Research Article

Abnormal Peripheral Neutrophil Transcriptome in Newly Diagnosed Type 2 Diabetes Patients

Qiuqiu Lin, Wenzhi Zhou, Yanfei Wang, Juan Huang, Xiaoyan Hui, Zhiguang Zhou, and Yang Xiao

1Department of Metabolism & Endocrinology, The Second Xiangya Hospital, Central South University, Changsha, Hunan 410011, China
2National Clinical Research Center for Metabolic Diseases, Changsha, Hunan, China
3Key Laboratory of Diabetes Immunology (Central South University), Ministry of Education, Changsha, Hunan 410011, China
4State Key Laboratory of Pharmaceutical Biotechnology, The University of Hong Kong, Hong Kong, China

Correspondence should be addressed to Yang Xiao; xiaoyang29@csu.edu.cn

Received 26 December 2019; Revised 15 February 2020; Accepted 21 February 2020; Published 23 April 2020

Aim.
There are increasing evidence demonstrating that neutrophil-mediated inflammation plays a role in the etiology of type 2 diabetes. However, the molecular mechanisms by which neutrophils contribute to type 2 diabetes remain largely unknown. The aim of the present work was to identify possible changes in circulating neutrophils to better elucidate neutrophil involvement in human type 2 diabetes.

Methods.
Patients newly diagnosed with type 2 diabetes (n = 5) and age- and sex-matched healthy controls (n = 5) were recruited. Neutrophils were purified from type 2 diabetes patients and controls, and RNA sequencing (RNA-seq) was used for comprehensive transcriptome analysis. Differentially expressed genes (DEGs) were screened, and Gene Ontology (GO) and KEGG pathway enrichment analyses were performed. Real-time polymerase chain reaction (qPCR) was used for validation in external samples of type 2 diabetes patients (n = 8) and healthy controls (n = 8).

Results.
Gene expression analysis showed that, compared with neutrophils from healthy controls, there were 1990 upregulated DEGs and 1314 downregulated DEGs in neutrophils from type 2 diabetes patients. GO analysis demonstrated that the DEGs were mainly involved in myeloid leukocyte activation, T cell activation, adaptive immunity, and cytokine production. The top 20 enriched KEGG pathways included the cytokine-cytokine receptor interaction pathway, NF-κB signaling pathway, cell adhesion molecules, and chemokine signaling pathway. Furthermore, qPCR of genes related to neutrophil activation revealed that the expression of SELL, SELP, CXCR1, and S100A8 was significantly increased in neutrophils from type 2 diabetes patients compared with that in neutrophils from controls.

Conclusions.
Our study reveals an abnormal activation of circulating neutrophils at the transcriptome level in type 2 diabetes patients. These findings suggest a potential involvement of neutrophil dysfunction in the pathologic process of type 2 diabetes and provide insight into potential therapeutic targets for type 2 diabetes.

1. Introduction

Low-grade inflammation is a common component in type 2 diabetes, particularly in the development of obesity-related insulin resistance [1]. Neutrophils are the most abundant type of white blood cell and are reported as active players in inflammation of obesity-related insulin resistance [2]. Additionally, neutrophil count, a marker of subclinical inflammation, has been shown to significantly increase in type 2 diabetes compared with healthy subjects [3–5]. Neutrophil-lymphocyte ratio significantly increases in prediabetes and diabetes and may be a predictive marker for prediabetes and diabetes mellitus [6]. Furthermore, several large-scale prospective studies demonstrated that the neutrophil count could be used as a predictor of the incidence of type 2 diabetes [7], suggesting the potential role of neutrophils in the development of type 2 diabetes.

Neutrophils eliminate extracellular pathogens by multiple strategies, including phagocytosis, degranulation to release lytic enzymes, and neutrophil extracellular traps (NETs),...
which are formed through a unique cell death process that is clearly different from both apoptosis and necrosis, termed “NETosis” [8–10]. However, improper activation of neutrophils may lead to tissue damage during exaggerated inflammatory responses [11]. Neutrophils from patients with type 2 diabetes reportedly produce more superoxide and cytokines [12, 13], and neutrophils from type 2 diabetes patients are more susceptible than those from healthy controls to PMA-induced NETosis [14]. Neutrophil serine proteases, which are crucial components of NET, including neutrophil elastase (NE) and proteinase 3 (PR3), have been shown to participate in the initiation of insulin resistance and type 2 diabetes [2, 15]. NE treatment elicits insulin resistance and glucose intolerance in mice, while neutrophil elastase deficiency results in improved tissue inflammation with less macrophage infiltration in adipose tissues in high-fat diet-induced obese mice [2]. Injection of recombinant PR3 alone is sufficient to induce hyperglycemia in mice, and inhibition of PR3 activity leads to an increase in glucose clearance [15]. However, the precise mechanism by which neutrophils induce type 2 diabetes remains elusive. Therefore, in this study, we aimed to identify the transcriptomic changes in circulating neutrophils from type 2 diabetes by RNA sequencing (RNA-seq) to better elucidate neutrophil involvement in type 2 diabetes.

2. Materials and Methods

2.1. Subjects. Thirteen patients with type 2 diabetes whose disease duration was less than one year were enrolled from the Second Xiangya Hospital, Central South University. The diagnosis of diabetes was based on the World Health Organization (WHO) criteria (1999). The exclusion criteria for type 2 diabetes were as follows: (1) acute infection, trauma, or surgery within one month; (2) use of antibiotics, glucocorticoids, or other immune regulators within one month; (3) severe cardiocerebrovascular, liver, kidney, or malignant disease; (4) pregnancy or lactation; (4) autoimmune diseases, such as hyperthyroidism; and (5) other types of diabetes. Thirteen gender- and age-matched controls were recruited and exhibited euglycemia using a 75 g glucose tolerance test. The diagnosis of diabetes was based on the World Health Organization (WHO) criteria (1999). The exclusion criteria for type 2 diabetes patients and healthy control subjects by using the DEG-seq algorithms [19]. An adjusted P value not greater than 0.001 and an absolute value of the log2 ratio greater than 1 indicated significant gene expression differences. All the samples were hierarchically clustered by the expression level of all genes. According to the DEGs, we next performed Gene Ontology (GO) classification including molecular biological function, cellular component, and biological process. With the DEGs, we performed KEGG pathway classification and functional enrichment by using phyper, which is a function of R package. We calculated the false discovery rate (FDR) for each P value, and the terms for which the FDR was not greater than 0.01 were defined as significantly enriched.

2.2. Measurements. Height and weight, waist circumference, hip circumference, blood pressure, body mass index (BMI), and weight/height ratio (WHR) were obtained for all patients. Fasting venous blood samples were obtained at 8:00 am. The following biochemical parameters were assessed in fasting venous blood samples: fasting glucose, cholesterol (TC), triglycerides (TGs), fasting blood glucose, fasting C-peptide (FCP), and hemoglobin A1C (HbA1c) levels. Circulating cell counts were analyzed by the automated hematology analyzer Sysmex XE-2100. Plasma glucose was measured by a Hitachi 7170 analyzer (Boehringer Mannheim, Germany). Serum cholesterol and TG levels were measured enzymatically. Serum levels of C-peptide were assessed by the Advia Centaur System (Siemens, Munich, Germany). HbA1c was determined by liquid chromatography (Bio-Rad Laboratories, Hercules, CA).

2.3. Neutrophil Isolation and RNA Extraction. Human neutrophils were isolated from venous blood of patients and healthy controls by density gradient centrifugation using Ficoll-Paque Plus according to the manufacturer’s protocol and then followed by positive magnetic separation for further purification using human CD16 Microbeads (Miltenyi Biotech). The cells were dissolved in TRIzol (Roche, America) in a volume of 5–10 × 10⁶ cells/mL, followed by storage at -80°C. Total RNA was extracted, and the concentration and purity of RNA were tested on a NanoDrop spectrophotometer, followed by reverse transcription using the High-Capacity cDNA Reverse Transcription Kit (Thermo Fisher Scientific, USA).

2.4. RNA-seq. In this study, we sequenced 5 samples from type 2 diabetes patients and 5 samples from controls on the BGISEQ-500 platform. A total of 19,718 genes were detected, averaging approximately 24.04 million reads per sample. Before downstream analyses, the raw sequencing reads, such as low-quality, polluted, and unknown base (N) reads, were filtered, followed by mapping of the clean reads to the reference genome using HISAT [16] and Bowtie2 [17]. The gene expression was calculated using a software package called RSEM [18]. The average mapping ratio to the reference genome was 92.93% (see Supplementary Tables 1 and 2), and the average mapping ratio to genes was 65.73%. According to the gene expression level, we identified differentially expressed genes (DEGs) between type 2 diabetes patients and healthy control subjects by using the DEG-seq algorithms [19]. An adjusted P value not greater than 0.001 and an absolute value of the log2 ratio greater than 1 indicated significant gene expression differences. All the samples were hierarchically clustered by the expression level of all genes. According to the DEGs, we next performed Gene Ontology (GO) classification including molecular biological function, cellular component, and biological process. With the DEGs, we performed KEGG pathway classification and functional enrichment by using phyper, which is a function of R package. We calculated the false discovery rate (FDR) for each P value, and the terms for which the FDR was not greater than 0.01 were defined as significantly enriched.

2.5. Real-Time PCR Analysis. Real-time quantitative (qPCR) was performed with Power SYBR green PCR master mix (Go Taq® qPCR, Promega Corporation, USA) on the Mini-Opticon real-time PCR detection system (ViiA™ 7 Real-Time PCR System containing the Optiflex™ Optics System). All the primers used for qPCR were designed and synthesized by TSINGKE (TSINGKE Biological Technology, China). The expression of each gene was quantified as a fold change against β-actin by 2−ΔΔCt method. Primer sequences of genes are shown in Table 1.

2.6. Statistical Analysis. Data are presented as the mean ± SD or median with interquartile range. Student’s t-test was used to identify the differentially expressed groups using GraphPad.
3. Results

3.1. Characteristics of Participants. The anthropometric characteristics of the discovery group including 5 type 2 diabetes patients and 5 control subjects and the validation group with 8 type 2 diabetes patients and 8 control subjects are presented in Tables 2 and 3, respectively. In the discovery group, HbA1c, postprandial glucose, fasting C-peptide, systolic pressure, and diastolic blood pressure were higher in type 2 diabetes patients than in healthy controls (*P < 0.05), and no differences were found with respect to BMI, WHR, fasting glucose, LDL-C, and TC between type 2 diabetes patients and control subjects.

| Gene | Sense (5' → 3') | Antisense (3' → 5') |
|------|----------------|-------------------|
| CXCR1 | TCAAGTGCCCTCTAGCTGTT | TGATCTAACTGGAAGCCAGCGC |
| CXCR2 | TCTGCTAGAGGTCTGACTAC | CTGGGCTTTTCACCTGTTAGGA |
| SELL | TCTGTGTGTAGTTCCGGCAC | CCACCCACGTCATATTCCT |
| SELP | CCCAGTGTGTAAAGCTATTTCGT | GCTCCTCTAGCATGAAACCT |
| PECAM1 | TTTTGGCGTCAGTGGTGTC | CTTGAACAGACGAAAGGTCAGCA |
| S100A8 | AGACCTGAAAGTTCTGTTTTCA | AGGACATCTCGTTCTTACAGGA |
| S100A11 | GCATCGGATCCCTGATTGCTT | AGGGTCCTTCTGTTCTTTCAG |
| S100A12 | ATTCCTGTGCATTGGGGTTA | TTGCAAAATGCCCCCTTCCAG |
| SLC2A3 | CGTGGAGAAAACTTGCTGCTG | TCAGAGCTGGGGTGACCCTCAG |
| HSPA1 | CGCAAGTGCATCTTTGAC | TCGGTTGTCTTGGCTTTCAG |
| BST2 | TGTCGCAATGTCACCCATCT | AGCCATTAGGGCCATACAG |
| CPNE3 | GACTCCCACGAAACTCAGGT | AACATTCAGCGCCACTCAG |
| β-Actin | GCATCCCCAAAGTTCCT | AGGAGTGGGACCTTCTCT |

Data are expressed by mean ± SD or median (25-75th percentile). BMI: body mass index; WHR: waist to hip ratio; DBP: diastolic blood pressure; SBP: systolic blood pressure; TG: triglycerides; HDL-C: high-density lipoprotein cholesterol; LDL-C: low-density lipoprotein cholesterol; *compared with HC. **P < 0.01 compared with HC.
and control subjects. For the validation set, HbA1c and fasting and postprandial glucose were higher in type 2 diabetes patients than in healthy controls ($P < 0.05$).

3.2. Bioinformatics Analysis. RNA-seq analysis of neutrophils was performed and then DEGs were identified. Compared with neutrophils from the healthy controls, there were 1990 upregulated DEGs and 1314 downregulated DEGs in neutrophils from type 2 diabetes patients. The DEG-seq algorithm was used to detect the DEGs between the two groups, and the significance of the difference was established based on the filtering criteria: fold change $\geq 2$ or fold change $\leq 0.5$ and adjusted $P$ value $\leq 0.001$ (Figures 1 and 2 and Supplementary Data 1).

3.3. Gene Ontology (GO) Classification. Based on these DEGs, a GO classification and a functional enrichment analysis were performed to determine the molecular functions (MM), cellular components (CC), and biological processes (BP) involved in the proteins encoded by these genes. As expected, most are important to neutrophil functioning. We discovered that GO categories of the top 8 upregulated biological functions were myeloid leukocyte activation ($log P = −22.62$), T cell activation ($log P = −14.94$), adaptive immune system ($log P = −14.48$), cytokine production ($log P = −11.59$), immune response-regulating signaling pathway ($log P = −11.40$), cytokine-mediated signaling pathway ($log P = −10.00$), immunoregulatory interactions between a lymphoid and a nonlymphoid cell ($log P = −10.00$), and immune response-regulating pathway ($log P = −9.09$) (Figure 3(a) and Supplementary Data 3). Among the downregulated GO categories, there was a close relationship with the response to molecules of bacterial origin, the biological process of the inflammatory response, interleukin- (IL-) 10 signaling, regulation of cytokine production, cytokine-cytokine receptor interaction, cellular response to lipids, and regulation of cell adhesion (Figure 3(b) and Supplementary Data 3).

3.4. KEGG Pathway Enrichment Analysis. KEGG pathway enrichment analysis of the important DEGs suggested that a wide range of biological pathways were altered in neutrophils from type 2 diabetes patients compared with controls. $P$ values were used to describe the significance level of pathway enrichment. There were 318 main pathways identified, and the top 20 differential pathways were primarily involved in the cytokine-cytokine receptor interactions ($P = 3.46 \times 10^{-10}$), NF-κB signaling ($P = 4.12 \times 10^{-9}$), tumor necrosis factor (TNF) signaling ($P = 6.23 \times 10^{-5}$), cell adhesion molecule (CAM) signaling ($P = 1.66 \times 10^{-5}$), Toll-like receptor signaling ($P = 2.82 \times 10^{-3}$), and chemokine signaling ($P = 3.25 \times 10^{-3}$) (Figure 4 and Supplementary Data 2). In the above most enriched pathways, especially cytokine-cytokine receptor interactions and cell adhesion molecule (CAM) signaling, the majority of the genes associated with the two pathways were upregulated in type 2 diabetes, such

### Table 3: Clinical and biochemical characteristics of the study participants for validation.

|                        | HC ($n = 8$) | T2D ($n = 8$) | $P$ value |
|------------------------|-------------|--------------|-----------|
| Sex (male/female)      | 8 (6/2)     | 8 (6/2)      | 1.000     |
| Age (years)            | 44.25 ± 8.41| 47.63 ± 10.74| 0.496     |
| BMI (kg/m²)            | 22.40 ± 2.10| 23.82 ± 2.83 | 0.273     |
| WHR                    | 0.85 ± 0.08 | 0.88 ± 0.05  | 0.388     |
| DBP (mmHg)             | 75.85 ± 7.85| 79.88 ± 8.01 | 0.316     |
| SBP (mmHg)             | 121.00 ± 11.20| 118.50 ± 16.45 | 0.733     |
| TG (mmol/L)            | 1.61 ± 0.81 | 1.54 ± 1.58  | 0.903     |
| TC (mmol/L)            | 4.77 (4.05–4.93)| 3.96 (2.50–4.56) | 0.050     |
| HDL-C (mmol/L)         | 1.14 ± 0.48 | 1.51 ± 0.92  | 0.332     |
| LDL-C (mmol/L)         | 2.75 ± 0.36 | 2.07 ± 0.89  | 0.065     |
| HbA1c (%)              | 5.30 (5.20–5.55) | 6.50 (6.05–7.70)*** | <0.001 |
| Fasting BS (mmol/L)    | 4.61 (4.21–5.11)| 5.97 (5.14–7.65)** | 0.002     |
| 2 h postprandial BS (mmol/L) | 4.88 ± 1.64 | 12.58±5.17*** | <0.001 |
| Fasting C-peptide (pmol/L) | 418.71 ± 90.49 | 369.44 ± 141.28 | 0.420     |
| 2 h postprandial C-peptide (pmol/L) | 1598.80 ± 711.38 | 1229.03 ± 548.98 | 0.264     |
| White cell count (10⁹/L) | 5.97 ± 0.81 | 5.73 ± 0.42  | 0.411     |
| Lymphoid cell count (10⁹/L) | 1.97 ± 0.54 | 1.83 ± 0.83  | 0.695     |
| Neutrophil count (10⁹/L) | 3.41 (3.12–3.50)| 3.42 (2.88–3.68) | 0.878     |
| Mononuclear count (10⁹/L) | 0.40 ± 0.08 | 0.30 ± 0.07  | 0.095     |

Data are expressed by mean ± SD or median (25-75th percentile). BMI: body mass index; WHR: waist to hip ratio; DBP: diastolic blood pressure; SBP: systolic blood pressure; TG: triglycerides; HDL-C: high-density lipoprotein cholesterol; LDL-C: low-density lipoprotein cholesterol; *compared by the Mann-Whitney U test. *$P < 0.05$ compared with HC. **$P < 0.01$ compared with HC. ***$P < 0.001$ compared with HC.
as chemokine (C-X-C motif) ligand (CXCL7, CXCL4), C-X-C chemokine receptor type (CXCR1, CXCR2), also named Interleukin 8 Receptor (IL8RA, IL8RB), as well as cell adhesion molecules integrin subunit alpha M (ITGAM), L-selectin (SELL), P-selectin (SELP), plate endothelial cell adhesion molecule-1 (PECAM1), and P-selection glycoprotein ligand-1 (PSGL-1). Pathway maps of cytokine-cytokine receptor interaction, cell adhesion molecules, and leukocyte transendothelial migration are shown in Figure 5.

3.5. Real-Time Quantitative PCR. We next expanded the analysis of the purified neutrophils. In the RNA-seq results, genes were directly with neutrophil activation, like the expression of adhesion molecules like SELL, SELP, PECAM1, and related ligands or receptors, such as CXCR1, CXCR2, calcium-binding protein (S100A8, S100A11, and S100A12), bone marrow stromal cell antigen 2 (BST2), heat shock protein family A member 1A (HSPA1), and Copine 3 (CPNE3) were increased in type 2 diabetes (Figure 6). For the validation group, we performed qPCR for these genes. As adhesion molecules, the expression of SELL and SELP in neutrophils from patients was increased compared with that in neutrophils from healthy controls ($P = 0.030$ and $P = 0.003$). In addition, CXCR1 expression was higher in neutrophils from type 2 diabetes patients than in those from healthy controls ($P = 0.022$). The levels of S100A8, which is derived mainly from neutrophils regarded as a mediator of inflammation, were higher in neutrophils from type 2 diabetes patients than in those from healthy controls ($P = 0.019$). However, comparable levels of PECAM1, CXCR2, SLC2A3, BST2, S100A11, S100A12, and CPNE3 were found in neutrophils from diabetes patients and controls ($P > 0.05$).

4. Discussion

Neutrophils are the first-line immune cells involved in inflammation, and circulating neutrophil counts are moderately increased in type 2 diabetes [3, 7]. However, the role of neutrophils in the pathogenesis of type 2 diabetes is largely unknown. Our study is the first to investigate the presence of DEGs and the biological functions associated with these genes in neutrophils from type 2 diabetes and healthy individuals. The study has shown that neutrophils from patients with type 2 diabetes presented increased neutrophil activation, responses to chemokines and neutrophil transendothelial cell migration at the transcriptome level.
In the current study, we used an RNA-seq dataset to assess the neutrophil gene expression changes at the transcriptome level between patients with type 2 diabetes and healthy controls. A total of 3304 DEGs were identified, including 1990 upregulated genes and 1314 downregulated genes. According to the GO analysis, myeloid leukocyte activation was the most significant among the top 20 enriched terms, which is consistent with previous studies showing increased leukocyte activation in patients with insulin resistance and type 2 diabetes [20, 21], indicating a potential role of leukocyte activation in the pathogenesis of type 2 diabetes.

Our data showed that the process of leukocyte-endothelial adhesion was activated in the analyzed neutrophils as the pathway map of leukocyte transendothelial migration shown. The adhesion genes ITGAM (CD11b), PECAM1, SELL, and SELP and the receptor PSGL-1 were upregulated in type 2 diabetes, and the mRNA levels of SELL and SELP were also increased in neutrophils from patients with type 2 diabetes, as assessed by real-time PCR; however, there was no difference in PECAM1 levels in neutrophils between patients and healthy controls. Selectins play unique roles in neutrophil recruitment by mediating recognition and adhesion between leukocytes and vascular endothelial cells. Mice lacking in L-selectin and PSGL-1 show worse neutrophil recruitment into the inflamed peritoneum than PSGL-1 knock-out mice. L-selectin (CD62L), which is expressed by most leukocytes, is involved in neutrophil trafficking [22]. P-selectin, encoded by SELP, can capture leukocytes from the circulation to the vessel wall, leading to the rolling of neutrophils [23], and polymorphisms of SELP are associated with vascular risk of type 2 diabetes [24]. The KEGG pathway analysis demonstrated that neutrophil rolling function, neutrophil activation, and adhesion were dysregulated in patients with type 2 diabetes compared with healthy controls. However, the literature also shows that CD62L is decreased in peripheral blood neutrophils in patients with diabetic microangiopathy, as assessed by flow cytometry [25]. This difference may be explained by differences between transcriptome and protein levels.

The migration of neutrophils across endothelial cells to the vascular wall is an essential step in tissue damage and the inflammatory response. Adhesion molecules mediate the adhesion of neutrophils to vascular endothelial cells. Several studies have confirmed the activation of neutrophils and elevated CD11b expression in diabetic patients [20, 21]. Moreover, it has been shown that the neutrophil-secreted enzyme NE (neutrophil elastase) impairs insulin signaling and increases insulin resistance. Conversely, obese mice without NE fed a high-fat diet showed

![Gene Ontology (GO) analysis with the GO enrichment for differentially expressed genes (DEGs) between type 2 diabetes patients and healthy controls. DEGs of T2D compared with healthy controls are defined as levels of fold change ≥ 2 for upregulated genes and ≤0.5 for downregulated genes. (a) presents the top 8 GO categories for upregulated genes. (b) presents the top 8 GO categories for downregulated genes.](image)
improvement in insulin sensitivity [2]. Furthermore, NE has been detected in the plasma of type 2 diabetes patients who had elevated levels of glycated hemoglobin [26].

CXCR1 and CXCR2 widely exit on the cell surface of neutrophils [27]. Neutrophils expressing CXCR1/2 can be recruited to the pancreas by murine β cells, and macrophages produce C-X-C motif ligand 2 (CXCL2) in autoimmune diabetes [28], which plays a vital role in the early stages of diabetes. The expression of CXCR1/2 decreased after bariatric surgery in female adipose tissue [29]. Moreover, CXCR2-deficient mice are resistant to diet-induced insulin resistance and diabetes, mainly because CXCL5 blocks insulin signaling in muscle by activating the muscle Jak/STAT/SOCS pathway through the CXCR2 receptor [30]. In accordance with this finding, our RNA-seq analysis revealed that the CXCR1 and CXCR2 genes were significantly upregulated in patients with type 2 diabetes compared with healthy controls. Consistent with this result, the real-time PCR results showed that CXCR1 mRNA levels were significantly increased and that there was a tendency toward increased CXCR2 mRNA levels in type 2 diabetes.

Type 2 diabetes is associated with worse outcomes and mortality caused by infection [31] due to impaired innate immune functions, including phagocytosis, cytokine and reactive oxygen species (ROS) production, bactericidal activity, and chemotaxis [32]. In our study, the GO analysis showed that the diabetic neutrophil response to molecules of bacterial origin, such as LPS, and inflammatory response were decreased. In addition, the gene expression of cytokines/chemokines, such as CXCL2, CXCL3, CXCL5, and

**Figure 4:** The top 20 KEGG pathways on the basis of all the differentially expressed genes (DEGs) between neutrophils from patients with type 2 diabetes and healthy controls. DEGs of T2D compared with healthy controls are defined as levels of fold change ≥ 2 for upregulated genes and ≤ 0.5 for downregulated genes. The X axis represents enrichment factor. The Y axis represents pathway name. The color indicates the q value (high: white, low: blue), and the lower q value indicates the more significant enrichment. Point size indicates DEG number (the bigger dots refer to larger amount). Rich factor refers to the value of enrichment factor, which is the quotient of foreground value (the number of DEGs) and background value (total gene amount). The larger the value, the more significant enrichment.
increased in neutrophils from type 2 diabetes patients compared to controls. (a) Cytokine-cytokine receptor interaction; (b) cell adhesion molecules; (c) leukocyte transendothelial migration. Red represents upregulated genes, and green represents downregulated genes.

Figure 5: Pathway map enriched by differentially expressed genes (DEGs) of neutrophils from type 2 diabetes patients compared with healthy controls. (a) Cytokine-cytokine receptor interaction; (b) cell adhesion molecules; (c) leukocyte transendothelial migration. Red represents upregulated genes, and green represents downregulated genes.

Figure 6: RT-qPCR validation of the RNA-seq results for a subset of 12 genes. In order to validate the RNA-seq transcriptome, 12 genes were selected from diverse biological functional categories, and RT-qPCR was performed on these genes. External samples including type 2 diabetes patients (n = 8) and healthy controls (n = 8) were selected for RNA-qPCR validation. Blue bars denote the RNA-seq fold induction values, while colored bars represent RT-qPCR fold induction values calculated using the 2^{-ΔΔCt} method, and data are presented as the mean ± SD. For RNA-qPCR validation, CXCR1, SELP, SELT, S100A8, S100A9, HSP1A1, BST2, PECAM1, S100A11, and S100A12 were significantly increased in neutrophils from type 2 diabetes patients compared with healthy controls (P < 0.05).

CXCL8, was downregulated. These data demonstrate that neutrophils from patients with diabetes may also exhibit impaired migration because of the downregulated chemotaxis, which may explain why some patients with diabetes have increased infection rates. Kuwabara et al. showed that neutrophils had an impaired response to LPS in a type 2 diabetes and obesity animal model, and neutrophils from the GK rats were not capable of migrating to the site of inflammation due to the impaired expression of adhesion proteins after LPS stimulation [32]. In addition, a defect in the chemotaxis of leukocytes in patients with diabetes has been identified, which could contribute to increased infections in these patients [33, 34], and high blood glucose induces a defective leukocyte-endothelial interaction in rats [35]. Our research is based on a small sample size, so the results may have limited generalizability. In addition the males are the majority among the participants, so our opinions may be generalizable mainly to males. Depending on the existing essay, the future research needs more representative sample and focus on making more depth analyses.

5. Conclusion

Our study investigated the presence of DEGs and the biological functions associated with these genes in neutrophils from type 2 diabetes patients and healthy controls. The study has shown that patients with type 2 diabetes have increased neutrophil activation, increased responses to chemokines, and increased neutrophil transendothelial cell migration at the transcriptome level. On the other hand, in patients with type 2 diabetes, neutrophil responses to molecules of bacterial origin, such as LPS, the cellular response to bacteria, and inflammatory reactions are reduced. These findings support the role of neutrophils in the pathogenesis of type 2 diabetes and
provide insight into potential therapeutic targets for type 2 diabetes.

Data Availability
The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest
The data are provided in the Supplementary Information files.

Acknowledgments
This work was supported by the National Natural Science Foundation of China (NSFC) grant (81870577 and 81670772 to Y.X.), National Key Research and Development Project (2016YFC1305000 and 2016YFC1305001 to Z.Z.), and Science and Technology Major Project of Hunan Province (2017SK1020 to Z.Z.).

Supplementary Materials

Supplementary 1. Supplementary Data 1: information of DEGs including gene ID, expression, and log2 ratio (T2D/HC).

Supplementary 2. Supplementary Data 2: information of main pathways identified in KEGG.

Supplementary 3. Supplementary Data 3: upregulated and downregulated gene annotations and identified biological processes in GO analysis.

Supplementary 4. Supplementary Table 1: RNA-seq sequence reads mapping to reference transcripts using Bowtie2. Supplementary Table 2: RNA-seq sequence reads mapping to reference genome using HISAT.

References

[1] M. Y. Donath and S. E. Shoelson, “Type 2 diabetes as an inflammatory disease,” Nature Reviews. Immunology, vol. 11, no. 2, pp. 98–107, 2011.

[2] S. Talukdar, D. Y. Oh, G. Bandyopadhyay et al., “Neutrophils mediate insulin resistance in mice fed a high-fat diet through secreted elastase,” Nature Medicine, vol. 18, no. 9, pp. 1407–1412, 2012.

[3] J. Huang, Y. Xiao, P. Zheng et al., “Distinct neutrophil counts and functions in newly diagnosed type 1 diabetes, latent autoimmune diabetes in adults, and type 2 diabetes,” Diabetes/Metabolism Research and Reviews, vol. 35, no. 1, article e3064, 2019.

[4] S. M. Cabrera, A. M. Henschel, and M. J. Hessner, “Innate inflammation in type 1 diabetes,” Translational Research, vol. 167, no. 1, pp. 214–227, 2016.

[5] E. Palella, R. Cimino, S. A. Pullano et al., “Laboratory Parameters of Hemostasis, Adhesion Molecules, and Inflammation in Type 2 Diabetes Mellitus: Correlation with Glycemic Control,” International Journal of Environmental Research and Public Health, vol. 17, no. 1, p. 300, 2020.

[6] C. Mertoglu and M. Gunay, “Neutrophil-lymphocyte ratio and platelet-lymphocyte ratio as useful predictive markers of prediabetes and diabetes mellitus,” Diabetes & Metabolic Syndrome: Clinical Research & Reviews, vol. 11, Supplement 1, pp. S127–S131, 2017.

[7] C.-T. C. Lee, S. B. Harris, R. Renfakaran et al., “White blood cell subtypes, insulin resistance and β-cell dysfunction in high-risk individuals - the PROMISE cohort,” Clinical Endocrinology, vol. 81, no. 4, pp. 536–541, 2014.

[8] J. Mestas and C. W. W. Hughes, “Of mice and not men: differences between mouse and human immunology,” Journal of Immunology, vol. 172, no. 5, pp. 2731–2738, 2004.

[9] E. Kolaczkowska and P. Kubies, “Neutrophil recruitment and function in health and inflammation,” Nature Reviews. Immunology, vol. 13, no. 3, pp. 159–175, 2013.

[10] V. Brinkmann, U. Reichard, C. Goosmann et al., “Neutrophil extracellular traps kill bacteria,” Science, vol. 303, no. 5663, pp. 1532–1535, 2004.

[11] A. Mocsai, “Diverse novel functions of neutrophils in immunity, inflammation, and beyond,” The Journal of Experimental Medicine, vol. 210, no. 7, pp. 1283–1299, 2013.

[12] B. Wierusz-Wysocka, A. Wykretowicz, H. Byks, K. Sadurska, and H. Wysocki, “Polymorphonuclear neutrophils adherence, superoxide anion (O2-) production and HBAl level in diabetic patients,” Diabetes Research and Clinical Practice, vol. 21, no. 2-3, pp. 109–114, 1993.

[13] F. Hanses, S. Park, J. Rich, and J. C. Lee, “Reduced neutrophil apoptosis in diabetic mice during staphylococcal infection leads to prolonged Tnfα production and reduced neutrophil clearance,” PLoS One, vol. 6, no. 8, article e23633, 2011.

[14] S. L. Wong, M. Demers, K. Martinod et al., “Diabetes primes neutrophils to undergo NETosis, which impairs wound healing,” Nature Medicine, vol. 21, no. 7, pp. 815–819, 2015.

[15] S. Bae, J. Choi, J. Hong et al., “Neutrophil proteinase 3 induces diabetes in a mouse model of glucose tolerance,” Endocrine Research, vol. 37, no. 1, pp. 35–45, 2012.

[16] D. Kim, B. Langmead, and S. L. Salzberg, “HISAT: a fast spliced aligner with low memory requirements,” Nature Methods, vol. 12, no. 4, pp. 357–360, 2015.

[17] B. Langmead and S. L. Salzberg, “Fast gapped-read alignment with Bowtie 2,” Nature Methods, vol. 9, no. 4, pp. 357–359, 2012.

[18] B. Li and C. N. Dewey, “RSEM: accurate transcript quantification from RNA-seq data with or without a reference genome,” BMC Bioinformatics, vol. 12, no. 1, p. 323, 2011.

[19] L. Wang, Z. Feng, X. Wang, X. Wang, and X. Zhang, “DEGseq: an R package for identifying differentially expressed genes from RNA-seq data,” Bioinformatics, vol. 26, no. 1, pp. 136–138, 2010.

[20] A. J. Van Oostrom, J. P. Van Wijk, T. P. Sijmonsma, T. J. Rabelink, and M. Castro Cabezas, “Increased expression of activation markers on monocytes and neutrophils in type 2 diabetes,” The Netherlands Journal of Medicine, vol. 62, no. 9, pp. 320–325, 2004.

[21] M. J. Sampson, I. R. Davies, J. C. Brown, K. Ivory, and D. A. Hughes, “Monocyte and neutrophil adhesion molecule expression during acute hyperglycaemia and after antioxidant treatment in type 2 diabetes and control patients,” Arteriosclerosis, Thrombosis, and Vascular Biology, vol. 22, no. 7, pp. 1187–1193, 2002.
[22] A. Shigeta, M. Matsumoto, T. F. Tedder, J. B. Lowe, M. Miyasaka, and T. Hirata, "An L-selectin ligand distinct from P-selectin glycoprotein ligand-1 is expressed on endothelial cells and promotes neutrophil rolling in inflammation," *Blood*, vol. 112, no. 13, pp. 4915–4923, 2008.

[23] G. R. Larsen, D. Sako, T. J. Ahern et al., "P-selectin and E-selectin. Distinct but overlapping leukocyte ligand specificities," *The Journal of Biological Chemistry*, vol. 267, no. 16, pp. 11104–11110, 1992.

[24] R. Kaur, J. Singh, R. Kapoor, and M. Kaur, "Association of SELP polymorphisms with soluble P-selectin levels and vascular risk in patients with type 2 diabetes mellitus: a case-control study," *Biochemical Genetics*, vol. 57, no. 1, pp. 73–97, 2019.

[25] K. Mastej and R. Adamiec, "Neutrophil surface expression of CD11b and CD62L in diabetic microangiopathy," *Acta Diabetologica*, vol. 45, no. 3, pp. 183–190, 2008.

[26] O. E. Sorensen and N. Borregaard, "Neutrophil extracellular traps - the dark side of neutrophils," *The Journal of Clinical Investigation*, vol. 126, no. 5, pp. 1612–1620, 2016.

[27] A. Citro, E. Cantarelli, and L. Piemonti, "The CXCR1/2 pathway: involvement in diabetes pathophysiology and potential target for T1D interventions," *Current Diabetes Reports*, vol. 15, no. 10, p. 68, 2015.

[28] A. Shiny, Y. S. Bibin, C. S. Shanthirani et al., "Association of neutrophil-lymphocyte ratio with glucose intolerance: an indicator of systemic inflammation in patients with type 2 diabetes," *Diabetes Technology & Therapeutics*, vol. 16, no. 8, pp. 524–530, 2014.

[29] C. Poitou, C. Perret, F. Mathieu et al., "Bariatric surgery induces disruption in inflammatory signaling pathways mediated by immune cells in adipose tissue: a RNA-seq study," *PLoS One*, vol. 10, no. 5, article e0125718, 2015.

[30] T. Fainsod-Levi, M. Gershkovitz, S. Völs et al., "Hyperglycemia impairs neutrophil mobilization leading to enhanced metastatic seeding," *Cell Reports*, vol. 21, no. 9, pp. 2384–2392, 2017.

[31] M. E. Falagas and M. Kompoti, "Obesity and infection," *The Lancet Infectious Diseases*, vol. 6, no. 7, pp. 438–446, 2006.

[32] W. M. T. Kuwabara, C. N. F. Yokota, R. Curi, and T. C. Alba-Loureiro, "Obesity and type 2 diabetes mellitus induce lipopolysaccharide tolerance in rat neutrophils," *Scientific Reports*, vol. 8, no. 1, article 17534, 2018.

[33] A. Mowat and J. Baum, "Chemotaxis of polymorphonuclear leukocytes from patients with diabetes mellitus," *The New England Journal of Medicine*, vol. 284, no. 12, pp. 621–627, 1971.

[34] M. A. Pereira, P. Sannomiya, and J. G. Leme, "Inhibition of leukocyte chemotaxis by factor in alloxan-induced diabetic rat plasma," *Diabetes*, vol. 36, no. 11, pp. 1307–1314, 1987.

[35] Z. B. Fortes, S. P. Farsky, M. A. Oliveira, and J. Garcia-Leme, "Direct vital microscopic study of defective leukocyte-endothelial interaction in diabetes mellitus," *Diabetes*, vol. 40, no. 10, pp. 1267–1273, 1991.