

Identification of Key Candidate Genes Related to Inflammatory Osteolysis Associated with Vitamin E-Blended UHMWPE Debris of Orthopedic Implants by Integrated Bioinformatics Analysis and Experimental Confirmation

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Purpose: This study aims to identify differentially expressed genes (DEGs) in macrophages exposed to ultra-high-molecular-weight polyethylene (UHMWPE) or vitamin E-blended UHMWPE (VE-UHMWPE) particles, thereby providing potential targets for the treatment of inflammatory osteolysis.

Methods: The GSE104589 dataset of genome expression in macrophages exposed to UHMWPE and VE-UHMWPE was downloaded from the Gene Expression Omnibus database to identify DEGs. Functional enrichment analysis was performed using DAVID, and the corresponding protein-protein interaction (PPI) network was constructed from the STRING database. Important modules were selected using the molecular complex detection algorithm, and hub genes were identified in cytoHubba. MicroRNAs targeting these DEGs were obtained from the TarBase, miRTarBase, and miRecords databases, while transcription factors (TFs) targeting DEGs were predicted from the ENCODE database. Finally, the top five DEGs were validated by quantitative real-time polymerase chain reaction (qRT-PCR).

Results: A total of 112 DEGs (44 upregulated and 68 downregulated DEGs) were screened. Immune and inflammatory responses were significantly related in gene ontology analysis, and 18 signaling pathways were enriched according to Kyoto Encyclopedia of Genes and Genomes pathway analysis. The PPI network involving 85 nodes and 266 protein pairs indicated that IL1 β , CXCL1, ICAM1, CCL5 and CCL4 showed higher degrees. qRT-PCR analysis of the top five DEGs revealed a decreasing trend in the VE-UHMWPE group compared with the UHMWPE group. Key microRNAs (hsa-miR-144, hsa-miR-21, and hsa-miR-221) and TFs (RELA and NFkB1) were predicted to be correlated with the pathogenesis of inflammatory osteolysis through microRNA-TF regulatory network analysis.

Conclusion: The present study helps shed light on the molecular mechanisms underlying the changes in the wear-induced inflammatory process after blending vitamin E with UHMWPE. Hub genes including IL1 β , CXCL1, ICAM1, CCL5, and CCL4, key microRNAs (hsa-miR-144, hsa-miR-21, and hsa-miR-221) and TFs (RELA and NFkB1) may serve as prognostic and therapeutic targets of inflammatory osteolysis.

Keywords: inflammatory osteolysis, aseptic loosening, bioinformatics analysis, VE-UHMWPE, macrophages



Introduction

While total joint arthroplasty (TJA) is still the gold standard for the treatment of pain and dysfunction in patients with end-stage arthritis and femoral head necrosis,^{1,2} determining the long-term behavior of implants remains challenging in the scientific community and healthcare service systems due to polyethylene wear, with resultant osteolysis and/or aseptic loosening.³ Revision surgery is more difficult and costly than primary TJAs because of poorer prognosis and higher risk of failure and complications.^{2–5} Inflammatory osteolysis initiates when macrophages recognize wear debris derived from the polyethylene (PE) liners of TJAs and release a series of cytokines and chemokines that promote the recruitment of other inflammatory cells, leading to the formation of inflamed granulomas.⁴ Despite advances achieved in pharmacological therapies for inflammation in other skeletal diseases, there are no agents that specifically target periprosthetic inflammatory osteolysis.

A better understanding of the molecular and cellular mechanisms of inflammatory osteolysis in TJA and the development of specific therapies are urgently needed to prevent implant loosening. Attempts such as the introduction of ultra-high-molecular-weight polyethylene (UHMWPE) and cross-linking of UHMWPE using gamma irradiation (HXLPE) have been made to improve the wear resistance of implant bearing materials (acetabular components in total hip arthroplasties and tibial inserts in total knee arthroplasties) and have revealed superior durability in mid-term and long-term follow-up.⁴ However, the cross-linking process of polyethylene generates free radicals, reducing long-term oxidative stability and enhancing material embrittlement of HXLPE.^{1,6} To address this issue, the new generation of HXLPE is stabilized with vitamin E (alpha-tocopherol), a natural lipid-soluble antioxidative substance exerting potent anti-inflammatory properties on cells.⁷ The infusion of vitamin E into UHMWPE could donate hydrogen atoms to free radicals in polyethylene, decrease the production of reactive oxygen species (ROS) and tumor necrosis factor α (TNF- α), and interrupt the chemical reaction cascade between the macromolecules and oxygen.⁸ Currently, there are two methods of integrating vitamin E into UHMWPE.² The first is to blend vitamin E with UHMWPE resin powder: after consolidation, the blend can be irradiated for cross-linking.⁵ However, the presence of vitamin E could reduce the cross-linking efficiency of UHMWPE and a concentration of vitamin E above 0.3% wt (mass fraction) in the irradiated surface

would significantly impair wear characteristics.^{9,10} The second is doping the polyethylene polymers after radiation cross-linking in vitamin E, by which a maximum concentration of approximately 0.7 wt% can be reached on account of the saturation upper limit of the HXLPE.^{2,5} Bichara and colleagues revealed that fabricated wear particles derived from VE-UHMWPE triggered lesser cellular responses and osteolytic potentials than conventional UHMWPE in *in vivo* experiments,¹¹ and a series of randomized controlled trials demonstrated lower wear rates in HXLPE/VitE cups than in HXLPE or UHMWPE cups in mid- or long-term follow-ups.^{1,5,7,12} However, some investigators obtained negative results for these two materials.^{12–14} Until now, whether the addition of vitamin E exerts a protective effect on the bearing materials of arthroplasties remained to be confirmed.

Given that macrophages are the first immune cells encountering wear particles, we attempted to analyze the genomic mechanisms of human macrophages after exposure to VE-UHMWPE and conventional UHMWPE particles. Notably, Terkawi and his colleagues originally submitted the GSE104589 dataset to the Gene Expression Omnibus (GEO) database. They screened correlated gene expression in human macrophages exposed to VE-UHMWPE and UHMWPE particles, demonstrating the involvement of IL-27 in the macrophage response to VE-UHMWPE particles and its regulatory role in osteolysis.¹⁵ However, they only investigated the function of one transcriptome gene (IL-27),¹⁵ and there were still multiple genes that could play vital roles in these processes. Therefore, to identify the differentially expressed genes (DEGs) and correlated pathways in macrophages after exposure to VE-UHMWPE and UHMWPE, and provide a more comprehensive view of the genetic mechanisms in inflammatory osteolysis, in the present study these microarray data were retrieved from the GEO database and reanalyzed by using various bioinformatics technologies and *in vitro* experiments. Furthermore, the present study constructed a transcription factor (TF)-DEG network and a microRNA (miRNA)-DEG network. The associations between the TFs and their targets were predicted by using the Encyclopedia of DNA Elements (ENCODE), while the regulatory associations between miRNAs and their targets were extracted from the TarBase, miRTarBase and miRecords databases. The integration of these two networks was then performed to establish a TF-microRNA coregulatory network from which key

microRNAs and TFs can be identified. Our analysis provided insight into genomics at the cellular level and explored the potential mechanism of inflammatory osteolysis.

Materials and Methods

High-Throughput Gene Expression Data

The high-throughput gene expression profile was downloaded from the GEO database (<https://www.ncbi.nlm.nih.gov/geo/>), under the accession number GSE104589. The GSE104589 dataset, based on GPL16791 (a platform for Illumina HiSeq 2500, Homo sapiens) contained 9 samples, including 3 samples from macrophages exposed to VE-UHMWPE particles, and 3 samples from macrophages exposed to UHMWPE and 3 control samples from macrophages cultured alone, which were all selected for deep analysis in this study.

Data Preprocessing and Identification of DEGs

Based on the annotation information in the platform, the probe sets were transformed into the corresponding gene symbol. The mean value of multiple probe sets was calculated if they corresponded to the same gene symbol. The data were normalized using quantile normalization with Illumina¹⁶ in R software (Version 3.6.2) and DEGs were identified using the limma package (Version 3.42.2) in R software. Individual *p*-values were calculated and converted to adjusted *p*-values (adj. *p*. val) for comparisons by false discovery rate correction of the Benjamini and Hochberg test. The cutoff point of adj. *p*. val < 0.05 and |log₂fold change (FC)| > 2 were used to select the DEGs. Then, the heatmap and volcano plot of the DEGs were generated using the ggplot2 package in R software.

Functional and Pathway Enrichment Analyses of DEGs

The up- and downregulated DEGs were uploaded to an online biological information database, the Database for Annotation, Visualization, and Integrated Discovery (DAVID) version 6.8 Beta (<https://david-d.ncifcrf.gov/>),^{17,18} for further analysis, including gene ontology (GO) and Kyoto Encyclopedia of Gene and Genome (KEGG) pathways, which were visualized in the R ggplot2 package. The significance criterion was set at *p*-val < 0.05.

Protein-protein Interaction Network Construction and Module Analysis

The PPI network of DEGs was predicted using the multiple protein online tool in the STRING database,¹⁹ a search tool for the Retrieval of Interacting Genes database (version 11.0, <http://string-db.org>) and visualized in Cytoscape software (Version 3.6.2).^{20,21} The most significant modules in the PPI network were identified and visualized using Molecular Complex Detection (MCODE), which is a tool in Cytoscape to select densely connected areas in the acquired network. The selection criteria were as follows: MCODE scores > 5, node score cutoff = 0.2, degree cutoff = 2, max depth = 100 and k-score = 2. The biological process analysis of the top 20 genes was performed and visualized using the Biological Networks Gene Ontology tool (BiNGO) (version 3.0.3) plugin in Cytoscape.²²

Hub Gene Selection

The hub genes (top 10) in the PPI network were identified by Maximal Clique Centrality (MCC), Degree, and Maximum Neighborhood Component (MNC) from the cytoHubba plugin in Cytoscape.

TF-DEG Network and miRNA-DEG Network Construction

The targeted TFs of DEGs were predicted using the ENCODE database in NetworkAnalyst (<https://www.networkanalyst.ca/faces/home.xhtml>), a visual analytics platform for comprehensive gene expression profiling and meta-analysis. Meanwhile, the microRNA-DEG pairs were identified using the TarBase, miRTarBase and miRecords databases. The generated pairs of TF-DEG and miRNA-DEG were retained for further analysis.

TF-miRNA Integrated Network Construction

The integrated network of TF-miRNA was constructed according to the TF-DEG and miRNA-DEG networks. The coregulated DEGs targeted by miRNAs and TFs were first selected, and subsequent miRNAs and TFs were extracted. Eventually, the integrated regulatory network was constructed and visualized in Cytoscape software (Version 3.6.2).

Preparation of Polyethylene Particles and Macrophage Culture

The materials were prepared from a single batch of medical-grade GUR 1050 resin (Ticona, Bayport, TX, USA). The VE-UHMWPE and UHMWPE particles were first fabricated using the method presented by Oral.²³ The human monocyte cell line THP-1 (ATCC; American Type Culture Collection, USA) was seeded in a 6-well plate at a density of 5×10^5 cells/well and cultured with RPMI-1640 medium (BSigma Aldrich, St. Louis, MO, USA) supplemented with 10% inactivated fetal bovine serum, 100 U/mL penicillin, 100 µg/mL streptomycin, and 2 mM L-glutamine. Cells were incubated in a humidified atmosphere with 5% CO₂ environment at 37°C. The cells were differentiated into macrophages by using 125 ng/well phorbol 12-myristate 13-acetate (PMA) (Sigma Aldrich, St. Louis, MO, USA) for 48 h. Then the cells were cultured with free-endotoxin VE-UHMWPE or UHMWPE particles at a ratio of 1:4000 µm³ for 24 h using an inverted cell culture technique that allows particles to directly interact with macrophages. Control macrophages were cultured alone.

Quantitative Real-Time Reverse Transcription Polymerase Chain Reaction (qRT-PCR)

Purified 0.5 mg DNA-free RNA samples were reverse transcribed using a One Step PrimeScript[®] miRNA cDNA Synthesis Kit (Takara, Japan, D350A), and the amplification reaction was conducted using a SYBR[®] Premix Ex TaqTM II kit (Takara, Japan, DRR820A). Gene expression of each target was calculated using the $\Delta\Delta CT$ method after normalization to the expression of GAPDH housekeeping genes.

ELISA

Secreted IL-1 β , IL-1, IL6 and TNF- α levels in the supernatant of macrophages in different groups were detected using quantitative ELISA kits (R&D Systems Inc. MN, USA) according to each manufacturer's instruction.

Statistical Analysis

Statistical analyses were performed using GraphPad Prism Software, version 7.0 (GraphPad Software Inc., USA). One-way analysis of variance (ANOVA) followed by Tukey's multiple-comparison procedure was used to compare the gene expression and levels of inflammatory

factors. The results are presented as the means \pm standard errors (SEM), and a p -value < 0.05 was considered statistically significant.

Results

Data Normalization

Data normalization and cross-comparability were evaluated via the box plot visualization of the gene expression of all samples and principal component analysis (PCA) for confirming biological variability between different samples. Figure 1 demonstrates that the black lines were almost in the same position after data normalization, indicating an excellent degree of standardization, which can ensure the accuracy of subsequent data processing. Figure 2A shows that all samples involving UHMWPE, VE-UHMWPE, and no particles were grouped separately, indicating globally distinct expression profiles.

Identification of DEGs

The high-throughput sequencing data from the GSE104589 dataset were analyzed to select DEGs according to the predefined criteria. In total, 112 DEGs were detected between the VE-UHMWPE group and the UHMWPE group, including 44 upregulated and 68 downregulated DEGs. In addition, a volcano plot and a heatmap of all DEGs were generated using the R ggplot2 package (Figure 2B and C). Meanwhile, 376 DEGs were identified between the UHMWPE group and the control group, including 182 upregulated and 194 downregulated DEGs, and the top 50 genes are listed in [Supplementary Table 1](#).

GO Function and KEGG Enrichment Analysis of DEGs

GO enrichment analysis of 112 DEGs between the VE-UHMWPE group and the UHMWPE group was performed to identify the most relevant biological processes (BPs), molecular functions (MFs), and cellular components (CCs). The top ten enriched terms in BP, CC, and MF are presented in Figure 3A–C. Additionally, based on KEGG pathway analysis, the DEGs were significantly enriched in 18 signaling pathways, such as the chemokine signaling pathway, cytokine–cytokine receptor interaction, rheumatoid arthritis, herpes simplex infection and influenza A, measles, and legionellosis (Figure 3D). The top 20 terms associated with 112 DEGs, enriched in BP, CC and MF are presented in Table 1. GO enrichment analysis and related pathways of 376 DEGs between the UHMWPE group and

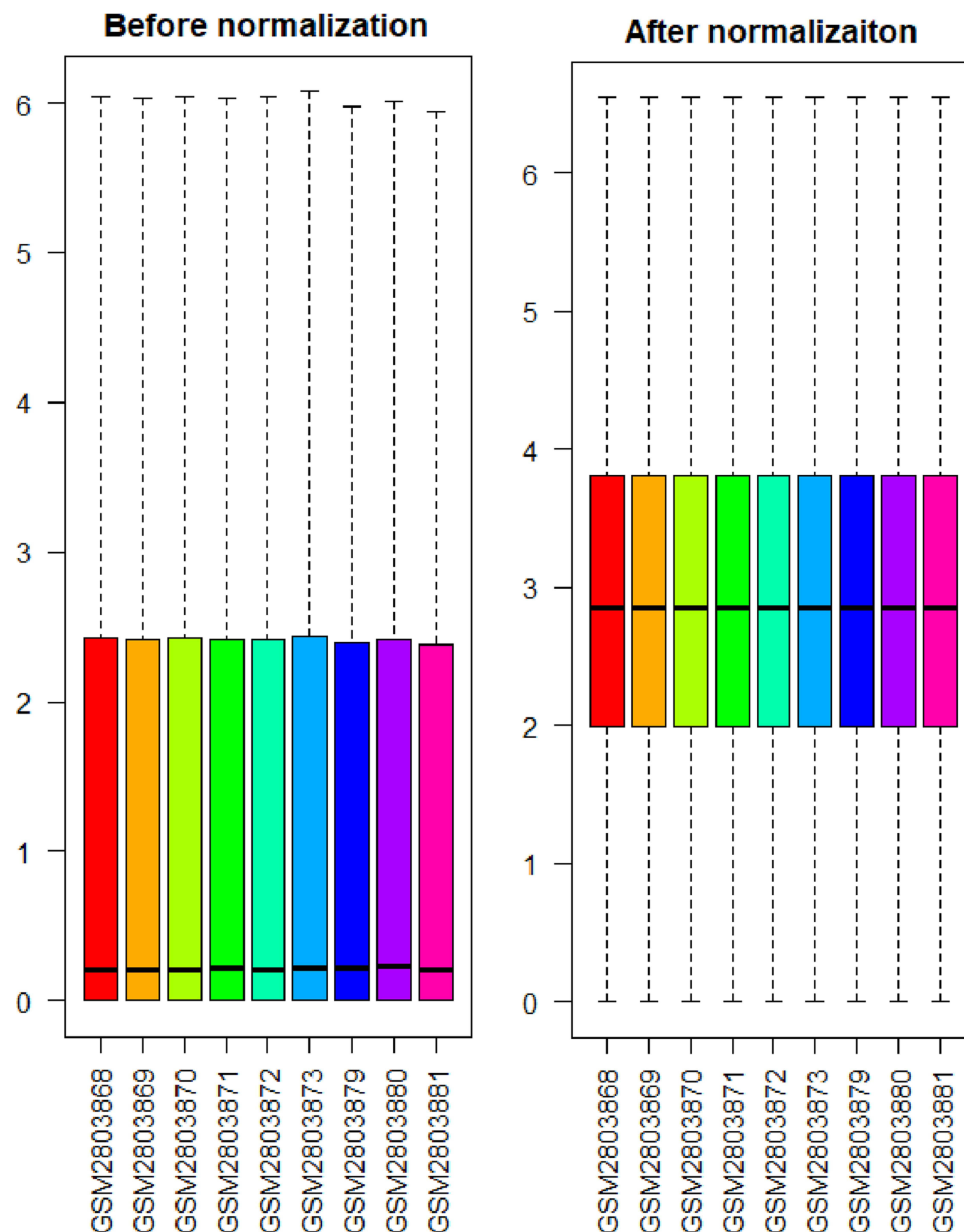


Figure 1 The distribution of expression of all samples before and after normalization.

the control group are presented in [Supplementary Figure 1](#) and [Supplementary Table 2](#). Based on KEGG pathway analysis, the DEGs were significantly enriched in 30 signaling pathways, such as the Toll-like receptor signaling and phagosome pathways.

PPI Network Construction and Module Analysis

The interactions between the proteins expressed from DEGs, which consisted of 85 nodes and 266 edges

(Figure 4A), were constructed from the STRING database and visualized using Cytoscape. In addition, 5 of the top genes with relatively high connectivity degrees (≥ 20) were IL1 β (degree = 23), CXCL1 (degree = 22), ICAM1 (degree = 21), CCL5 (degree = 20), and CCL4 (degree = 20). Four significant modules (Figure 4B) were obtained by module analysis in the PPI network using MCODE from Cytoscape based on the aforementioned criteria. The crucial nodes with a high MCODE score in these four modules were IL1RN, C3AR1, C5AR1, ICAM1, HCAR2, IDO1, CCL5, CXCL9, CXCL1, CCL20,

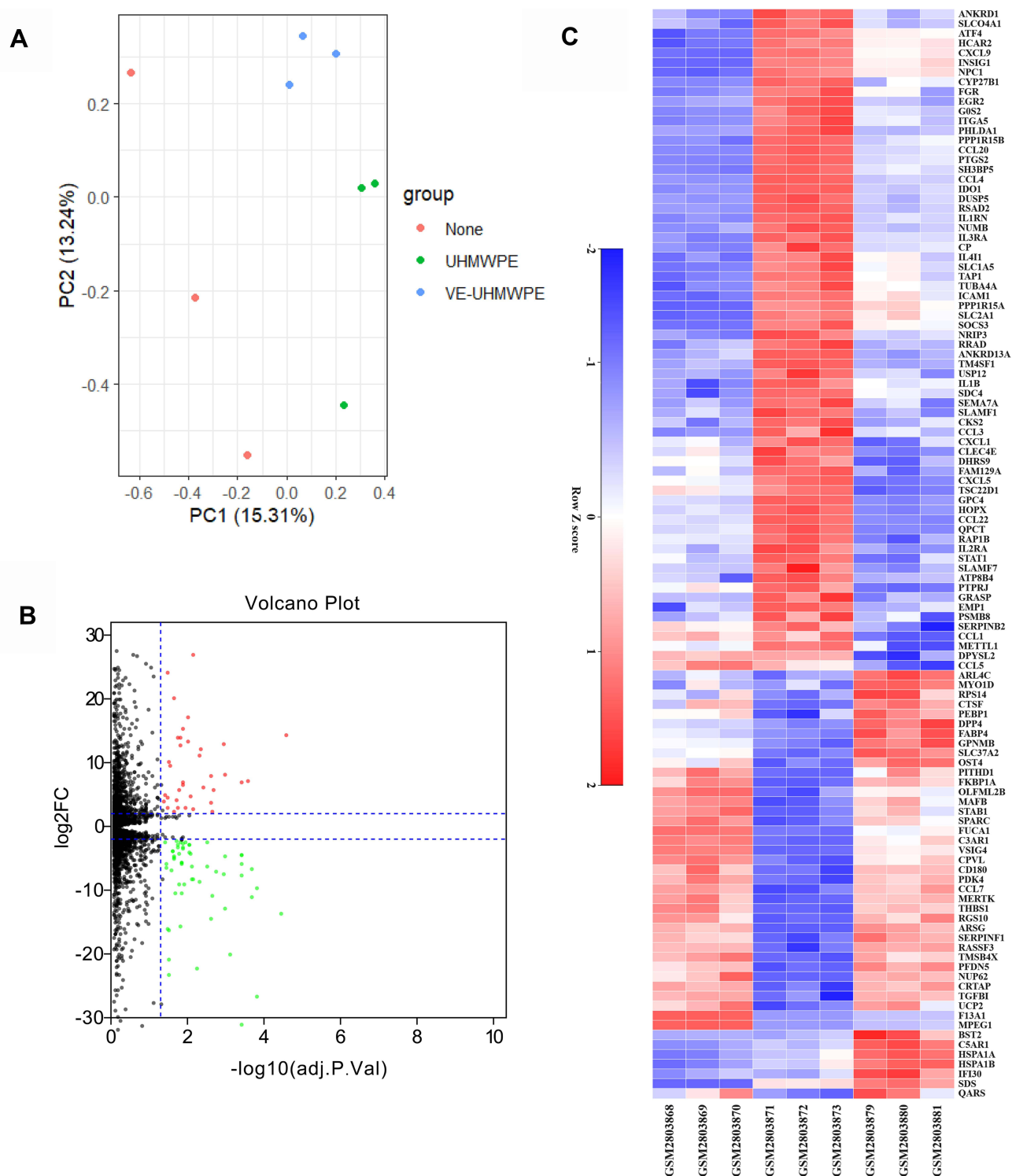


Figure 2 (A) The distribution of expression of samples involving principal component analysis (PCA) for confirming biological variability between different samples; (B) 112 DEGs are shown in the volcano plot, containing 44 upregulated genes in red and 68 downregulated genes in green; (C) the heatmap shows the 112 most significant DEGs. Red indicates relatively high expression, and blue indicates relatively low expression. DEGs were identified by the criteria of $|\log_2\text{fold change (FC)}| > 2$ and $\text{adj. } p\text{-val} < 0.05$. **Abbreviation:** DEGs, differentially expressed genes.

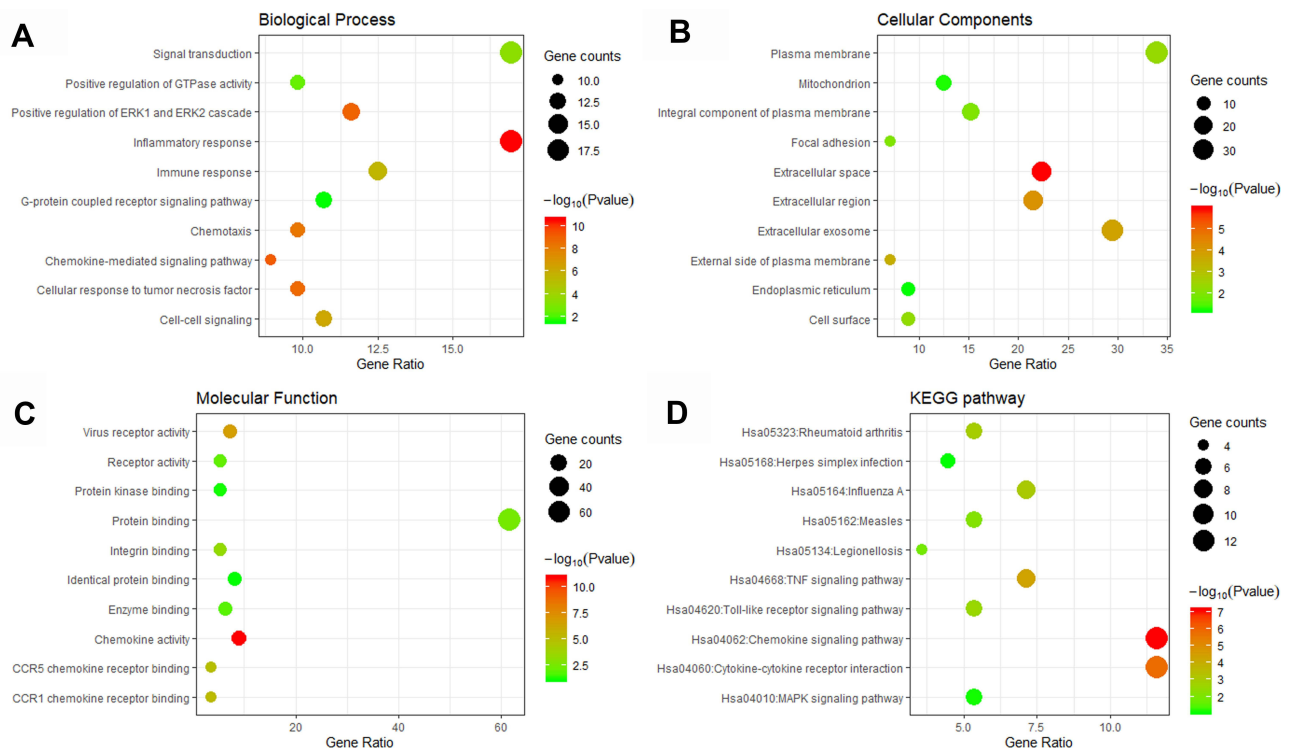


Figure 3 GO including biological process (A), cellular components (B) and molecular function (C) analysis, and KEGG (D) enrichment analysis of the DEGs between the VE-UHMWPE group and the UHMWPE group.

Abbreviations: DEGs, differentially expressed genes; GO, Gene Ontology; KEGG, Kyoto Encyclopedia of Genes and Genomes.

STAT1, CCL4, CCL3, CXCL5, PTGS2, IL1B, and CCL1. The biological process analysis of the top 20 genes was performed and visualized using BiNGO in Cytoscape, which is shown in Figure 5.

Hub Gene Selection

Hub genes were selected by CytoHubba. The top 10 hub genes, which were selected based on the 3 most commonly used classification methods in cytoHubba, are presented in Table 2. By overlapping the first 13 genes, 7 central genes (IL1 β , CXCL1, CCL5, CCL4, CCL20, CXCL9 and C3AR1) were consequently identified as presented in Figure 6A. After being marked on selected modules, we observed that the hub genes were downregulated and gathered in module 1 (Figure 6B).

Construction of the TF-DEG Network Analysis

According to TF binding site data and genetic coordinate position information provided in ENCODE, a potential regulatory network between DEGs and TFs was constructed to analyze the functional roles of selected DEGs. A total of 73 associations between 22 TFs and 13 DEGs were predicted. As

shown, KLF16 regulated 5 DEGs (eg, C5AR1 and CCL20), TFDP1 regulated 5 DEGs (eg, IL2RA and THBS1), and NFIC regulated 4 DEGs (eg, CCL3 and CCL44) (Figure 7).

Construction of miRNA-DEG Network Analysis

The microRNA-DEG pairs were identified through network analysis of 20 DEGs using the TarBase, miRTarBase and miRecords databases. Ultimately, a total of 64 associations between 25 microRNAs and only 18 DEGs were identified, and then the network was visualized in Cytoscape (Figure 8). THBS1 regulated 8 interacting microRNAs, hsa-let-7b-5p, hsa-let-7f-5p, hsa-mir-17-5p, hsa-mir-1-3p, hsa-mir-155-5p, hsa-mir-410-3p, hsa-mir-4302 and hsa-mir-2467-3p. A hub microRNA, hsa-mir-335-5p, was predicted to interact with five DEGs, including PTGS2, ICAMI1, CCL20, C5AR1 and CXCL9.

TF-miRNA Interaction Network

The TF-microRNA interaction network was constructed through network analysis of 20 DEGs in Cytoscape (Figure 9), which included 16 DEGs, 28 TFs, and 13

Table I Top 20 Biological Processes of DEGs

Term	Description	Genes	Gene Counts	P-value
GO:0005515	Protein binding	PTGS2, RRAD, QARS, ANKRD1, SDC4, SLC1A5, SEMA7A, TGFBI, INSIG1, SLC2A1, USPI2, FAMI29A, DPP4, PTPRJ, ICAM1, NRIP3, EGR2, BST2, SOCS3, NPC1, SERPINF1, RPS14, PFDN5, TUBA4A, PEBPI, G0S2, TMSB4X, PPP1R15B, PPP1R15A, ARL4C, EMPI, CCL3, FGR, CXCL9, IFI30, RSAD2, FKBP1A, HSPA1A, HSPA1B, CCL5, CCL4, CCL7, TSC22D1, CCL20, METTL1, TAPI, NUMB, THBS1, GPNMB, PHLDA1, MAFB, IL1RN, DPYSL2, SPARC, STAT1, SLAMF1, PSMB8, SH3BP5, RASSF3, DUSP5, ATF4, NUP62, STAB1, ITGA5, UCP2, HOPX, CKS2, RAPIB, MERTK	69	0.003967971
GO:0005886	Plasma membrane	C3AR1, FGR, IFI30, RRAD, SDC4, GPC4, RGS10, SLC1A5, CLEC4E, SEMA7A, NUMB, SLC2A1, TGFBI, FAMI29A, GPNMB, HCAR2, DPP4, ATP8B4, PTPRJ, ICAM1, C5AR1, IL2RA, BST2, SLC04A1, IL1RN, SLAMF7, SPARC, ANKRD13A, RASSF3, ITGA5, STAB1, SERPINB2, RAPIB, MERTK, ARL4C, IL3RA, EMPI, GRASP	38	0.004446979
GO:0070062	Extracellular exosome	FGR, FKBP1A, SDC4, GPC4, SLC1A5, TGFBI, SLC2A1, IL1B, FAMI29A, THBS1, DPP4, PTPRJ, ICAM1, BST2, IL1RN, MYO1D, DPYSL2, SLAMF1, CPVL, FUCA1, PSMB8, QPCT, NPC1, SERPINF1, RPS14, SLC37A2, PEBPI, TUBA4A, FABP4, RAPIB, CP, VSIG4, CTSF	33	1.62E-04
GO:0005615	Extracellular space	CCL1, CXCL1, ICAM1, CCL3, CXCL5, ARSG, CRTAP, IL1RN, CXCL9, SPARC, CCL5, CCL4, CCL7, GPC4, CCL22, SERPINF1, CCL20, SEMA7A, TGFBI, SERPINB2, IL1B, CP, MERTK, THBS1, CTSF	25	1.06E-06
GO:0005576	Extracellular region	CCL1, CXCL1, CCL3, CXCL5, OLFML2B, F13A1, CXCL9, IFI30, IL4I1, SPARC, CCL5, CCL4, CCL7, NPC1, CCL22, SERPINF1, CCL20, TGFBI, SERPINB2, TUBA4A, IL1B, TMSB4X, CP, THBS1	24	6.67E-05
GO:0006954	Inflammatory response	CCL1, CXCL1, C3AR1, CCL3, IL2RA, C5AR1, CXCL5, PTGS2, CXCL9, CCL5, CCL4, CD180, CCL7, CCL22, CCL20, STAB1, SEMA7A, IL1B, THBS1	19	2.87E-11
GO:0007165	Signal transduction	CCL1, CXCL1, C5AR1, CXCL5, CXCL9, RRAD, SPARC, DPYSL2, SDC4, CCL4, SLAMF1, CCL7, SH3BP5, RASSF3, NPC1, CCL22, CCL20, IL1B, GRASP	19	3.70E-04
GO:0005887	Integral component of plasma membrane	PTPRJ, ICAM1, C3AR1, C5AR1, BST2, SLC04A1, SDC4, CD180, GPC4, SLC1A5, NPC1, STAB1, SLC2A1, TAPI, TM4SF1, MERTK, GPNMB	17	0.0098955
GO:0006955	Immune response	CXCL1, CCL3, IL2RA, C5AR1, CXCL5, IL1RN, CXCL9, CCL5, CCL4, CCL22, CCL20, SEMA7A, IL1B, THBS1	14	2.75E-06
GO:0005739	Mitochondrion	PKD4, RSAD2, QARS, HSPA1A, HSPA1B, SPARC, DPYSL2, SH3BP5, CYP27B1, SDS, RPS14, TAPI, G0S2, PPP1R15A	14	0.05425545
GO:0070374	Positive regulation of ERK1 and ERK2 cascade	CCL1, ICAM1, CCL22, CCL3, C5AR1, CCL20, SEMA7A, RAPIB, CCL5, GPNMB, SLAMF1, CCL4, CCL7	13	1.18E-09
GO:0007267	Cell-cell signaling	CCL22, CCL3, CXCL5, BST2, CCL20, STAB1, CXCL9, IL1B, CCL5, MERTK, CCL4, CCL7	12	6.58E-07
GO:0007186	G-protein coupled receptor signaling pathway	CXCL1, CCL1, C3AR1, CCL22, CCL3, CXCL5, CCL20, CXCL9, CCL5, HCAR2, CCL4, CCL7	12	0.028053381

(Continued)

Table 1 (Continued).

Term	Description	Genes	Gene Counts	P-value
GO:0071356	Cellular response to tumor necrosis factor	CCL1, ICAMI, CCL22, CCL3, CCL20, FABP4, ANKRD1, THBS1, CCL5, CCL4, CCL7	11	1.99E-09
GO:0006935	Chemotaxis	CXCL1, CCL1, C3AR1, CCL22, CCL3, C5AR1, CXCL5, CCL20, CXCL9, CCL5, CCL7	11	5.48E-09
GO:0043547	Positive regulation of GTPase activity	CCL1, ICAMI, RGS10, CCL22, CCL3, IL2RA, CCL20, CCL5, CCL4, CCL7, IL3RA	11	0.00324408
GO:0070098	Chemokine-mediated signaling pathway	CXCL1, CCL1, CCL22, CCL3, CXCL5, CCL20, CXCL9, CCL5, CCL4, CCL7	10	6.60E-10
GO:0008283	Cell proliferation	CXCL1, GPC4, IL2RA, BST2, SERPINF1, TGFBI, INSIG1, CKS2, RAP1B, EMP1	10	5.52E-04
GO:0009986	Cell surface	PTPRJ, ICAMI, C5AR1, BST2, ITGA5, SPARC, THBS1, SDC4, SLAMF1, DPP4	10	0.005226755
GO:0008009	Chemokine activity	CXCL1, CCL1, CCL22, CCL3, CXCL5, CCL20, CXCL9, CCL5, CCL4, CCL7	10	1.50E-11

Abbreviation: DEGs, differentially expressed genes.

microRNAs, with 70 associations between the TFs and DEGs and 19 associations between the miRNAs and DEGs. We separately analyzed the degree of 16 DEGs in the TF-DEG network and the microRNA-DEG

network (Table 3). We found that RELA and NFKB1 regulated six interacting DEGs. Simultaneously, hsa-miR-144, hsa-miR-21, and hsa-miR-221 regulated three interacting DEGs.

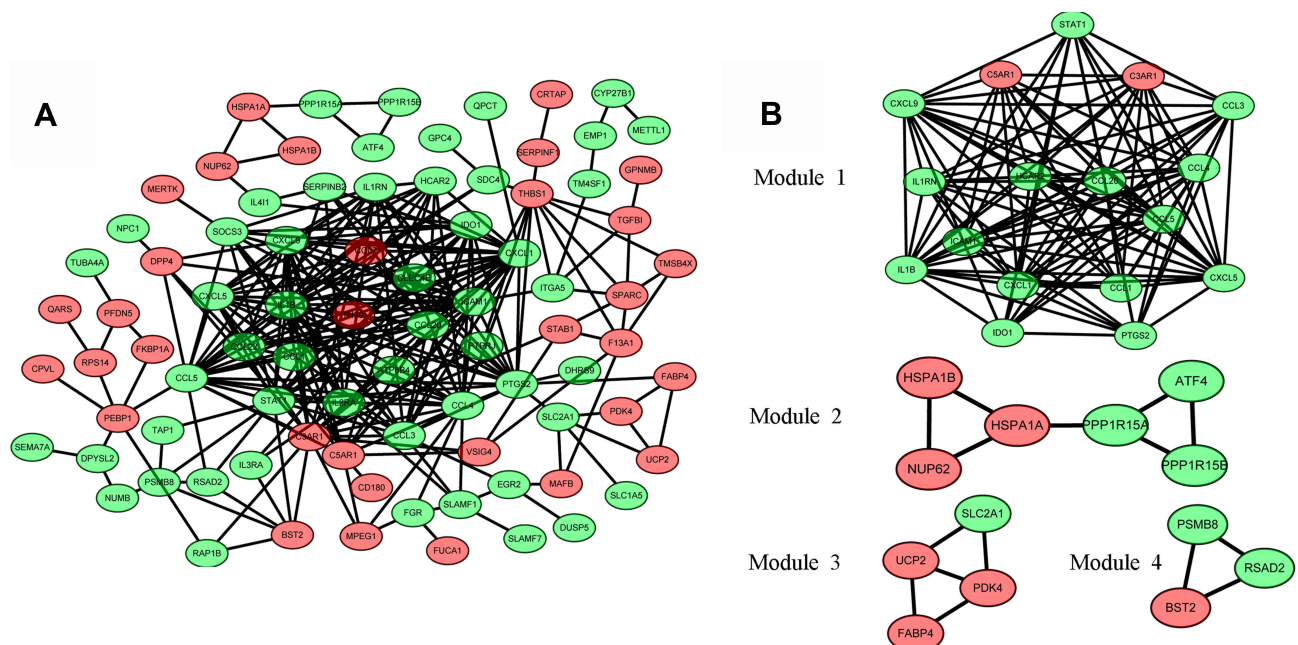


Figure 4 Protein-protein interaction (PPI) network analysis. (A) The PPI network of DEGs visualized in Cytoscape. Red indicates upregulated genes, and green indicates the downregulated genes. (B) Four significant modules in the PPI network.

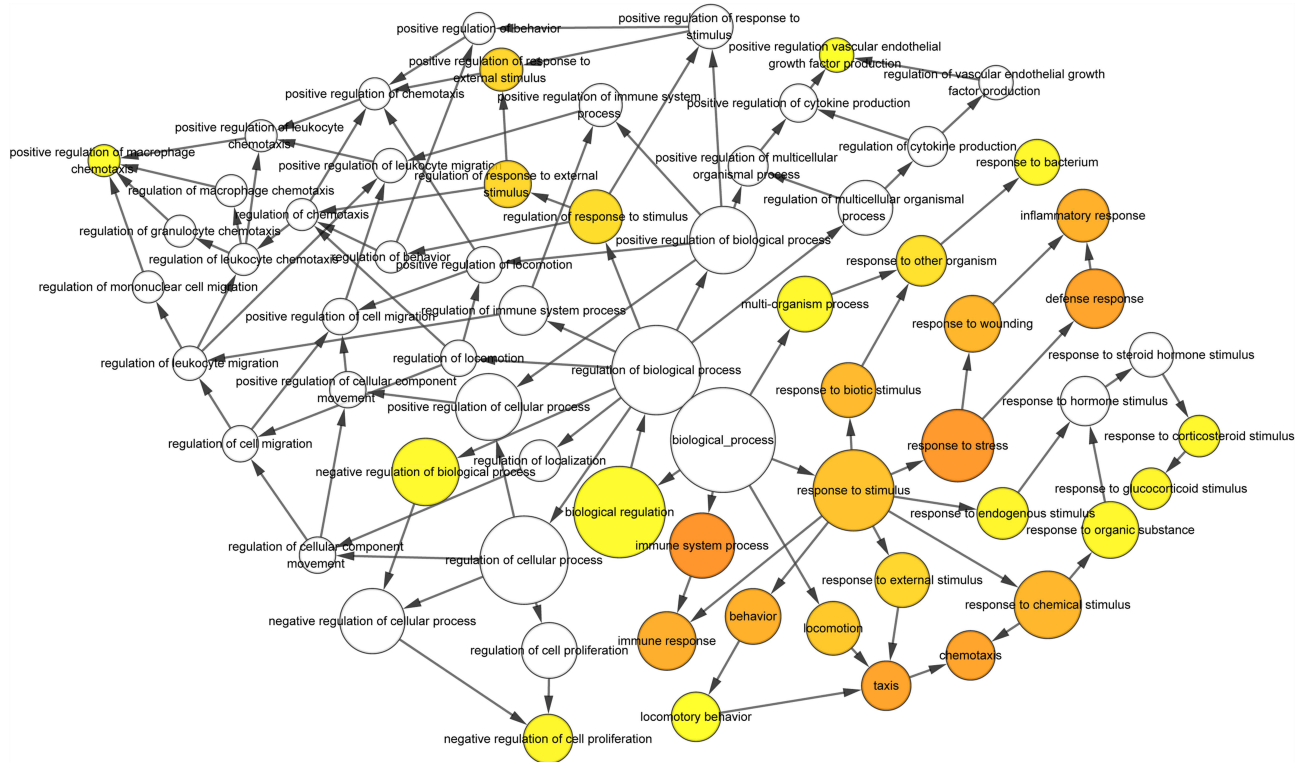


Figure 5 The biological process analysis of the top 20 DEGs was constructed using BiNGO. The color depth of nodes refers to the corrected *p*-value of ontologies. The size of nodes refers to the numbers of genes that are involved in the ontologies. *P* < 0.01 was considered statistically significant.

Assessment of Polyethylene Particles, qRT-PCR and ELISA

As visualized with a transmission electron microscope, VE-UHMWPE particles were engulfed by macrophages (Figure 10A and B). To further validate the transcriptional RNA-seq data, the expression of 5 hub genes (IL1 β , CXCL1, ICAM1, CCL5, and CCL4) with high

connectivity degrees (≥ 20) was examined by qRT-PCR (Figure 10C). Notably, the results of the gene expression examined by qRT-PCR and high-throughput sequencing were highly correlated. Moreover, ELISA revealed a significant increase in the levels of inflammatory factors including IL-1 β , IL-2, IL-6 and TNF- α in the supernatant of the UHMWPE group in comparison with the control groups, while the levels of these inflammatory factors were decreased in the VE-UHMWPE group in comparison with the UHMWPE groups (Figure 10D).

Table 2 List of the Top 10 Hub Genes Selected by MCC, MNC and Degree Methods in cytoHubba

MCC	MNC	Degree
IL1B	IL1B	IL1B
CXCL1	CXCL1	CXCL1
CCL20	ICAM1	ICAM1
CCL4	CCL5	CCL5
CXCL5	CCL4	CCL4
CXCL9	STAT1	STAT1
CCL5	CCL20	C3AR1
C3AR1	CXCL9	CCL20
HCAR2	CXCL5	CXCL9
C5AR1	C3AR1	PTGS2

Abbreviations: MCC, maximal clique centrality; MNC, maximum neighborhood component.

Discussion

Bearing in mind that wear particles continuously released from the surfaces of implant-bearing materials could result in local infiltration and activation of inflammatory cells, an ideal approach to minimize peri-prosthetic osteolysis at the site of implantation should be emphasized to avoid an inflammatory response. As the major source of inflammatory mediators associated with impaired bone metabolism and increased activity of resorptive osteoclasts, macrophages can colonize the site of implantation and initiate an inflammatory cascade.²⁴ Several studies have focused on the

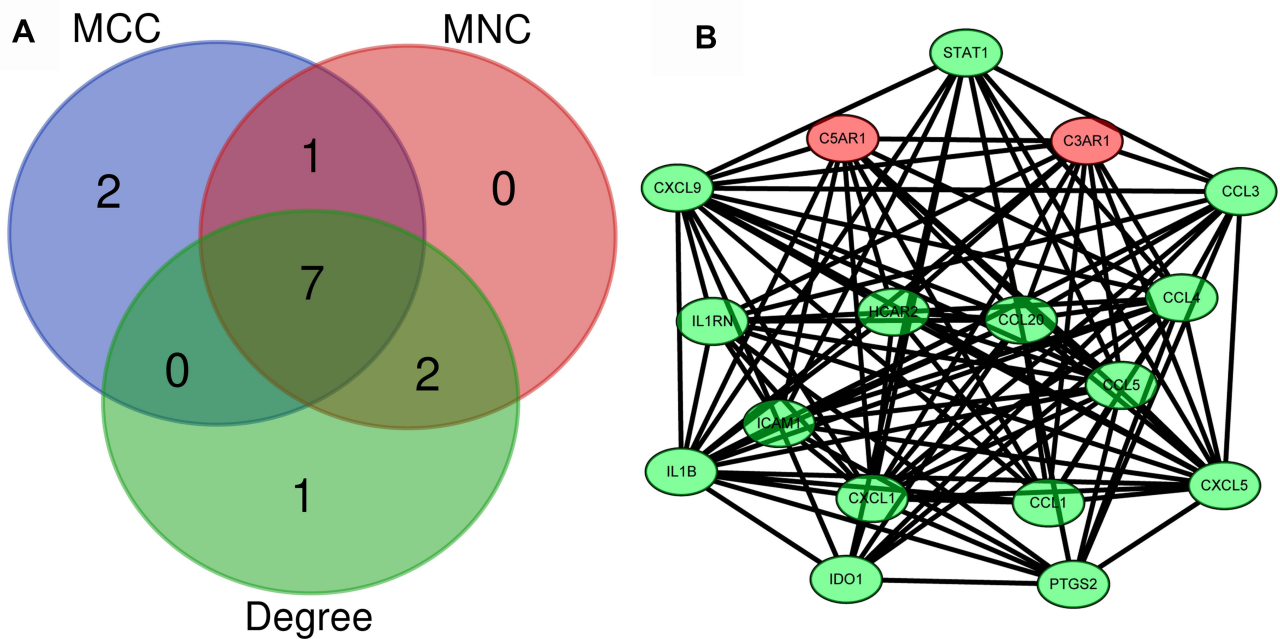


Figure 6 Identification of the hub genes. (A) Hub genes were identified by the overlap of the three methods in CytoHubba; (B) the selected hub genes were all included in the most significant module.

characterization of the immune response of macrophages to implant particles in order to understand the molecular mechanisms of inflammatory osteolysis.^{25–27} Terkawi revealed that the differentiated gene profile of macrophages stimulated by conventional UHMWPE particles mainly involves common gene expression signatures for inflammation and rheumatoid arthritis.¹⁵ By using informatics analysis of the gene profile of macrophages stimulated by UHMWPE and VE-UHMWPE particles, it was revealed that VE-UHMWPE triggered reduced biological activity and inflammatory responses in comparison with UHMWPE. A total of 112 DEGs were identified between the two groups, which may aid in advancing our knowledge on regulatory factors with respect to aseptic loosening of orthopedic implants.

Integrated analysis of differential gene expression indicated that the top 10 DEGs (IL1 β , CXCL1, CCL20, CCL4, CXCL5, CXCL9, CCL5, C3AR1, HCAR2 and C5AR1) in the PPI network were the hub genes affecting the inflammatory response of macrophages to VE-UHMWPE, all showing a decreasing trend compared to the UHMWPE group. We found that most of these DEGs were chemokines (IL1 β , CXCL1, CCL20, CCL4, CXCL5, CXCL9 and CCL5) and three (C3AR1, HCAR2 and C5AR1) were inflammation-

related receptors. These findings agreed with the fact that proinflammatory mediators including SPP1, CCL2, CCL3, CCL4, CCL5 and CCL3 were highly expressed in macrophages cultured with UHMWPE in comparison with the blank control.¹⁵ Importantly, large numbers of DEGs identified between the UHMWPE group and the VE-UHMWPE group also serve regulatory roles in osteoclastogenesis and bone resorption.^{15,28} Some inflammatory diseases are associated with increased bone resorption and fracture rates because inflammatory cytokines produced by innate and adaptive immune cells could not only facilitate inflammation but also activate bone degeneration and inhibit bone formation.²⁹ The degree of inflammation is related to the extent of local and systemic bone loss.³⁰ For instance, chemokines including CCL3, CCL4, CCL5, CXCL9, CCL5 and CCL20 are involved in the remodeling of bone tissue through mediating the development and function of osteoblasts and osteoclasts.^{15,24,31,32} CXCL1 could contribute to osteolysis and facilitate metastasis of prostate cancer and osteosarcoma,^{33,34} while several published results confirmed the role of IL1 β inducing bone resorption and osteoporosis.²⁹ C5AR1 signaling in osteoblasts plays a detrimental role in bone regeneration by interacting

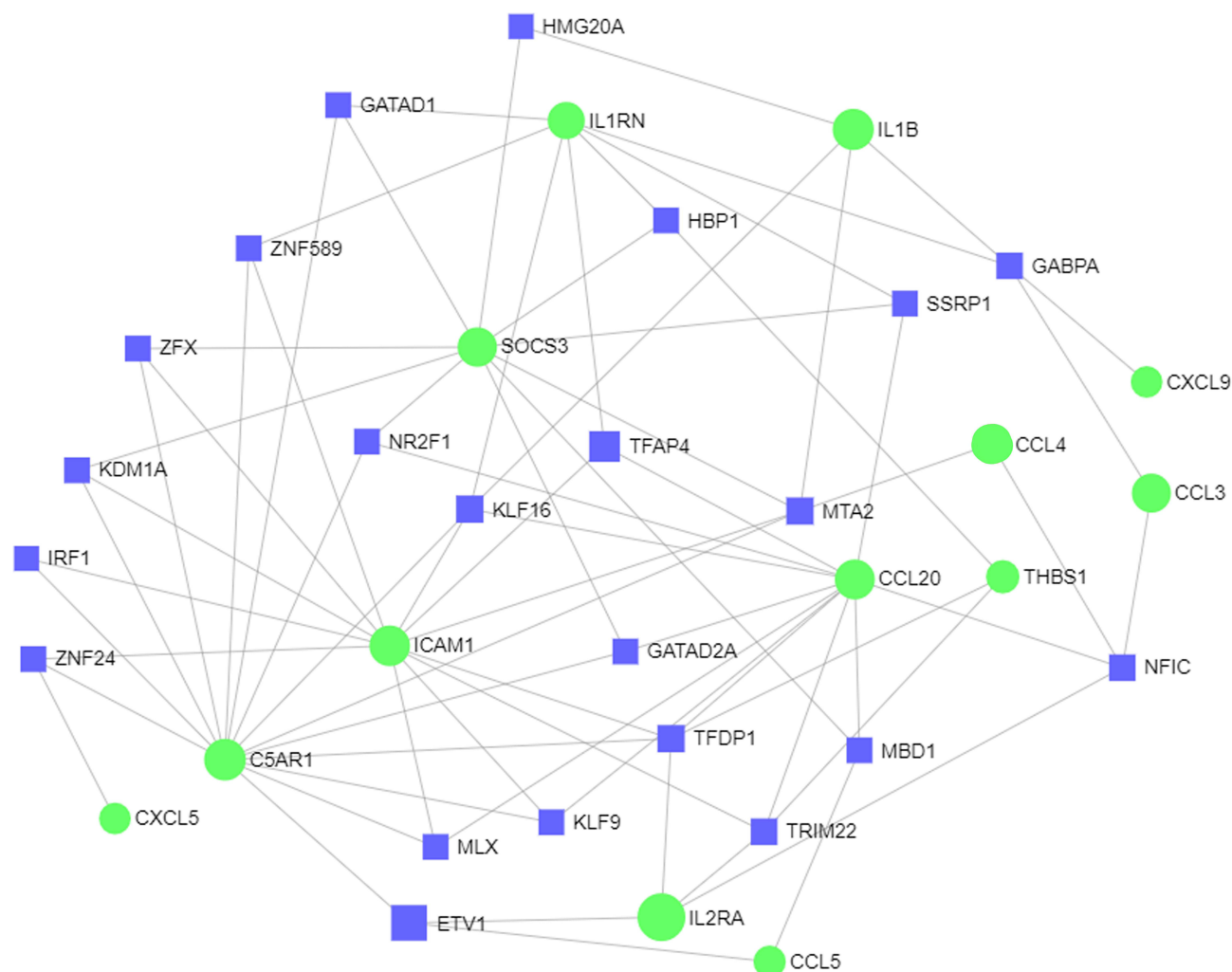


Figure 7 The network of TF-DEG was obtained from the ENCODE database.

Abbreviations ENCODE, Encyclopedia of DNA Elements; TF, transcription factor; DEG, differentially expressed gene.

with TLR2 and stimulating CXCL10.²⁸ The expression of HCAR2 in peripheral blood mononuclear cells was significantly lower in nonunion patients than in those with healed long bone fractures, thus acting as a valuable biomarker for nonunion diagnosis.³⁵ Tian and his colleagues demonstrated that C3AR1 might participate in the osteocyte apoptosis induced by myeloma cells.³⁶ The expression of the top 5 hub genes (IL1 β , CXCL1, ICAM1, CCL5, and CCL4) with connectivity degrees ≥ 20 showed a consistent trend by qRT-PCR assay and high-throughput sequencing. Moreover, ELISA revealed that UHMWPE particles stimulated a significant increase in the levels of IL-1 β , IL-2, IL-6 and TNF- α , which was consistent with

a study by Terkawi et al⁴ and diffusion of vitamin E to UHMWPE could reduce the secretion of these inflammatory factors.

We then used a variety of bioinformatics analysis methods to analyze these DEGs to obtain a comprehensive understanding of these genes. As a result, GO terms of 112 DEGs for biological process categories included positive regulation of the signal transduction process, inflammatory response and immune response. Furthermore, the KEGG pathway enrichment analysis revealed that these DEGs were predominantly associated with the chemokine signaling pathway and cytokine–cytokine receptor interactions. The DEGs in the chemokine signaling pathway were

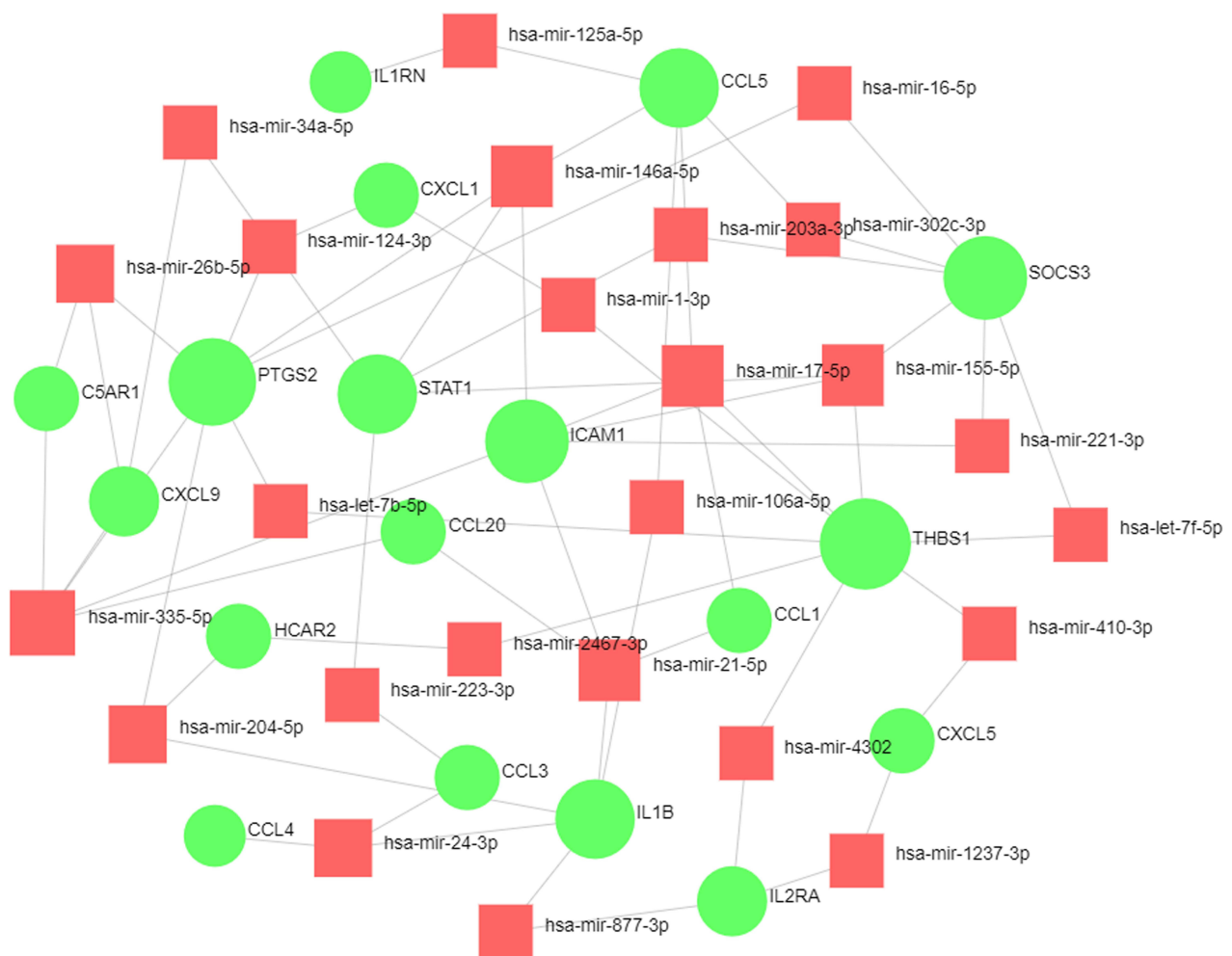


Figure 8 The network of DEG-miRNA was obtained from the tarbase, miRTarBase, and miRecords databases.

Abbreviations: DEG, differentially expressed gene; miRNA, microRNA.

CXCL1, CCL1, CCL3, CXCL5, FGR, CXCL9, STAT1, CCL5, CCL4, CCL7, CCL22, CCL20 and RAP1B, while the DEGs in the cytokine–cytokine receptor interaction pathway were CCL1, CXCL1, CCL3, IL2RA, CXCL5, CXCL9, CCL5, CCL4, CCL7, CCL22, CCL20, IL1B and IL3RA. Nineteen DEGs were involved in the inflammatory response and fourteen were related to the immune response. Our bioinformatics analysis demonstrated the involvement of the rheumatoid arthritis pathway and measles pathway of DEGs in the response of macrophages to VE-UHMWPE and UHMWPE particles. Of significance, the cytokine–cytokine receptor interaction pathway, which was reported to be the most significantly enriched in patients with rheumatoid arthritis and

measles, was the major pathway involved in the response of macrophages to UHMWPE particles.³⁷ Many genes that were downregulated in macrophages encountering VE-UHMWPE particles in comparison with the UHMWPE group such as IL-1 α , IL-27, CCL3, CCL20, SOCS3 and CCL5 were abundantly expressed in rheumatoid synovial tissues and played substantial roles in the pathogenesis of rheumatoid arthritis.^{38,39} There is overwhelming evidence that macrophages serve a prominent role in the pathogenesis of rheumatoid arthritis.³⁸ They are the most abundant cell type in the rheumatoid synovium and can release an array of inflammatory cytokines resulting in the recruitment of other inflammatory cells, T-cell polarization, fibroblast activation and osteoclast

Table 3 Co-DEGs Regulated by TF and miRNAs

TF	DEGS	Gene Counts
RELA	Socs3, Ptgs2, Icam1, Cxcl1, Ccl5, Ccl20	5
NFKB1	Socs3, Ptgs2, Icam1, Cxcl1, Ccl5, Il1rn	5
SPI	Ptgs2, Icam1, Cxcl1, Ccl20	4
REL	Ptgs2, Icam1, Cxcl1, Ccl5	4
JUN	Ptgs2, Icam1, Cxcl1, Ccl5	4
TP53	Ptgs2, Icam1, Thbs1	3
SPII	Icam1, Ccl5, Il1rn	3
5915	Ptgs2, Icam1, Il1rn	3
NFKB2	Ptgs2, Icam1, Ccl5	3
CTCF	Ptgs2, Thbs1, Ccl20	3
CREB1	Ptgs2, Icam1, Ccl5	3
CEBPB	Ptgs2, Icam1, Ccl5	3
STAT1	Socs3, Icam1	2
USF1	Ptgs2, Thbs1	2
STAT5A	Socs3, Il1rn	2
SRF	Ptgs2, Thbs1	2
MYC	Icam1, Ccl20	2
FOS	Ptgs2, Ccl5	2
EGR1	Ccl5, Thbs1	2
CEBPD	Ptgs2, Ccl5	2
CEBPA	Ptgs2, Ccl5	2
ATF1	Ptgs2, Ccl5	2
STAT3	Socs3	1
TFAP2A	Ptgs2	1
STAT2	Socs3	1
STAT5B	Socs3	1
IRF5	Ccl5	1
miRNA	DEGS	Gene counts
hsa-miR-144	Ptgs2, Thbs1, Ccl20	3
hsa-miR-21	Icam1, Thbs1, Ccl20	3
hsa-miR-221	Socs3, Icam1, Thbs1	3
hsa-let-7b	Ptgs2, Thbs1	2
hsa-miR-17	Icam1, Thbs1	2
hsa-miR-98	Ccl5, Thbs1	2
hsa-let-7a	Thbs1	1
hsa-miR-203	Socs3	1
hsa-miR-30e	Socs3	1
hsa-miR-590-5p	Ccl20	1

Abbreviations: DEGs, differentially expressed genes; miRNAs, microRNAs.

of RELA and NFKB1 were enriched predominantly with respect to terms such as protein binding, inflammatory response, immune response, chemotaxis, chemokine-mediated signaling pathway, chemokine activity, etc. Thus, RELA and NFKB1 may be new drug targets for regulating inflammatory osteolysis in orthopedic implants. Additionally, the most connected microRNA node

identified was hsa-miR-335-5p, which targeted five DEGs (PTGS2, ICAM1, CCL20, C5AR1, and CXCL9). A series of studies have demonstrated that an increased level of miR-335-5p could facilitate bone formation, which may also be related to the decreased potential of osteoclastogenesis.^{46,47} Meanwhile, microRNA-21 could act as a functional factor in the pathogenesis of particle-induced osteolysis.⁴⁸ Although there is no direct evidence to prove the role of hsa-miR-144 and hsa-miR-221 in the pathogenesis of inflammatory osteolysis until now, the target genes of these microRNAs such as ICAM1 and CCL20 are considered candidate genetic markers associated with the inflammatory process.

The current study provided several advances in the field. At present, a systematic bioinformatics analysis on the different performances of VE-UHMWPE and UHMWPE in the pathological process of osteolysis is still lacking. We performed a rigorous informatics analysis based on multiple databases and tools and provided results on this topic as comprehensively as possible, proposing multiple key genes, transcription factors, and miRNAs that may play important roles in the process of reduction of inflammatory osteolysis by VE-UHMWPE. These findings could provide research directions and molecular targets for further basic and clinical trials. However, it should be admitted that the current investigation is only fundamental research. Although multiple hub genes, key miRNAs and TFs were selected, the detailed functions of these molecules in inflammatory osteolysis should be further identified by in vivo and in vitro studies. Currently, whether the addition of vitamin E to UHMWPE could decrease the incidence of aseptic loosening of orthopedic implants has not been verified by randomized controlled trials. Analyses of protein expression and changes in osteoclastogenesis in the VE-UHMWPE and UHMWPE groups are warranted to further explore the roles of these materials in the pathogenesis of inflammatory osteolysis of orthopedic implants.

Conclusion

Combined with various informatics analyses and in vitro experiments, IL1 β , CXCL1, ICAM1, CCL5, and CCL4 may serve as prognostic and therapeutic targets of inflammatory osteolysis in orthopedic devices. Additionally, the TF-microRNA interaction regulatory network revealed that key microRNAs (hsa-miR-144, hsa-miR-21, and hsa-miR-221) and TFs (RELA and NFKB1) were important regulators at the transcriptional level in this process. Further experiments are required to probe the functions of these genes in the

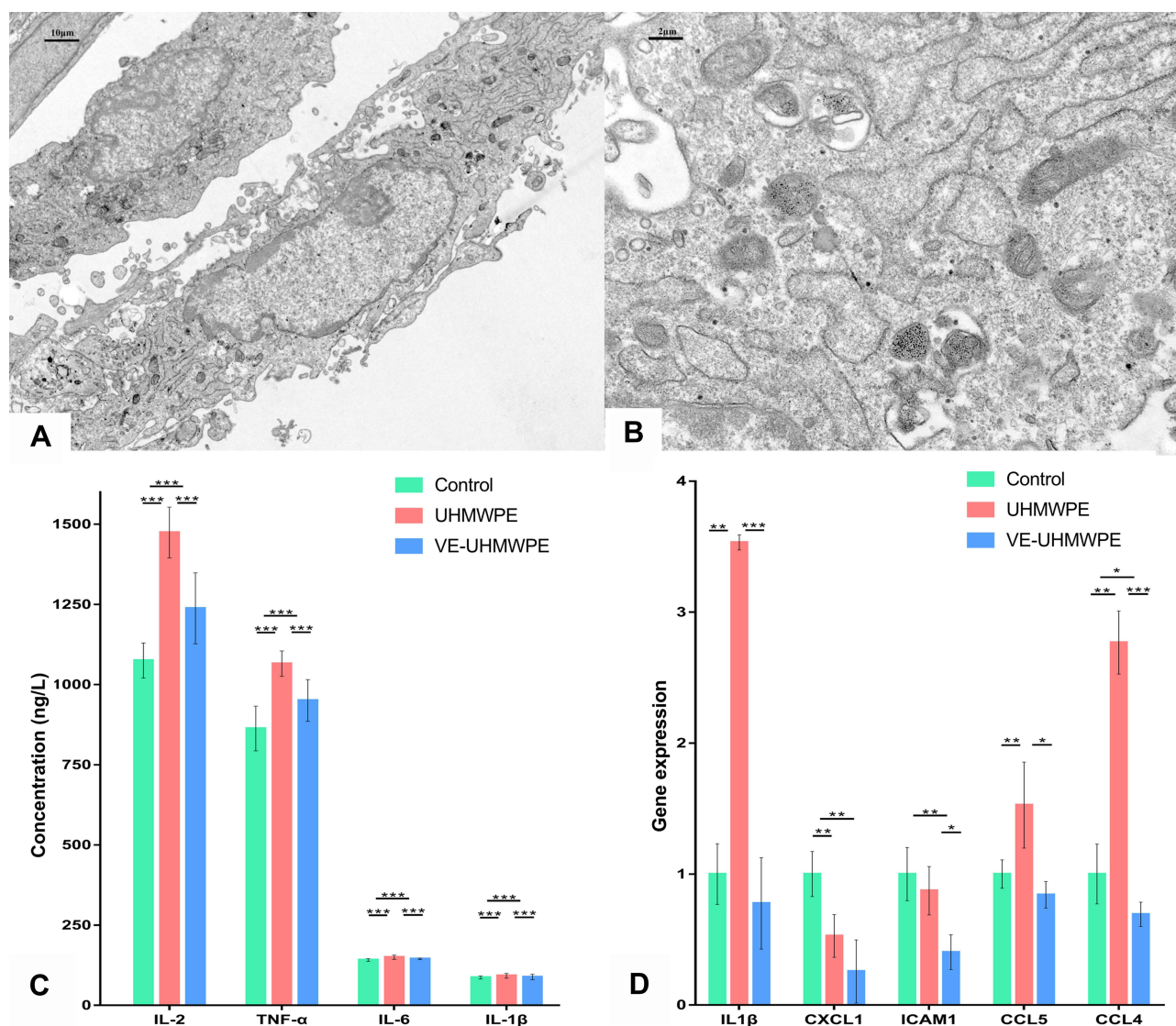


Figure 10 Macrophage intervention, qRT-PCR and ELISA. **(A)** Macrophage-engulfed VE-UHMWPE particles observed under a transmission electron microscope (3000 \times magnification); **(B)** macrophage-engulfed VE-UHMWPE particles observed under transmission electron microscope (15,000 \times magnification); **(C)** comparison of the top five hub genes with high connectivity degree by qRT-PCR and **(D)** comparison of inflammatory factors in supernatant by ELISA. * p -value < 0.05, ** p -value < 0.01 and *** p -value < 0.001.

pathogenesis of inflammatory osteolysis of orthopedic implants.

Data Sharing Statement

All data generated or analysed during this study are included in this published article.

Ethics Approval

This investigation involved the THP-1 cell lines that were purchased commercially from American Type Culture Collection in the USA and no further approval was required.

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Disclosure

The authors declare there are no conflicts of interest regarding the publication of this paper.

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