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Transcriptional analysis of long non-coding RNAs in facet joint osteoarthritis†

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It is recognized that facet joint osteoarthritis (FJOA) is commonly induced by the degeneration of articular cartilage of the facet joint. However, the specific pathological mechanisms underlying facet joint osteoarthritis has not yet been elucidated. To obtain the differential expression patterns and putative functions of long noncoding RNAs (lncRNAs) in FJOA, in the current study, we detected the expression levels of lncRNAs in patients with varying degrees of facet cartilage degeneration (control group: normal or mild facet cartilage degeneration; FJOA: moderate to severe facet cartilage degeneration) by RNA deep sequencing. Differentially expressed lncRNAs were screened and the accuracy of sequencing data was further validated by using quantitative reverse transcription-polymerase chain reaction (qRT-PCR). Target genes of differentially expressed lncRNAs were predicted by antisense and/or *cis*-regulatory module prediction. Predicted target genes were further analyzed by Gene Ontology (GO) and Kyoto Enrichment of Genes and Genomes pathway analysis (KEGG) to discover enriched cellular component, molecular function, biological process, and signaling pathways. Our results provided a general view of the expression changes of lncRNAs in FJOA and thus might facilitate the illumination of the underlying mechanisms of FJOA.

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Introduction

Facet joint osteoarthritis (FJOA) is a common disease that causes low back and lower extremity pain, especially severe low back pain in the morning.^{1–3} FJOA is generally diagnosed by physical examinations and imaging, including radiographs, magnetic resonance imaging, computed axial tomography scanning, single photon emission scanning, and radionuclide bone scanning.⁴ Common treatments of this disease contain nonsteroidal anti-inflammatory drugs, steroid medications, muscle relaxers, and physical therapies.^{5,6} It has been demonstrated that FJOA is mainly caused by the degeneration of articular cartilage of the facet joint. But we know little about the special underlying mechanisms of FJOA, especially the molecular changes of FJOA. To fill this gap of knowledge, it is necessary to obtain a global perspective of genetic changes of FJOA.

Non-coding RNAs are a group of RNA molecules that do not translate into proteins. Previously, non-coding RNAs were considered as non-functional RNAs or even junk RNAs. Nowadays, it has been showed that non-coding RNAs, especially microRNAs (miRNAs) and long non-coding RNAs (lncRNAs), are

important transcriptional and post-transcriptional regulators.^{7–9} lncRNAs are non-coding RNA transcripts with greater than 200 nucleotides in length.¹⁰ lncRNAs play essential roles in genetic regulation as well as multiple epigenetic regulations such as genetic imprinting, histone modification, X-chromosome inactivation, and chromatin dynamics.^{11,12} Emerging studies showed that lncRNAs are involved in a variety of diseases, including cancer, metabolic disease, cardiovascular disease, neurodegenerative disorder, and immune system dysfunction.^{13–15}

In view of the importance of lncRNAs, in the current work, RNA deep sequencing analysis was performed and the expressions of lncRNAs in human facet joints were determined and validated. Differentially expressed lncRNAs were discovered and antisense or *cis*-regulatory module predicted target genes of these differentially expressed lncRNAs were enriched by Gene Ontology (GO) and Kyoto Enrichment of Genes and Genomes (KEGG) pathway analysis (Fig. 1).

Materials and methods results

RNA deep sequencing

The current work was approved by the Human Ethics Committee of No. 2 People Hospital Affiliated to Nantong University and participating patients signed the informed consent agreement. Control and morbid human facet joint samples were collected from patients with vertebral fracture

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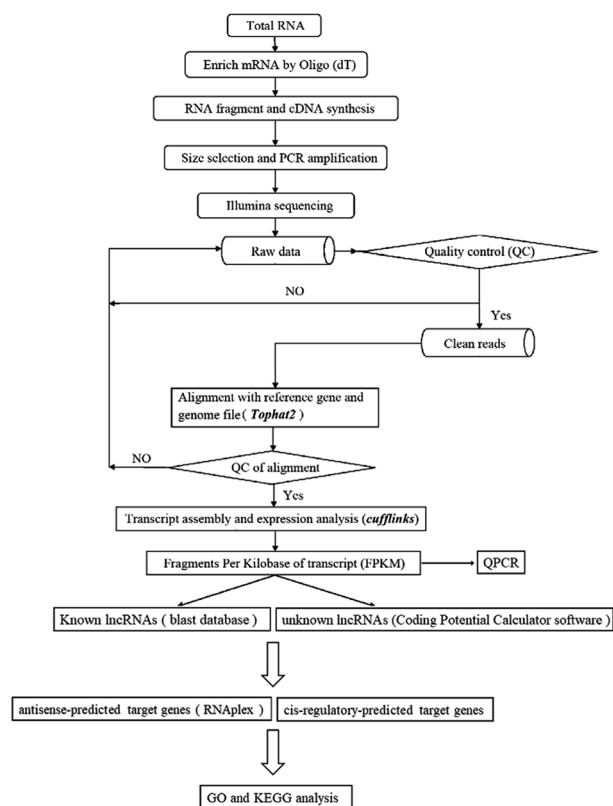


Fig. 1 Schematic of the experimental protocols of deep sequencing and bioinformatic analysis.

(CTRL1, CTRL2, and CTRL3) and patients with FJOA (FJOA1, FJOA2, and FJOA3) separately. RNAs were isolated from facet joint samples, purified to remove contaminating DNA, and then sequenced by using Illumina HiSeq. For strand-specific library construction and sequencing, after total RNA was extracted, rRNAs were removed to retain mRNAs and ncRNAs. The enriched mRNAs and ncRNAs were fragmented into short fragments by using fragmentation buffer and reverse transcribed into cDNA with random primers. Second-strand cDNA were synthesized by DNA polymerase I, RNase H, dNTP (dUTP instead of dTTP) and buffer. Next, the cDNA fragments were

purified with QiaQuick PCR extraction kit, end repaired, poly(A) added, and ligated to Illumina sequencing adapters. Then UNG (Uracil-*N*-Glycosylase) was used to digest the second-strand cDNA. The digested products were size selected by Agarose gel electrophoresis, PCR amplified, and sequenced using Illumina HiSeq X Ten by Gene Denovo Biotechnology Co. (Guangzhou, China). Then for library examination, RNA concentration of library was measured using Qubit® RNA Assay Kit in Qubit® 2.0 to preliminary quantify and then dilute to $1 \text{ ng } \mu\text{l}^{-1}$. Insert size was assessed using the Agilent Bioanalyzer 2100 system (Agilent Technologies, CA, USA), and after the insert size consistent with expectations, qualified insert size was accurate quantitative using Taqman fluorescence probe of AB Step One Plus Real-Time PCR system (Library valid concentration $> 2 \text{ nM}$). For library clustering and sequencing, the qualified libraries were sequenced by an Illumina HiSeq 2500 platform and generate 50 bp single-end reads.

Identification and quantitative analysis of lncRNAs

Sequencing raw data were subjected to quality filter and filtered clean data were quantitative analyzed by expected number of fragments per kilobase of transcript sequence per million base pairs sequenced (FPKM). Transcripts with class code of "I, j, x, u, o" and a length $\geq 200 \text{ bp}$ were screened. Known lncRNAs were identified by comparing screened transcripts with blast database and novel lncRNAs were identified by filtering transcripts with coding potential using Coding Potential Calculator (CPC) software.

qRT-PCR

Isolated RNAs were also subjected to qRT-PCR validation. RNAs were reverse-transcribed to cDNA by using a Prime-Script RT reagent Kit (TaKaRa, Dalian, China). qRT-PCR was conducted by using SYBR Premix Ex Taq (TaKaRa) on an ABI system (Applied Biosystems, Foster City, CA). The primers used in this study were as listed in Table 1. The reliability of primer sets and the quality of qRT-PCR experiments were validated by a single peak melt curve representing a single PCR product. The expression levels of target lncRNAs were calculated by using the $\Delta\Delta C_t$ method with GAPDH as the reference.

Table 1 Primers of real-time PCR

Primer names	Primer (5' to 3')	Sequence	Annealing temperature (°C)
lncRNA MIR22HG	Forward	GCTGCTTCCCCATCATCTG	50.0
	Reverse	TCTCCAACCTTGCCCAAAACG	50.0
lncRNA RERG-AS1	Forward	ATCTGTTCCTGCCTCTGTC	50.0
	Reverse	AACACCATCTGCAAGCCAAG	50.0
lncRNA HECTD2-AS1	Forward	TCAGGGACAGTGGTTGCTTT	50.0
	Reverse	ACCGCAAATGTCGAGTGTTT	50.0
XLOC_091678	Forward	AACTTTGCCCGGAGAAAACG	50.0
	Reverse	TCCCGTTCAAATTAACCGC	50.0
linc01783	Forward	GGTCCCCTTTGCTGATTGAG	50.0
	Reverse	TACTCCACCTGCTGTCCCTTG	50.0
XLOC_091845	Forward	GTTTGCTGCATACCCTAGC	50.0
	Reverse	ACAGCTCCCTCAACGTCTTT	50.0

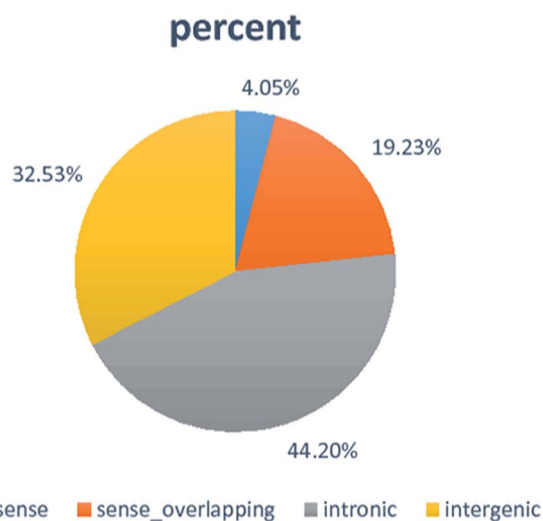


Fig. 2 The percentage of lncRNA classification. The percentages of antisense, intergenic, intronic, and sense overlapping, were listed.

Prediction and functional analysis of target genes of lncRNAs

The potential target genes of lncRNAs were predicted by anti-sense prediction or *cis*-regulatory module prediction. For anti-sense prediction, RNAplex software was applied to calculate the free energy between genes and to predict complementary association of antisense lncRNAs and mRNAs. For *cis*-regulatory module prediction, adjacent coding genes of lncRNAs (10k upstream and 10k downstream) were selected. Target genes of significantly differentially expressed lncRNAs were subjected to GO and KEGG bioinformatic analysis.

Statistical analysis

Statistical analysis was performed by using SPSS for Windows 11.0.1 (SPSS Inc., Chicago, IL). Student's *t*-test was used for statistical comparison and *p*-value < 0.05 was considered as significant different.

Results

Identification of lncRNA in facet joints

Clean data obtained from RNA deep sequencing were filtered to screen candidate lncRNA. A total of 2839 known lncRNAs were identified in control facet joint samples and/or FJOA samples. Moreover, a total of 35 099 novel lncRNA were newly discovered. All identified lncRNAs were listed in Table S1.† Moreover, screened lncRNAs were quantitatively analyzed by cuffdiff software (<https://cole-trapnell-lab.github.io/cufflinks/cuffdiff/index.html>) and FPKM reading of lncRNAs in facet joints were obtained to determine lncRNA expressions.

The expression levels of lncRNAs in FJOA samples were then compared with their expression levels in control facet joint samples. lncRNAs that obtained a \log_2 ratio ≤ -1 or ≥ 1 and a *q*-value ≤ 0.05 were defined as significantly differentially expressed. A total of 8506 lncRNAs were found to be differentially expressed while 650 lncRNAs were up-regulated in FJOA samples and 7856 lncRNAs were down-regulated (Fig. 2). The expression patterns of all these differentially expressed lncRNAs were listed in Table S2† and displayed in a heatmap (Fig. S1†). Top 10 up-regulated and down-regulated lncRNAs were listed in Table 2. These lncRNAs were further categorized into antisense,

Table 2 Top 10 differentially expressed lncRNAs^a

lncRNA	Locus	\log_2 ratio	<i>p</i> -value
Up-regulated			
XLOC_010753	chr10: 30108870–30110823	13.61421521	0.0016
XLOC_016219	chr11: 49359512–49359740	13.43758304	5.00×10^{-5}
XLOC_119379	chrX: 48040578–48040811	11.9579835	0.00245
XLOC_108373	chr8: 10052458–10052689	11.54294963	0.0024
XLOC_092577	chr6: 80727431–80727681	11.09226162	0.0018
XLOC_021340	chr12: 4995659–4995913	10.94871034	0.00215
XLOC_036022	chr15: 72624890–72625144	10.80519571	0.00225
XLOC_007580	chr1: 118757149–118757399	10.70536456	0.0024
XLOC_043063	chr17: 14360831–14361098	10.48474235	0.0028
XLOC_020221	chr11: 93224734–93224998	10.29314974	0.0019
Down-regulated			
XLOC_030524	chr14: 21712360–21712741	–13.35201583	5.00×10^{-5}
XLOC_086316	chr5: 91484799–91485000	–13.10078387	0.0016
XLOC_073826	chr3: 189505508–189505713	–13.05758058	0.0022
XLOC_056381	chr2: 128837149–128837348	–13.02509803	5.00×10^{-5}
XLOC_070170	chr3: 25127728–25127927	–12.98960205	0.00255
XLOC_089811	chr5: 123342227–123342430	–12.97417077	0.0016
XLOC_088041	chr5: 6201207–6201406	–12.96810155	0.00245
XLOC_100362	chr7: 45784516–45784716	–12.92429292	0.00255
XLOC_083411	chr4: 121799502–121799703	–12.84602312	0.00245
XLOC_114984	chr9: 99395260–99395462	–12.79600309	0.00255

^a The top 10 up-regulated and down-regulated lncRNAs were listed.

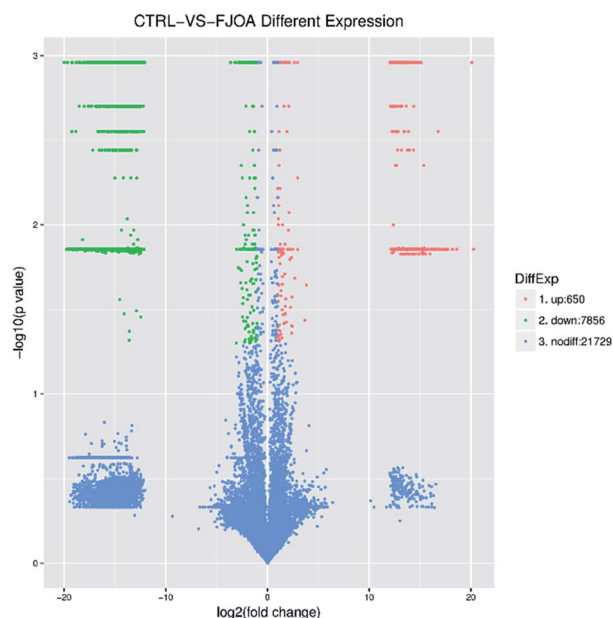


Fig. 3 The volcano plot of differentially expressed lncRNAs. Up-regulated lncRNAs were labeled in red and down-regulated lncRNAs were labeled in green.

intergenic, intronic, and sense overlapping subtypes according to their genomic locations (Fig. 3).¹⁶

qRT-PCR validation

After the identification of lncRNA, qRT-PCR was performed to validate RNA deep sequencing results. A total of 6 lncRNAs (lncRNA MIR22HG, lncRNA RERG-AS1, lncRNA HECTD2-AS1, XLOC_091678, linc01783, and XLOC_091845) that were expressed in both the control facet joint samples and FJOA samples were randomly selected for validation. The characteristics of these lncRNAs are shown in Table 3. The expression levels of these 6 lncRNAs in both control facet joint samples and FJOA samples were determined. Result from qRT-PCR showed that the expression patterns of these lncRNAs were mainly in consistent with their expression patterns observed from RNA deep sequencing (Fig. 4). Both FJOA groups and control groups had three independent samples and all the samples were processed in triplicate.

Table 3 The characteristics of lncRNAs for validation^a

lncRNA	Locus	log ₂ ratio	q-Value	Strand	Class
Up-regulated					
lncRNA MIR22HG	chr17:1711504–1716272	1.20	1.09 × 10 ⁻³	–	Intergenic
lncRNA RERG-AS1	chr12:15152468–15155316	1.63	1.98 × 10 ⁻³	–	Antisense
lncRNA HECTD2-AS1	chr10:91415256–91417528	1.06	1.39 × 10 ⁻²	+	Antisense
Down-regulated					
XLOC_091678	chr6:27688190–27688522	–1.09	1.09 × 10 ⁻³	+	Intergenic
linc01783	chr1:16534514–16535347	–1.92	1.09 × 10 ⁻³	–	Intergenic
XLOC_091845	chr6:31820654–31821480	–1.35	1.09 × 10 ⁻³	+	Intergenic

^a lncRNA, Locus, log₂ ratio, q-value, strand, class were listed.

Functional enrichment analysis of antisense-predicted target genes

Antisense lncRNAs may regulate gene silencing, gene transcription, and mRNA stability by binding to sense mRNA. Therefore, based on the complementary pairing of lncRNAs and genes, potential target genes of lncRNAs were discovered by antisense prediction. Antisense-predicted target genes of differentially expressed lncRNAs were then analyzed by GO terms to find possible biological processes, molecular functions, and cellular components in FJOA (Fig. 5A). GO biological process analysis showed that ontologies related with transport (negative regulation of transport, metal ion transport, ion transport, and inorganic ion transmembrane transport) occupied a large proportion of enriched GO terms. In agreement with GO biological process outcomes, GO molecular function analysis showed that top enriched ontologies were generally related with transporter activity. GO cellular component analysis showed that plasma membrane-related ontologies were significantly enriched. KEGG pathway analysis was also performed to find possible signaling pathways in FJOA (Fig. 5B). However, only 8 signaling pathways were enriched and only one gene were involved in each signaling pathway.

Functional enrichment analysis of cis-regulatory module-predicted target genes

Considering that lncRNAs also target and regulate their adjacent mRNAs, potential target genes of lncRNAs were discovered by *cis*-regulatory module prediction. Similar as antisense-predicted target genes, target genes of differentially expressed lncRNAs predicted by *cis*-regulatory module were investigated by GO and KEGG analysis (Fig. 6). GO biological process analysis revealed that ontologies related with cellular and organismal development, including system development, single-organism developmental process, organ development, multicellular organismal development, and developmental process, were top enriched. Enriched GO molecular function terms were related with molecular binding and signaling cascade activity and enriched GO cellular component terms were related with multiple organelles (Fig. 6A). KEGG pathway analysis demonstrated that many disease-related signaling pathways (tuberculosis, salmonella infection, renal cell carcinoma, pathways in

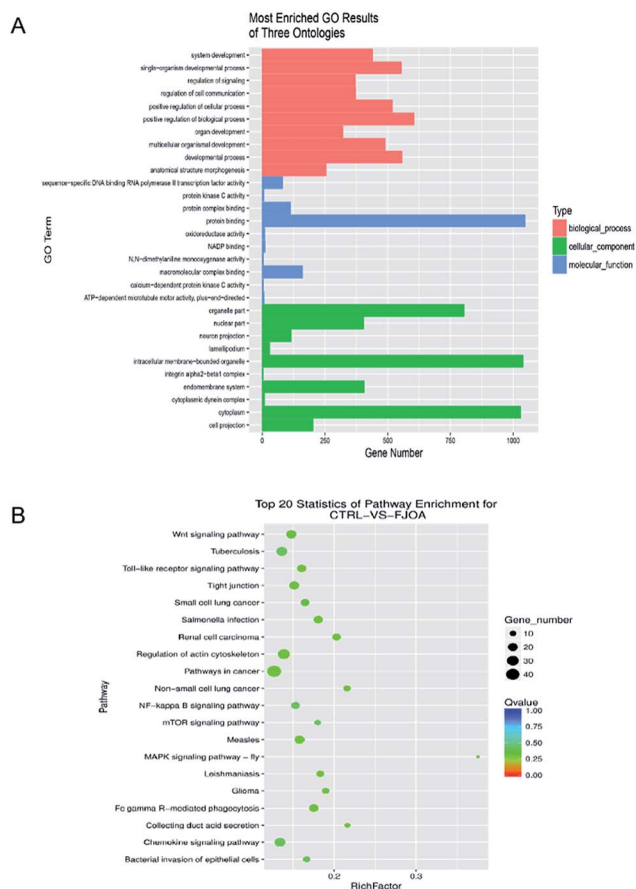


Fig. 6 (A) GO cellular component, molecular function, and biological process of *cis*-regulatory module predicted target genes. (B) Top enriched KEGG pathways of *cis*-regulatory module predicted target genes.

a limited sample size (sample size = 3), the accuracy of predicted potential target genes of lncRNA might not be high.

For the *cis*-effect mechanism, it is suggested that the biological functions of lncRNAs were performed by regulating their neighboring protein coding genes. Therefore, we also predicted the potential target genes of lncRNAs by *cis*-regulatory module prediction and analyzed the functions of these potential target genes by GO and KEGG analysis. Enriched GO terms and KEGG pathways were quite different from the function analysis of potential target genes predicted by antisense prediction. Notably, many enriched KEGG pathways were related to immune response. In our previous study, we analyzed differentially expressed mRNAs and classified these differentially expressed mRNAs into KEGG pathways. Our previous obtained bioinformatic data suggested that many immune and inflammation-related signaling pathways, including B cell receptor signaling pathway, primary immunodeficiency, Fc gamma R-mediated phagocytosis, natural killer cell mediated cytotoxicity, T cell receptor signaling pathway, Wnt signaling pathway, NF-kappa B signaling pathway, leukocyte trans-endothelial migration, Fc epsilon RI signaling pathway, NOD-like receptor signaling pathway, and phagosome, were among the top 30 activated signaling pathways.¹⁹ Many signaling

pathways, including Wnt signaling pathway, NF-kappa B signaling pathway, and Fc gamma R-mediated phagocytosis, were significantly involved in both differentially expressed mRNAs and lncRNAs, suggesting the importance of immune response in FJOA. Recently, it has been demonstrated that neighboring genes can also be regulated by antisense lncRNAs.²⁷ This largely increased the complexity of the regulatory mechanisms of lncRNAs and the subsequent function study of target genes of lncRNAs.

In summary, in the current work, we systematically identified lncRNAs in facet joints, discovered differentially expressed lncRNAs in FJOA, and predicted target genes and biological functions of these differentially expressed lncRNAs. Our work provided a preliminary overview of lncRNAs in FJOA and might benefit the further understanding of the mechanisms of FJOA.

Conclusions

In the current work, we systematically identified lncRNAs in facet joints, discovered differentially expressed lncRNAs in FJOA, and predicted target genes and biological functions of these differentially expressed lncRNAs. Our work provided a preliminary overview of lncRNAs in FJOA and might benefit the further understanding of the mechanisms of FJOA.

Conflicts of interest

There are no conflicts to declare.

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