Identification of Biomarker $IL2R\beta$ in Ankylosing Spondylitis via Multi-Chip Integration Analysis of Gene Differential Expression

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Abstract

Background: Ankylosing spondylitis (AS) is a chronic inflammatory autoimmune disease that affects axial joints such as the spine. Early diagnosis is essential to improve treatment outcomes. The purpose of this study is to uncover underlying genetic diagnostic features of AS.

Methods: We downloaded gene expression data from the Gene Expression Omnibus (GEO) database for three studies of groups of healthy and AS samples. After preprocessing and normalizing the data, we employed linear models to identify significant differentially expressed genes (DEGs) and further integrated the differential genes to acquire reliable differential transcriptional markers. Functional enrichment analysis was conducted to obtain enriched pathways and regulatory gene interactions were extracted from pathways to further elucidate pathway networks. Seventy-three reliably differentially expressed genes (DEGs) were integrated by differential analysis. Utilizing the regulatory relationships of the 21 Kyoto Encyclopedia of Genes and Genomes (KEGG) Pathway genes that were enriched in the analysis, a regulatory network of 622 genes was constructed and its topological properties were further analyzed.

Results: Functional enrichment analysis found 73 DEGs that were strongly associated with immune pathways like Th17, Th1 and Th2 cell differentiation. Using KEGG combined with DEGs, six hub genes ($KLRD1$, $HLA-DRB3$, $HLA-DRB5$, $IL2R\beta$, $CD247$, and $CXCL10$) were suggested from the network. Of these, the $IL2R\beta$ gene was significantly differentially expressed compared with the normal control.

Conclusion: $IL2R\beta$ (Interleukin-2 receptor beta) is strongly associated with the onset and progression of autoimmune joint diseases, and may be used as a potential biomarker of AS. This study offers new characteristics that can help in the diagnosis and individualized therapy of AS.

Keywords: ankylosing spondylitis; $IL2R\beta$; crosstalk; biomarkers; support vector machines; pathway network

1. Introduction

Ankylosing spondylitis (AS) is a chronic inflammatory disease mediated by immunity that primarily impacts the axial joints like the spine [1]. AS can lead to severe stiffness, back pain and the formation of new bone, often resulting in progressive joint stiffness that decreases the quality of life [2]. Although the current evaluation of the disease state of AS measures disease activity, its progression and prognosis remain difficult to predict [3]. Furthermore, the potential molecular processes that promote AS progression remain largely unclear. Consequently, it is urgent to investigate the AS pathogenesis.

The rapid development of high-throughput transcriptomic profiling has made available massive repositories of data, enhancing efforts to improve the diagnosis of target diseases. In the past decade, several studies have used gene chips to determine candidate genes associated with AS [4–7] focusing on a single data set of samples. Specifically, Pimentel-Santo et al. [4] reported a group of differentially expressed genes (DEGs) highly correlated with AS that also modulate the joint destruction and inflammatory process. Similarly, Zhao et al. [7] utilized a bioinformatics approach to predict genes related to AS, including $MRPL22$, $RPL17$, $PSMA4$ and $PSMA6$. However, to date, most studies evaluating AS genetics have been concerned with individual genes or individual pathways, and multi-dataset integration analysis and multi-gene joint diagnosis are often overlooked.

A pathway contains a large amount of information about the interaction between genes and the synergy among them. Abnormality in some genes may lead to changes in other genes in the same pathway. As a result, establishing a pathway network and mining for disease markers has been an effective method in disease research.
This study integrates multi-platform high-throughput expression profiling data to (1) study altered gene expression patterns between AS and healthy women, (2) pre-screen diagnostic markers for AS, and (3) analyze these genes for their functions. These AS candidate biomarkers were further investigated by building Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway networks and an AS diagnostic classifier was established to assist clinical diagnosis.

2. Materials and Methods

Our method consisted of the following steps: collection of data, standardization, differential gene integration analysis, enrichment analysis, pathway integration analysis, pathway network construction, feature selection, and the construction and verification of the classifier. Fig. 1 summarizes the workflow.

2.1 Data Collection

We searched the Gene Expression Omnibus (GEO) database for studies on patients with AS and expression profiles from human blood samples were obtained. Three studies were identified and from them a dataset created of 76 cases and 45 healthy controls. Details are in Table 1.

2.2 Data Processing

For the two sets of gene expression data from the Illumina platform (http://www.ncbi.nlm.nih.gov/geo/), we downloaded the expression data after quantile normalization. Since there is no quantile normalization available for the GSE11886 dataset (http://www.ncbi.nlm.nih.gov/geo/), we downloaded the raw data and processed the data using the R package affy (version 4.2.0) [8] to acquire the probe expression profiles, which were further normalized using the RMA approach [9]. The result was an expression matrix of 54,675 gene probes for the 17 samples. The probes were annotated with R package hgu133plus2.db; probes matching multiple genes were deleted, and where multiple probes were matched to one gene, the median of the probes was used as the gene expression value. Subsequently, 23,313 gene expression profiles were acquired.

2.3 Integrated Analysis of Microarray Datasets

The R software package limma [10] was used to log₂-transform and normalize the matrix data in each GEO dataset. DEGs between normal and AS samples were then detected using the limma package. The DEGs recognized from the three datasets were genetically integrated with the R package “RobustRankAggreg” [11], which is
Table 1. Studies and data included in this analysis.

<table>
<thead>
<tr>
<th>Study GEO accession</th>
<th>Sample size</th>
<th>Sample source</th>
<th>Platform</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 GSE2510</td>
<td>16</td>
<td>Blood</td>
<td>GPL6947 Illumina HumanHT-12 V3.0 expression BeadChip</td>
</tr>
<tr>
<td>2 GSE73754</td>
<td>52</td>
<td>Blood</td>
<td>GPL10558 Illumina HumanHT-12 V4.0 expression BeadChip</td>
</tr>
<tr>
<td>3 GSE11886</td>
<td>8</td>
<td>Blood</td>
<td>GPL570 Affymetrix Human Genome U133 Plus 2.0 Array</td>
</tr>
</tbody>
</table>

Table 2. Characteristics of patients with ankylosing spondylitis and healthy controls.

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>AS patients</th>
<th>Healthy controls</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>35.73 ± 14.29</td>
<td>33.64 ± 12.23</td>
<td>0.6687</td>
</tr>
<tr>
<td>Male (%)</td>
<td>73%</td>
<td>64%</td>
<td>1.000</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>72.82 ± 19.06</td>
<td>69.82 ± 15.73</td>
<td>0.7053</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>167 ± 11.01</td>
<td>168.7 ± 7.254</td>
<td>0.5568</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>28.09 ± 7.354</td>
<td>24.45 ± 5.507</td>
<td>0.2315</td>
</tr>
<tr>
<td>Disease duration (years)</td>
<td>11.7 (1, 20)</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>HLA-B27 (+)</td>
<td>55%</td>
<td>—</td>
<td>—</td>
</tr>
</tbody>
</table>

Table 3. Primer sequences utilized in quantitative Real-Time PCR (qRT-PCR).

<table>
<thead>
<tr>
<th>Genes</th>
<th>F/R</th>
<th>Sequences (5'-3')</th>
</tr>
</thead>
<tbody>
<tr>
<td>KLRD1</td>
<td>Forward</td>
<td>CAGGACCCAACATAGAACTCCA</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>GAAAGAAATGAACTAAGTGCACC</td>
</tr>
<tr>
<td>HLA-DRB3</td>
<td>Forward</td>
<td>CGGGGCTGGTGAGAGCTTC</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>AACCACCTGACTTCAATGCTG</td>
</tr>
<tr>
<td>HLA-DRB5</td>
<td>Forward</td>
<td>CGGGGCTGGTGAGAGCTTC</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>AACCACCTGACTTCAATGCTG</td>
</tr>
<tr>
<td>IL2RB</td>
<td>Forward</td>
<td>CTTGAGGAGGAGGGTTA</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>GGTTGAGTAAAGTACACTCT</td>
</tr>
<tr>
<td>CD247</td>
<td>Forward</td>
<td>GCCAGAACAGCTCTTATAAGG</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>GCCAGACGTCTTGTCCAA</td>
</tr>
<tr>
<td>CXCL10</td>
<td>Forward</td>
<td>GTGGACATTCAAGGGAGTACCCT</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>TGATGCCCTCGTCTTGAGTT</td>
</tr>
</tbody>
</table>

2.4 Functional Enrichment Analyses

R package clusterProfiler [14] was exploited for analyzing KEGG and Gene Ontology (GO) pathway enrichment to determine GO terms that appeared disproportionately in three categories (cellular component, molecular function, and biological processes). KEGG pathways were also analyzed by visualization with R package DOSE [15]. In both analyses, q-values (adjusted p-values for multiple tests) less than 0.05 indicate statistical significance.

2.5 Development of KEGG Pathway Gene Interaction Network

We downloaded the XML file of each KEGG pathway that was enriched in the DEGs from the KEGG website [16]. The R package XML was employed to extract entries, relationships, and group relationships from these XML files. The information on gene interactions was subsequently extracted with a combination of scripts to build the KEGG gene interaction network to visualize the network on Cytoscape [17], and examine the topological characteristics of the cytohubba network [18].

2.6 Network Topological Property Analysis to Screen AS Key Diagnostic Genes

Visualization was performed using Cytoscape [17], with the degree and closeness of the network computed through the use of the Cytoscape [17] plug-in cytohubba [18]. Genes of degree >20 and closeness >5 were selected and intersected with the differential genes to identify the final characteristic genes.

2.7 Development of Predictive Models for AS Diagnosis and Assessment of Model Predictive Capability

A support vector machine (SVM) [19] was used to build a diagnostic predictive model using genes to predict healthy versus AS individuals. As a supervised learning model, SVM uses machine learning algorithms for data analysis and pattern recognition. SVM builds a hyperplane that can be applied to categorization and regression in infinite-dimensional or high-dimensional spaces. A group of training samples, each labeled as belonging to one of two classes is used by the SVM training algorithm to create a model, and new instances are then assigned to a category according to the model, thereby making it a non-probabilistic binary linear categorization. The three datasets were categorized into two validation datasets and one training dataset. The model was built with the training dataset, and its classification capability was confirmed using the 10-fold cross-validation approach. The constructed model was, in turn, used to predict the category of samples based on the Robust Rank Aggregation (RRA) approach. Those genes that consistently ranked superior to expectations based on the null hypothesis of input irrelevance could be screened with the RRA approach. Therefore, an integration of gene expression values for samples from various datasets was not performed. In common with many who have utilized the RobustRankAggreg package [12,13], we did not correct for batch effects.

p < 0.05 together with 1.2-fold change in DEGs were deemed to be statistically significant.
Fig. 2. Determination of differentially expressed genes (DEGs) between Healthy Control and AS Samples. (A) Box plot of the overall expression level of the normalized samples (green bars: normal samples, red bars: AS samples). (B) Heatmap of the down- and up-regulated DEGs from the integrated microarray analysis. Each column and row denote a dataset and a gene, separately. In each rectangle, the color denotes the log$_2$-FC value. The gradient from green to red goes from descending to ascending expression. (C) The enrichment results of log$_2$-FC in each dataset of differentially expressed genes. FC, fold change.

in the validation dataset. The model’s predictive power was evaluated using the area under the receiver operating characteristic curve (ROC) curve (AUC), and the model was analyzed for its specificity and sensitivity in predicting AS.

2.8 Sample Preparation

Subjects were AS patients aged between 16 and 61 who were hospitalized in Jishuitan Hospital for the first time. The inclusion criteria were: (1) the diagnosis was in accordance with the New York classification revised in 1984; and (2) patients had not yet received medical treatment, including non-steroidal anti-inflammatory drugs or disease-modifying anti-rheumatic drugs. The exclusion criteria were: (1) individuals with a history of severe infection, surgery, or tumor; (2) Individuals with severe heart, lung, liver, and kidney dysfunction; and (3) individuals with a history of autoimmune diseases such as rheumatoid arthritis. Included in the study was a control group consisting of healthy volunteers between 24 and 60 without acute or chronic diseases, such as rheumatoid arthritis, and no long-term use of any drugs. Table 2 shows the characteristics of the AS patients and healthy control subjects.

Our experiments were in line with the principles of the Declaration of Helsinki and were approved by the ethics committee of Beijing Jishuitan Hospital (ethics code: 201611-03). From all participants, fasting whole blood was
collected in sodium heparin-coated tubes (Greiner Bio-one, Chonburi, Thailand) and stored at $-80^\circ$C until further analysis. All RNA samples had an A260/A280 ratio between 1.66 and 2.58 and concentrations ranging from 10.9–143 ng/mL and thus met the criteria for gene expression analysis.

2.9 Quantitative Real-Time PCR (qRT-PCR)

Extraction of total RNA from blood samples was implemented with Trizol reagent (Tiangen Biochemical, Beijing, China) and later extracted RNA was reverse transcribed to cDNA using FastQuant RT Kit containing gDNase (Tiangen Biochemical). Amplification was performed on an ABI7500 real-time fluorescence quantitative PCR instrument utilizing the Power SYBR® Green PCR Kit (Applied Biosystems, Foster City, CA, USA). Table 3 gives the primer sequences. The procedure followed was: pre-denaturation 95 $^\circ$C for 10 min; 95 $^\circ$C for 15 s; and 60 $^\circ$C for 60 s for 40 cycles. Three replicates were employed. Relative expression levels were determined via the $2^{-\Delta\Delta CT}$ method.

2.10 Statistical Analysis

All data are expressed as mean ± standard deviation (SD) calculated using the GraphPad Prism 9.0 software (GraphPad Software Inc., San Diego, CA, USA). An unpaired two-tailed Student’s t-test was performed to compare the differences between healthy and AS groups. $p < 0.05$ was considered statistically significant.

3. Results

3.1 Determination of DEGs between Healthy Control and AS Samples

Gene chip data from three datasets, GSE2510, GSE73754, and GSE1188655, were acquired from the GEO database. Following standardization, the gene expression distribution of each sample appeared similar (Fig. 2A). Genes with differential expression between AS and healthy samples from all three data sets were then further integrated and analyzed using limma. Altogether, 73 DEGs were identified (Supplementary Table 1), with 31 and 42 genes increased in the healthy and AS group, respectively, as indi-
Fig. 4. KEGG pathway gene interaction network analysis. (A) Crosstalk among genes; color, and size between KEGG pathways indicate the number of pathways in which a particular gene is involved. (B) The interaction network; color and size of KEGG pathway genes indicate the number of pathways in which genes participate.

Fig. 5. Network topological property analysis to screen AS key genes. (A) Distribution of the number of pathways involved in the pathway gene network. (B) Degree distribution of the network. (C) Closeness distribution of the network. (D) Betweenness distribution of the network.

cated by the differential multiple heatmaps in each dataset (Fig. 2B). The enrichment results of the differential multiple GSEA [20] in all data sets were analyzed (Fig. 2C). In each data set, up- and downregulated genes were predominantly enriched in groups with high and low differential multiples, respectively. The only exception was observed for the upregulated genes in data set GSE2510 where they were enriched in the group with lower differential multiplicity, which may be caused by sampling error.
3.2 Functional Enrichment Analysis of DEGs

To gain insight into the functional implications of these 73 DEGs, GO and KEGG functional enrichment analyses were implemented on the 31 upregulated and 42 downregulated DEGs. While 42 increased genes were not enriched in any KEGG pathways and GO terms, 31 decreased genes were identified as enriched in multiple KEGG pathways and GO terms. In the bioprocess group, five enriched GO terms were noted, including “cellular defense response”, “cytolysis”, and “T cell costimulation” (Fig. 3A), suggesting that these genes are associated with the cellular defense response. Classification by cellular component displayed seven enriched GO terms, for example, “lumen side of endoplasmic reticulum membrane” and “MHC protein complex” (Fig. 3B). Seven molecular functions were found to be mediated by these genes and were primarily related to “heparin-binding” and “antigen-binding” (Fig. 3C). Furthermore, 21 KEGG pathways were identified including “Type I diabetes mellitus”, “Th17 cell differentiation”, and “Th1 and Th2 cell differentiation” (Fig. 3D). Previous studies have also found a close relationship between “Th17 cell differentiation” and AS [21–23]. Together these data suggest that the downregulated DEGs are closely related to immunity.

3.3 KEGG Pathway Gene Interaction Network

For the 21 KEGG pathways identified, the XML files were downloaded from the KEGG website, and genetic interaction information was extracted using the R package XML. Specifically, gene ID was transformed into a gene symbol, a KEGG pathway gene interaction network was built and 622 genes were derived. Genetic analysis of the crosstalk between these pathways indicates that a close relationship exists among most pathways, and some genes are involved in multiple pathways (Fig. 4A). A total of 1663 genetic interactions were obtained based on the regulatory information of genes in the KEGG pathway (Fig. 4B). In addition, most of the genes involved in multiple pathways are closely related to other genes in the network.
3.4 Network Topological Property Analysis to Screen AS Key Genes

We analyzed the pathways involved in the KEGG gene interaction network and found that in total, 248 (39.8%) genes were involved in only one pathway (Fig. 5A). A small number of genes are involved in multiple pathways. We also observed a power-law degree distribution in our network (Fig. 5B), where the degree of most genes is less than 20. Additionally, closure of the network revealed that the majority of the nodes have lower closeness and only a few have high closeness (Fig. 5C). The network distribution of betweenness was also computed and the majority of the nodes exhibit a lower betweenness (Fig. 5D). Nodes with high degree, high betweenness or high closeness are viewed as significant in a network.

We considered a node to be the hub gene of the access network when it met the criteria of degree $>20$, closeness $>5$, and betweenness $>0$. Six genes met this criteria (KLRD1, HLA-DRB3, HLA-DRB5, IL2Rβ, CD247, CXCL10). Among these genes, HLA-DRB3 and HLA-DRB5 are HLA class II histocompatibility antigens. Major histocompatibility complex (MHC) is an important risk factor for AS; IL2RB is an important regulator of cytokine signaling and RET signaling in the immune system and has been reported as a potential transcriptional marker for AS signaling and RET signaling in the immune system and has been reported as a potential transcriptional marker for AS and IL2RB is an important regulator of cytokine signaling and RET signaling in the immune system and has been reported as a potential transcriptional marker for AS.

3.5 Establishment and Testing of the Diagnostic Model

To construct a diagnostic model, GSE2510 was used as the training dataset ($n=32$, AS=16) while GSE73754 ($n=72$, AS=52) and GSE11886 ($n=17$, AS=8) were used as validation datasets. Six previously chosen hub genes were utilized as features in the training dataset to acquire their corresponding expression profiles. After the z-score, the support vector machine classification model was established. A 10-fold cross-validation indicated the classification accuracy was 93.8%, with 30 of the 32 samples categorized correctly (Fig. 6A). The model’s specificity and sensitivity for AS were both 93.8%. The AUC was 0.94 (Fig. 6D). Validation of the model was then conducted with the GSE73754 dataset; 62 out of 72 samples were categorized correctly for an 86.1% classification accuracy. The model’s specificity and sensitivity for AS in this dataset were 60% and 96.2%, respectively (Fig. 6B) and the AUC was 0.78 (Fig. 6E). Finally, for the GSE11886 dataset, 15 of 17 samples were correctly categorized with 88.2% classification accuracy. The model’s specificity and sensitivity for AS were 88.9% and 87.5%, separately (Fig. 6C) and the AUC was 0.88 (Fig. 6F).

We validated these gene expressions in clinical samples. The findings demonstrated a significant difference in $IL2Rβ$ expression versus normal controls, which agrees with the findings of our bioinformatics analysis (Fig. 7).

![Fig. 7. Expressions of $IL2Rβ$ in AS samples.](image)

These findings suggest that the diagnostic prediction model established in this research can differentiate effectively between healthy individuals and those with AS and that $IL2Rβ$ can be used as a reliable biomarker for AS diagnosis.

4. Discussion

The comprehensive bioinformatics analysis focuses on the molecular screening of differential gene expression, and the network-based hub gene discovery has been extensively applied to determine underlying biomarkers relevant to the AS diagnosis and therapy. For example, Xu et al. [27] screened for hub genes by combining DEG screening, functional enrichment analysis, and PPI network construction, and further verified their outcomes by qRT-PCR. Ma et al. [28] used a set of GEO datasets to identify AS-related genes. Kanwal et al. [29] used protein interaction network analysis to find a CD4-centered AS anomaly-related regulatory network. Zhao et al. [7] used a set of GEO datasets for DEG screening, miRNA target gene analysis, and PPI network construction in efforts to screen for AS-related genes. Compared with previous studies, the current research not only integrates a relatively large sample size of microarray data from multiple GEO datasets but also constructs a pathway-based gene regulatory network in addition to an AS diagnostic model to identify potential diagnostic biomarkers for AS.

In our study, 73 consistent DEGs were identified, of which 42 were increased and 31 were decreased. The
genes downregulated were enriched in multiple GO terms and KEGG pathways. These pathways and GO terms were primarily correlated with immunological functions and the major histocompatibility complex (MHC), which is in keeping with prior findings that the immune pathway is affected in AS patients [6,30]. CD2 is the most upregulated gene, a T-cell surface antigen that optimizes immune recognition by interacting with LAE3 (CD58) on antigen-presenting cells and is linked to the onset and progress of various diseases [31–33]. The upregulated genes are not enriched in any of the KEGG pathways and GO terms, and thus their exact contribution to AS development is unknown. Further investigation is essential to clarify the roles of these genes as potential transcriptomic markers for AS. We identified six key genes, KLRE1, HLA-DRB3, HLA-DRB5, IL2RB, CD247, and CXCL10 in the downregulated KEGG pathway gene regulatory network. Among these, IL2RB has been previously reported as a potential transcriptional marker for AS [24] and CXCL10 is a distinguishing marker of autoimmune joint diseases [25,26]. All six genes are strongly correlated with the onset of autoimmune joint diseases.

In addition, as a common rheumatic disease, AS can cause inflammatory back pain, which greatly reduces quality of life [34]. Therefore, early diagnosis together with personalized medical interventions for AS are critical. In the past several years, gene expression profiling has been broadly applied to recognize biomarkers correlated with the disease [35–37]. Some of these identified biomarkers are functionally similar but have poor reproducibility, indicating they may not have accurate classification capabilities. In the current study, SVMs were employed to build and verify classifiers based on the expression profiles of six genes related to AS. The AUC of the training set reached 0.94 with our model, showing that these genes classify AS well. Additionally, in the verification datasets, we achieved AUCs of 0.88 and 0.78, which suggests these six genes could be applied as diagnostic markers for AS. Our results in clinical samples show that IL2RB is significantly differentially expressed in individuals with AS compared to healthy, which is consistent with our bioinformatics analysis. There were no significant differences in other genes, which may reflect the small sample size of the study. These findings reveal that the diagnostic prediction model built in this research can differentiate patients with AS from healthy controls effectively and that IL2RB can be a reliable biomarker for the diagnosis of AS. Moreover, IL2RB may be a potential target for the treatment of AS and/or a marker for monitoring disease activity and progression in AS patients.

Our work employed bioinformatics tools to identify underlying candidate genes involved in AS pathogenesis with large samples. However, limitations remain. First, the sample lacked other information about the clinic. Consequently, we did not account for the factors like whether the patients had other diseases or health conditions. Second, the clinical sample size was small in validation experiments. Hence, further experimental and genetic studies with larger sample sizes and preclinical validation are required.

5. Conclusion

IL2RB is strongly associated with the onset and progression of autoimmune joint disease and may be a potential biomarker of AS. This study offers new genetic characteristics to help diagnosis and individualized therapy of AS. Our findings further provide potential targets and biomarkers for clinicians and researchers.

Abbreviations

AS, ankylosing spondylitis; GEO, Gene Expression Omnibus; DEGs, differentially expressed genes; GO, Gene Ontology; KEGG, Kyoto Encyclopedia of Genes and Genomes; SVM, support vector machine.

Availability of Data and Materials

The datasets generated or analyzed during this study are available from the corresponding author on reasonable request.

Author Contributions

CAW, BX and CW designed the study. BX, PLC, ZMW and YZZ carried out data acquisition and analysis. ZMW, YZZ and HCL carried out the verification experiment. BX and PLC wrote the manuscript. CW and CAW were revising the paper. HCL and CW contributed to preparing and making figures. All authors contributed to editorial changes in the manuscript. All authors read and approved the final manuscript. All authors have participated sufficiently in the work and agreed to be accountable for all aspects of the work.

Ethics Approval and Consent to Participate

All procedures for sample collection and use were approved by the Ethics Committee of Beijing Jishuitan Hospital, Beijing, China (ethics code: 201611-03). All authors approved to participate.

Acknowledgment

Not applicable.

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Conflict of Interest

The authors declare no conflict of interest.
Supplementary Material

Supplementary material associated with this article can be found, in the online version, at https://doi.org/10.31083/j.fbl2812343.

References

