | 31.26 ± 3.23 | 44.58 ± 3.42a | 28.85 ± 3.22 | 47.59 ± 4.98* | 25.20 ± 7.56 | 62.54 ± 4.85a |

*Note: Continuous variables are expressed as mean ± SD
BMI body mass index, SBP systolic blood pressure, DBP diastolic blood pressure, FVC forced vital capacity, FEV1.0 forced expiratory volume in one second, WBC white blood cell

*P < 0.05 vs Healthy Controls; # P < 0.01 vs Healthy Controls. P values determined by t-test and Chi-squared test
Key molecules and signaling pathways were screened through annotation of DEGs in the silicosis

GO analysis was performed to explore potential roles of differentially expressed mRNAs in the silicosis. GO analysis showed that the DEGs between patients with healthy controls and patients with silicosis were mainly involved in cellular component (including organelle, extracellular region, extracellular matrix, and extracellular region part), oxidation reduction (such as catalytic and oxidoreductase activity), transcription factor activity (such as transcription factor activity, protein binding, and nucleic acid binding transcription factor activity), regulating metabolic processes, immune system processes, response to stimulus transcription, regulating signal transduction, and biological adhesion (Fig. 2C).

KEGG pathway analysis showed that the DEGs between patients with silicosis and healthy controls were primarily enriched in pathways associated with regulating immune-related pathways (Graft-versus-host disease, Rheumatoid arthritis, Allograft rejection, Systemic lupus erythematosus, Inflammatory bowel disease, Asthma, Human T-cell lymphotropic virus type 1 infection, Th1 and Th2 cell differentiation, and Th17 cell differentiation), regulating metabolic processes (amino acid metabolism, lipid metabolism, and metabolism of cofactors and vitamins), cell growth and death, cellular community, signal transduction (including the cGMP-dependent protein kinase G signaling pathway and the TGF-β signaling pathway), and transport and catabolism processes (cell adhesion molecules signaling pathway and the phagosome signaling pathway) (Fig. 2D).

NRF2-regulated key DEGs in PBMCs from patients with silicosis

Based on the above results, NRF2 played a key role in silicosis. GSEA, a powerful tool to infer biological function, was performed and showed that genes associated with GSH metabolism, the TGF-β signaling pathway, and ECM receptor interaction signaling pathway, which were closely related to silicosis [40], were significantly enriched in the group of patients with silicosis (Fig. 3A). These observations suggested that NRF2 may be closely related to the lymphocyte oxidative stress state in patients with silicosis.

We screened 27 DEGs that genes associated with GSH metabolism, the TGF-β signaling pathway, and ECM receptor interaction signaling pathway from 1158 DEGs (Table 3). To further define the interaction between the screened 27 DEGs, the STRING database was used to construct a PPI network. The PPI network consisted of 37 nodes interacting with 144 edges (Fig. 3B). The PPI database was filtered using a combined score > 0.4 and the following top 10 hub genes were identified: SMAD2 (degree = 15), MAPK3 (degree = 15), THBS1 (degree = 15), SMAD3 (degree = 14), ITGB3 (degree = 13), integrin alpha-V (ITGAV) (degree = 13), von Willebrand factor (VWF) (degree = 13), BMP4 (degree = 13), CD44 (degree = 12), and SMAD7 (degree = 11).

RT-qPCR was performed to determine the expression of NRF2-regulated DEGs in PBMCs from patients with silicosis, and confirm selected RNA-Seq findings. Steady-state mRNA levels of key genes associated with the TGF-β signaling pathway were determined. SMAD2, a downstream molecule of the TGF-β signaling pathway, was significantly higher in PBMCs from patients with silicosis. However, SMAD3 mRNA levels were significantly lower than those in healthy controls. Next, it was confirmed that gene expression of BMP4 and MAPK3 were higher in PBMCs from patients with silicosis. Furthermore, the transcriptional factor related to cell-to-matrix interactions, THBS1, was significantly higher in PBMCs from patients with silicosis. Additionally, ITGB3 was higher and CD44 was significantly lower in PBMCs from patients with silicosis compared to health controls (Fig. 3C). The results of correlation analysis showed that the mRNA expression levels of Nrf2 are significantly positively correlated with that of BMP4, MAPK3, and THBS1, and CD44 is irrelevant with that of CD44 (Fig. 3D).

Discussion

Nuclear factor erythroid 2-related factor 2 (NRF2) is a transcription activator that regulates the expression of target genes by binding antioxidant response elements (AREs) [41]. NRF2 is important for the coordinated upregulation of genes in response to oxidative stress. Silica causes the significant accumulation of ROS and activates the antioxidative protein NRF2 and its downstream proteins in the early stages of exposure to silica [42]. Immune cells with anti-inflammatory properties, especially
Fig. 2 (See legend on next page.)
monocytes/macrophages are probably involved in the regulation of cellular immune on silicosis fibrosis [43–45]. The present study focused on the lymphocyte oxidative stress state and found that the number of monocytes and the expression of NRF2 and NRF2-dependent antioxidative genes was significantly increased in PBMCs from patients with silicosis. Additionally, it was confirmed that respiratory function was significantly lower in the patients with silicosis.

The respiratory dysfunction in patients with silicosis is closely related to the severity of pulmonary fibrosis [46]. The TGF-β pathway can play an important direct inducer role in the process of collagen transcription in the development of silicosis [47], as TGF-βs are overexpressed in fibrosis [48]. TGF-β signaling plays a key role in extracellular matrix remodeling, the epithelial-mesenchymal transition, and cell growth, migration, differentiation in fibrosis [49]. Generally, TGF-β signaling is modulated by the phosphorylation of the cytoplasmic SMAD signaling molecules, which results in their translocation to the nucleus [50]. TGF-β1 signaling molecules play a key role by promoting transdifferentiation the fibroblast into myofibroblasts, which promote collagen synthesis and ECM deposition in the pathology of silicosis [48].

The ECM, complex mixture of structural and functional macromolecules, has an important role in tissue fibrosis and in the maintenance of cell and tissue structure and function [51]. Integrins and other transmembrane molecules mediate specific interactions between cells and the ECM [52, 53]. These interactions have the direct and indirect effects of cellular activities, lead to adhesion, proliferation, apoptosis, migration, and differentiation. In addition, integrins act as ‘mechanoreceptors’ that would provide a specific physical link between the cytoskeleton and the ECM [53].

In the present study, the differential expression of mRNAs in the silicosis was identified using RNA-Seq analysis. A total of 1158 dysregulated mRNAs were identified in PBMCs from patients with silicosis, including 475 upregulated and 683 downregulated mRNAs. GO analysis revealed that the functions of dysregulated mRNAs in PBMCs from patients with silicosis were related to the ECM, catalytic activity, oxidoreductase activity, transcription factor activity, metabolic processes, immune system processes, response to stimulus transcription, and biological adhesion. In KEGG pathway analysis, the dysregulated mRNAs were involved in regulating immune-related pathways, regulating metabolic processes, cellular community, cell adhesion molecules signaling pathway and the phagosome signaling pathway. Therefore, it is possible that the dysregulated mRNAs in these processes are involved in the pathogenesis of silicosis.

To confirm the results obtained by RNA-Seq analysis, seven differentially expressed mRNAs were selected to verify their expression in PBMCs from patients with silicosis using RT-qPCR. The results indicated that the expression of SMAD2, MAPK3, THBS1, ITGB3, and BMP4 was increased, while the expression of SMAD3 and CD44 was significantly low expression in PBMCs from patients with silicosis, indicating consistent results with the RNA-Seq data. The TGF-β family of cytokines signals through receptor serine/threonine kinases to control cell behavior and fate [54]. These signals are propagated through the transcription factors SMAD2 and SMAD3 downstream of TGFβ. SMAD2 and SMAD3 belong to the SMAD protein family that mediate multiple signaling pathways as transcriptional modulators [55]. These findings indicate that dysregulation of the TGF-β1-SMAD signaling pathway may play an important role in the pathological process of silicosis, and are consistent with findings in animal experiments [56, 57]. SMAD proteins mediate signaling of TGF-β through its interaction with the SMAD anchor for receptor activation (SARA) protein [58]. In response to a TGF-β signal, SMAD proteins are phosphorylated by TGF-β receptors [59]. This study revealed that BMP4 was a differential gene hub in PBMCs from patients with silicosis. BMP4 encodes a secreted ligand of the TGF-β superfamily of proteins and activates SMAD family transcription factors that regulate gene expression [60]. These results showed that SMAD2 and SMAD3 had different trends in the two groups. Recently, SMAD2 and SMAD3 have shown different roles in the TGF-β signaling pathway during embryonic development [55]. The cell experiments revealed that BMP7 is associated with inhibiting silica-induced fibrosis through activated BMP7/SMAD and suppressed TGF-β/SMAD pathways [61, 62]. To date, BMP4 in the BMP protein family has not been reported in silicosis.

MAPK3 is especially involved in activation towards microtubule-associated protein-2 and the control of cell survival, proliferation and differentiation [63]. In the present study, MAPK3 (ERK1) was significantly increased in PBMCs from patients with silicosis compared
Fig. 3 (See legend on next page.)
to that in the control group. Dysregulation of MAPK3 plays a significant role in the pathological processes of silicosis [16]. Crystal compounds in silicosis activate ROS, which activate the inflammasome through MAPK3 [16, 64]. In agreement with these findings, activation of MAPK3 (ERK1) and NF-κB in PBMCs is reported during oxidative stress [65].

The transcription factor thrombospondin-1 (THBS1) is an adhesive glycoprotein that mediates cell-to-cell and cell-to-matrix interactions. THBS1 binds to cell surface receptors, including fibrinogen, fibronectin, laminin, type

**Table 3** The screened 27 DEGs associated with lymphocyte oxidative stress state by GSEA analysis of NRF2. 27 key DEGs were identified from 1158 DEGs, which belong to the gene sets from GSH metabolism, the TGF-β signaling pathway, and ECM receptor interaction signaling pathway.

| Gene Symbol | Description | Log2FC | t | P.Value |
|-------------|-------------|--------|---|---------|
| GCLC        | glutamate-cysteine ligase catalytic subunit | 0.9761 | 3.5071 | 0.0130 |
| THBS1       | thrombospondin 1 | 1.8189 | 3.3611 | 0.0155 |
| ITGA2B      | integrin subunit alpha 2b | 1.4937 | 7.8035 | 0.0003 |
| BMP6        | bone morphogenetic protein 6 | 1.4981 | 3.6512 | 0.0110 |
| ZFYVE9      | zinc finger FYVE-type containing 9 | −1.2890 | −2.8443 | 0.0299 |
| ITGA2       | integrin subunit alpha 2 | 1.1618 | 2.6027 | 0.0410 |
| GP9         | glycoprotein IX platelet | 1.2380 | 4.6464 | 0.0036 |
| ITGA4       | integrin subunit alpha 4 | −1.7863 | −3.3991 | 0.0148 |
| CD44        | CD44 molecule | −1.2415 | −4.3524 | 0.0050 |
| SMURF1      | SMAD specific E3 ubiquitin protein ligase 1 | −0.6038 | −2.5541 | 0.0438 |
| TGFR2       | transforming growth factor beta 2 | 1.5877 | 2.6729 | 0.0374 |
| GSTM4       | glutathione S-transferase mu 4 | 1.8561 | 2.4830 | 0.0482 |
| GLCM        | glutamate-cysteine ligase modifier subunit | 0.6821 | 2.5854 | 0.0420 |
| SDC4        | syndecan 4 | 0.6770 | 2.7085 | 0.0357 |
| AGRN        | agrin | 0.9977 | 2.8008 | 0.0316 |
| MAPK3       | mitogen-activated protein kinase 3 | 0.8754 | 2.5837 | 0.0421 |
| ODC1        | ornithine decarboxylase 1 | 0.7449 | 2.9195 | 0.0271 |
| ACVR1       | activin A receptor type 1 | 1.8224 | 2.9468 | 0.0261 |
| GP6         | glycoprotein VI platelet | 1.2393 | 3.1317 | 0.0207 |
| LTB1P       | latent transforming growth factor beta binding protein 1 | 1.6414 | 4.7492 | 0.0033 |
| BMPR1B      | bone morphogenetic protein receptor type 1B | −2.4111 | −2.7113 | 0.0355 |
| GSR         | glutathione-disulfide reductase | 0.7830 | 2.6736 | 0.0374 |
| SP1         | SP1 transcription factor | 1.1197 | 2.5086 | 0.0465 |
| ITG4B       | integrin subunit beta 3 | 1.1695 | 4.6047 | 0.0383 |
| GP1BA       | glycoprotein IB platelet alpha subunit | 0.8702 | 2.7218 | 0.0351 |
| MGST3       | microsomal glutathione S-transferase 3 | 0.7892 | 2.9403 | 0.0264 |
| SRM         | spermidine synthase | −1.2838 | −3.3023 | 0.0167 |
V collagen and integrins, such as ITGB3 [66]. These studies suggest that THBS1 in human peripheral blood lymphocytes is involved in the regulation of pesticide-induced immune dysfunction [67].

The protein encoded by CD44 is a cell-surface glycoprotein involved in cell-cell interactions, cell migration and adhesion [68]. It also interacts with other ligands, such as matrix metalloproteinases, and collagens [69]. CD44 blockade alleviates silica-induced fibrosis and improves pulmonary function in vivo [70].

These findings identify activation of the NRF2-mediated oxidative stress in PBMCs as a key contributor to the pathogenesis of patients with silicosis. Nonetheless, it has to be underlined that only the patients with silicosis stage I are included in this study and our results must be confirmed in larger trial. The other limitation of this study is that the present study remains unclear that changes specific to different cell types present in PBMCs. It will be necessary to study these patients further by establishing an in vitro model of PBMCs exposed to silica particles.

Conclusions

Genome-wide mRNA profiling from PBMCs in patients with silicosis is identified. Overexpression of NRF2 and related signaling molecules are regulating the oxidative stress state in PBMCs from patients with silicosis. Therefore, NRF2 might serve as a novel preventive and therapeutic candidate for silicosis. The novel information provided by this study contribute to understand of the oxidative stress mechanism of silicosis.

Abbreviations

AREs: Antioxidant response elements; BMI: Body mass index; BMP4: Bone morphogenetic protein 4; CAT: Catalase; DEGs: Differentially expressed genes; FEV1: Forced expiratory volume in one second; FVC: Forced vital capacity; GAPDH: Glyceraldehyde 3-phosphate dehydrogenase; GCLM: Glutamate-cysteine ligase modifier subunit; GO: Gene ontology; GPXs: Glutathione peroxidases; GR: Glutathione reductase; GSEA: Gene set enrichment analysis; GSH: Glutathione; ITGB3: Integrin beta-3; ITGAV: Integrin alpha-V; KEAP1: Kelch-like ECH-associated protein 1; KEGG: Kyoto Encyclopedia of Genes and Genomes; MAPK3: Mitogen-activated protein kinase 3; NK: Natural killer; NRF2: Nuclear factor erythroid 2-related factor 2; PBMCs: Peripheral blood mononuclear cells; PPI: Protein-protein interaction; RNA-Seq: RNA sequencing; ROS: Reactive oxygen species; SMAD3: Mothers against decapentaplegic homolog 3; SODs: Superoxide dismutases; STRING: Search Tool for the Retrieval of Interacting Genes/Proteins; THBS1: Thrombospondin 1; TGF: Transforming growth factor; vWF: von Willebrand factor

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Authors’ contributions

Conceptualization, Yingzheng Zhao and Sanqiao Yao; Methodology, Guangcui Xu, Haibin Li, Meiyu Chang, Cheng Xiong and Yunchun Liu; Investigation, Yingzheng Zhao, Guangcui Xu, Haibin Li, Meiyu Chang, Cheng Xiong, Yunchun Li, Yi Guan, and Yingjun Tao; Software, Meiyu Chang and Yingzheng Zhao; validation, Yingzheng Zhao and Guangcui Xu; data curation, Guangcui Xu, Haibin Li, Yingjun Tao and Meiyu Chang; writing—original draft preparation, Yingzheng Zhao and Guangcui Xu; writing—review and editing, Sanqiao Yao and Yingzheng Zhao; supervision, Yingzheng Zhao and Haibin Li; project administration, Sanqiao Yao; Manuscript submission, Yingzheng Zhao. All authors have read and agreed to the published version of the manuscript.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Declarations

Ethics approval and consent to participate

All subjects signed an informed consent before being enrolled. The study was approved by the Ethics Committee of the Xinxiang Medical University (protocol number XYLL-2017086).

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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