Long Intergenic Noncoding RNAs Mediate the Human Chondrocyte Inflammatory Response and Are Differentially Expressed in Osteoarthritis Cartilage

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Objective. To identify long noncoding RNAs (lncRNAs), including long intergenic noncoding RNAs (lincRNAs), antisense RNAs, and pseudogenes, associated with the inflammatory response in human primary osteoarthritis (OA) chondrocytes and to explore their expression and function in OA.

Methods. OA cartilage was obtained from patients with hip or knee OA following joint replacement surgery. Non-OA cartilage was obtained from postmortem donors and patients with fracture of the neck of the femur. Primary OA chondrocytes were isolated by collagenase digestion. LncRNA expression analysis was performed by RNA sequencing (RNAseq) and quantitative reverse transcriptase–polymerase chain reaction. Modulation of lncRNA chondrocyte expression was achieved using LNA longRNA GapmeRs (Exiqon). Cytokine production was measured with Luminex.

Results. RNAseq identified 983 lncRNAs in primary human hip OA chondrocytes, 183 of which had not previously been identified. Following interleukin-1β (IL-1β) stimulation, we identified 125 lincRNAs that were differentially expressed. The lincRNA p50-associated cyclooxygenase 2–extragenic RNA (PACER) and 2 novel chondrocyte inflammation–associated lincRNAs (CILinc01 and CILinc02) were differentially expressed in both knee and hip OA cartilage compared to non-OA cartilage. In primary OA chondrocytes, these lincRNAs were rapidly and transiently induced in response to multiple proinflammatory cytokines. Knockdown of CILinc01 and CILinc02 expression in human chondrocytes significantly enhanced the IL-1–stimulated secretion of proinflammatory cytokines.

Conclusion. The inflammatory response in human OA chondrocytes is associated with widespread changes in the profile of lncRNAs, including PACER, CILinc01, and CILinc02. Differential expression of CILinc01 and CILinc02 in hip and knee OA cartilage, and their role in modulating cytokine production during the chondrocyte inflammatory response, suggest that they may play an important role in mediating inflammation-driven cartilage degeneration in OA.

Osteoarthritis (OA), typified by degenerative loss of cartilage integrity and joint space narrowing, is a leading cause of pain, disability, and shortening of adult working life throughout the world (1–3). Unfortunately, at present there is no approved treatment that can modify the disease progression, resulting in limited therapeutic options for patients (4).

In attempting to identify novel therapeutics, inflammation is increasingly being recognized as an important driver of OA cartilage pathology. Histologic analysis, ultrasound, and magnetic resonance imaging have all demonstrated evidence of synovitis in OA joints (5–7), with increased cellular infiltration of activated B cells and T lymphocytes. Indeed, synovitis is reported not only in established OA, but also at the onset of OA, being present in patients with only minimal radiographic signs of the disease (8). Several proinflammatory cytokines are elevated in the synovial fluid of OA joints...
compared to normal healthy joints (9), and cytokine stimulation of ex vivo cartilage tissue mimics the pathologic changes observed within the OA joint (9,10). However, the key regulators of the cellular inflammatory response in cartilage tissue are not well defined.

There is now overwhelming evidence that the microRNA (miRNA) family of short noncoding RNAs can regulate the inflammatory response (11,12). Indeed, our group previously identified differentially expressed miRNAs in human OA cartilage tissue that mediated the production of matrix metalloproteinase 13 (MMP-13) and tumor necrosis factor (TNF) (13). Of interest, earlier reports suggest that these miRNAs may also be central regulators of biologic processes (16–19), including the inflammatory response (20). In support of those findings, we recently identified miRNAs that were differentially expressed upon lipo-poly saccharide (LPS)-induced activation of the human innate response and demonstrated that these regulated interleukin-1β (IL-1β) and IL-8 production (21).

Currently, little is known about the expression and functional role of IncRNAs in OA joint tissue. Their potential importance is indicated in a recent report by Fu et al (22), who identified ~4,700 IncRNAs that were differentially expressed in cartilage from patients with knee OA (compared with controls) using a microarray-based approach. Although that preliminary study did not examine the function of these IncRNAs, another recent study has identified a lincRNA located upstream of the gene PTGS2 (cyclooxygenase 2 [COX-2]). This was shown to be increased in phorbol myristate acetate– and LPS-stimulated monocytes and to positively regulate COX-2 expression (23) by binding to, and relieving the action of, the repressive p50 component of the NF-κB complex (23). As a result of this action, the lincRNA was renamed p50-associated COX-2–extragenic RNA (PACER). Importantly, COX-2 is a key regulator of the arachidonic acid pathway and subsequent prostaglandin E2 production (24), which is a putative mediator of inflammation and pain in OA cartilage tissue (25,26). Given these observations, and the key role of inflammation in OA cartilage pathology, we hypothesized that IncRNAs, including PACER, are central regulators of the inflammatory response in cartilage tissue.

The aim of this study was therefore to perform RNAseq in order to identify IncRNAs that are associated with the inflammatory response in primary human OA chondrocytes isolated from the articular cartilage of patients with hip OA. We then proceeded to assess their potential involvement in OA by examining the expression of several “inflammation-associated” IncRNAs (including PACER) in human articular cartilage from patients with and those without hip or knee OA, profiling their expression in response to multiple proinflammatory cytokines and determining the functional effect of modulating the expression of an inflammation-associated lncRNA on the chondrocyte inflammatory response.

PATIENTS AND METHODS

Patients and tissue samples. Following ethics approval (UK National Research Ethics Committee 14/ES/1044), patients with hip OA (mean ± SEM age 69 ± 3 years; n = 9), patients with knee OA (age 70 ± 3 years; n = 12), and patients with fracture of the neck of the femur without OA (age 74 ± 2 years; n = 6) were recruited prior to elective joint replacement surgery at either The Royal Orthopaedic Hospital (Birmingham, UK) or Russell’s Hall Hospital (Dudley, UK). Patients with hip OA had Kellgren/Lawrence (K/L) grades of 3 or 4, patients with knee OA all had K/L grades of 4, and patients with fracture of the neck of the femur had K/L grades of 0. Cartilage from femoral condyles (from knee OA patients) and femoral heads (from hip OA patients) was collected. Ethics approval was also obtained (Derby Research Ethics Committee 1 [11/H0405/2]) to collect non-OA knee cartilage from postmortem donors (mean ± SEM age 74 ± 5 years; n = 4) (Kings Mill Hospital, Sutton-in-Ashfield, UK) with no history of joint pain or evidence of cartilage fibrillation based on chondropathy assessment (28). Consent was obtained from all patients or families. Patient demographic data are provided in Supplementary Table 1, available on the Arthritis & Rheumatology web site at http://onlinelibrary.wiley.com/doi/10.1002/art.39520/abstract. A protocol was in place to ensure that samples were all handled in the same way and processed in the same timeframe. For tissue processing, upon separation of cartilage from bone tissue, the cartilage was immediately snap-frozen in liquid nitrogen.

Isolation of primary chondrocytes from articular cartilage. Articular cartilage was separated from the subchondral bone using a scalpel and digested using filter-sterilized collagenase IIA (2 mg/ml; Sigma-Aldrich) for 5 hours at 37°C. Digested cartilage was then filtered by passing through a 40-μm cell strainer (BD Biosciences), and the filtrate was centrifuged. Primary chondrocytes were then resuspended in growth media (Dulbecco’s modified Eagle’s medium supplemented with 10% fetal calf serum [FCS], penicillin [100 units/ml], streptomycin [100 μg/ml], l-glutamine [2 mM], nonessential amino acids [5% volume/volume] [all from Life Technologies], and amphotericin [2 μg/ml; Sigma-Aldrich]) and grown to 70–80% confluence before being used in subsequent studies.

RNAseq analysis. Primary hip OA chondrocytes (n = 3 patients) were left unstimulated or stimulated with IL-1β (1 ng/ml) for 4 hours in 0.1% FCS culture media in the absence of antibiotics and amphotericin. Total RNA was isolated using TRIzol reagent (Life Technologies), further purified (RNaseasy column; Qiagen), and the RNA integrity number (RIN) was assessed (Agilent Bioanalyzer). All RIN
values were >7, and 260:280 ratios (measured by NanoDrop) were >1.7. Ribosomal RNA was removed using Ribozero (Epicentre Technologies), and RNA-seq (100-bp paired-end, stranded sequencing) was performed on an Illumina HiSeq 2000 sequencer. Subsequent analysis was undertaken using Tophat2/Cufflinks with alignment against the hg19 reference genome (Figure 1A). LncRNAs were identified using Cufflinks and then compared with known lncRNAs previously annotated in Gencode version 19 and the Human LincRNAs Catalog (29). CuffDiff was used to compare control and IL-1β–treated cells to identify differentially expressed transcripts (false discovery rate [FDR], 0.05, fold change >2, and change in fragments per kilobase of transcript per million mapped reads [FPKM] >1). Sequence data are available through the GEO database under series number GSE74220.

Analysis of lncRNA expression in primary chondrocytes and cartilage by quantitative reverse transcriptase–polymerase chain reaction (qRT-PCR). Articular hip and knee cartilage was snap-frozen in liquid nitrogen and pulverized using a 6770 Freezer/mill (Spex Sample Prep). Total RNA was extracted from both powdered cartilage and primary chondrocytes using TRIzol and further purified using RNeasy columns. RIN values were >7, and 260:280 ratios were >1.7. Custom primers and FAM-labeled probes were designed using Primer Express 3 software (Life Technologies) for qRT-PCR. The qRT-PCR was performed from 25 ng of total RNA in a one-step reaction (QuantiFast One-Step RT-PCR kit; Qiagen) using a Roche LightCycler 480 II. The relative expression of lncRNAs was determined using the ΔΔCt method, following normalization to 18S RNA. GAPDH expression (relative to 18S RNA) was comparable between non-OA and OA cartilage in both hip and knee samples (see Supplementary Figure 1, available on the Arthritis & Rheumatology web site at http://onlinelibrary.wiley.com/doi/10.1002/art.39520/abstract).

Inhibition of lncRNA expression in human chondrocytes using locked nucleic acid (LNA) GapmeRs. The human chondrocyte cell line TC28, which was previously characterized by Goldring et al (30), and provided to us as a gift from AstraZeneca, was transfected with either LNAs targeting CILinc01 or CILinc02 (30 nM) or with LNA control (30 nM) using LipoIce Transfectamine 2000 (Life Technologies). Following 24-hour transfection, cells were stimulated (in 0.1% FCS culture media in the absence of antibiotics and amphotericin) for either 4 hours or 24 hours with IL-1β (1 ng/ml). Supernatants were collected for subsequent cytokine analysis with Luminex. Cells were lysed with RLT (Qiagen) for subsequent RNA extraction to examine knockdown of CILinc01 expression.

Analysis of cytokine production in human chondrocyte supernatants. Supernatants from human chondrocyte–transfected cells and media controls were assayed for the concentration of 17 human proinflammatory cytokines using a human cytokine 17-plex immunoassay (Bio-Plex Pro; Bio-Rad). The interassay variability is <15%; intrasay variability is <10%. Cross-reactivity is <1%, and the dynamic range is

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**Figure 1.** Regulation of long noncoding RNA (lncRNA) expression by interleukin-1β (IL-1β) in human osteoarthritis (OA) chondrocytes. A, Pipeline for predicting lncRNAs from Cufflinks-assembled transfrag. FPKM = fragments per kilobase of transcript per million mapped reads. B, Release of IL-6 from primary human hip OA chondrocytes left unstimulated or stimulated with IL-1β for 4 hours or 24 hours, as measured by enzyme-linked immunosorbent assay. IL-6 release indicates activation of the inflammatory response. Bars show the mean ± SEM. *** = P < 0.001. C, Volcano plot displaying differentially expressed mRNAs (n = 3 IL-1β-stimulated hip OA chondrocytes and 3 unstimulated hip OA chondrocytes.). D, Pathway analysis of differentially expressed mRNAs. FDR = false discovery rate. E, Overlap of lncRNAs in OA chondrocytes, Gencode version 19, and the HumanBodyMap catalogs. F, Breakdown of differentially expressed lncRNAs based on positional classifications.
between 1 and 2,500 pg/ml. Briefly, nondiluted chondrocyte cell culture supernatants were incubated with a magnetic Bio-Plex bead cocktail consisting of beads specific for IL-1β, IL-2, IL-4, IL-5, IL-6, IL-7, IL-8, IL-10, IL-12 (p70), IL-13, IL-17, granulocyte colony-stimulating factor (G-CSF), granulocyte–macrophage colony-stimulating factor, interferon-γ, monocyte chemotactic protein 1, macrophage inflammatory protein 1β (MIP-1β), and TNF. A Bio-Plex Pro Wash Station was used to wash the beads between incubation steps using the wash buffer supplied with the kit. A biotinylated secondary antibody was added, and quantification was carried out using a streptavidin–phycoerythrin substrate with fluorescence detected on a Bio-Plex 200 System (Bio-Rad/Luminex).

Statistical analysis. Data were analyzed using SPSS software. Analysis of variance was performed throughout, followed by Fisher’s least significant difference post hoc test, where appropriate. In all cases, data are presented as the mean ± SEM, and P values less than 0.05 were considered significant.

RESULTS

RNAseq transcriptome profile of primary human OA chondrocytes in response to stimulation with IL-1β. IL-1β stimulation of primary human hip OA chondrocytes (n = 3 patients) induced a rapid release of IL-6 protein that peaked at 4 hours and remained elevated at 24 hours (Figure 1B). IL-1β stimulation also induced a significant increase in the release of MMP-13 at 24 hours (see Supplementary Figure 2, available on the Arthritis & Rheumatology web site at http://onlinelibrary.wiley.com/doi/10.1002/art.39520/abstract). Analysis of RNAseq data for Gencode-annotated messenger RNAs (mRNAs) showed that 499 protein-coding genes were differentially expressed upon IL-1β stimulation (382 up-regulated and 117 down-regulated) (Figure 2C).
1C and Supplementary Table 2, available on the Arthritis & Rheumatology web site at http://onlinelibrary.wiley.com/doi/10.1002/art.39520/abstract). As expected, the up-regulated genes from this set were significantly enriched (FDR, 0.05) in Kyoto Encyclopedia of Genes and Genomes pathways involved in the inflammatory response (Figure 1D). There were no significantly enriched pathways in down-regulated genes. This initial evaluation therefore demonstrated rapid and wide-spread induction of inflammatory gene expression following IL-1β stimulation of human chondrocytes.

Identification of novel lncRNAs in chondrocytes by RNAseq. Using the computational analysis pathway described in Figure 1A, we identified 983 lncRNAs in human chondrocytes, which could be divided into 642 lincRNAs, 124 antisense RNAs, and 217 pseudogenes (see Supplementary Table 3, available on the Arthritis & Rheumatology web site at http://onlinelibrary.wiley.com/doi/10.1002/art.39520/abstract). Of these assembled genes, 158 lincRNAs and 25 antisense RNAs had not previously been identified in Gencode version 19 or HumanBodyMap lncRNA (Figures 1E and F). As previously reported (14,15), the mean FPKM, length, and exon number for lncRNAs were smaller than those for mRNAs (mean FPKM 4.7 for lncRNAs and 29.6 for mRNAs, mean length 1.2 kb for lncRNAs and 2.8 kb for mRNAs, and mean exon number 3.6 for lncRNAs and 16.4 for mRNAs).

Based on sequencing in ~400 human cell types including chondrocytes, the FANTOM project has recently released an atlas of 43,011 enhancer regions that are characterized by bidirectional transcription of single-exon efference RNAs (eRNAs) (31). Interestingly, we found that <4% of our identified lncRNAs overlapped with putative eRNA regions (see Supplementary Table 3, available on the Arthritis & Rheumatology web site at http://onlinelibrary.wiley.com/doi/10.1002/art.39520/abstract). Furthermore, visual inspection and the fact that our transcripts were unidirectional and multieXonic indicated that these lncRNAs did not represent eRNAs.

Induction of widespread changes in lncRNA expression by IL-1β stimulation. Following IL-1β stimulation, we identified 125 lncRNAs that were differentially expressed (P, 0.05), including 93 lincRNAs (74%), 13 antisense RNAs (11%), and 19 pseudogenes (15%) (see Supplementary Table 4, available on the Arthritis & Rheumatology web site at http://onlinelibrary.wiley.com/doi/10.1002/art.39520/abstract). Of these, we observed 106 up-regulated and 19 down-regulated lncRNAs, of which 37 (30%) were novel lncRNAs. Using the Integrative Genomics Viewer (Broad Institute), the transcription start sites (TSS) for the majority of the 92 differentially expressed lncRNAs were found to be genomically located >5 kb from the TSS of a coding mRNA (Figures 2A and B). Previously, we have referred to these as mRNA-flanking lncRNAs (21), and it has been suggested that they may regulate the expression of the nearby mRNA. In support of this notion, we found a significant positive correlation between the fold change in expression of an mRNA-flanking lncRNA and the fold change in expression of its nearest coding mRNA (Figure 2C). In addition, detailed examination of these differentially expressed mRNA-flanking lncRNAs identified one as being PACER (Table 1). As previously described, PACER is located upstream of the PTGS2 (COX-2) gene, is transcribed in a bidirectional manner from the same promoter region, and is known to positively regulate PTGS2 expression (23) (Figure 2A). However, whether

<table>
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<th>LincRNA</th>
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* LincRNA = long intergenic noncoding RNA; OA = osteoarthritis; lncRNA = long noncoding RNA; TSS = transcription start site; IL-1β = interleukin-1β; PACER = p50-associated COX-2–extragenic RNA.
† = P < 0.001 versus unstimulated control chondrocytes.
‡ = P < 0.01 versus unstimulated control chondrocytes.
this is true of other mRNA-flanking lincRNAs remains to be elucidated.

**Differential expression of inflammation-associated lincRNAs in human hip OA and knee OA cartilage.** We next wished to further characterize the expression of PACER as well as 7 additional chondrocyte inflammation–associated lincRNAs (named CILinc01–CILinc07) that were selected based on being significantly induced in response to IL-1β stimulation (Table 1) and their nearest coding mRNA being a gene with purported evidence of a role in either inflammation or OA pathology (e.g., IL-7 and ADAMTS-5, respectively).

We initially determined the potential clinical relevance of these chondrocyte inflammation–associated lincRNAs by measuring their expression in human OA hip cartilage compared to non-OA hip cartilage. All 8 lincRNAs were found to be significantly down-regulated in OA hip cartilage (n = 9 patients) compared to non-OA hip cartilage (n = 6 patients) (Figure 3A). The lincRNAs PACER, CILinc01, and CILinc02 were also significantly down-regulated (>2-fold) in OA knee cartilage (n = 12) compared to non-OA knee cartilage (n = 4) (Figure 3B).

**Rapid, transient induction of lincRNAs by multiple proinflammatory cytokines.** Based on their induction in response to IL-1β stimulation, and their differential expression in both hip OA and knee OA cartilage, we next examined the time course of expression of PACER, CILinc01, and CILinc02 in primary OA chondrocytes in response to a panel of proinflammatory cytokines implicated in the pathogenesis of OA. Following stimulation with IL-1β, TNF, visfatin, and leptin, we observed a rapid and time-dependent induction of expression of all 3 lincRNAs (Figure 4A).
lincRNA expression at ~2 hours, which had dropped toward baseline levels by 24 hours. Stimulation with IL-1β or visfatin led to a slightly more prolonged induction of lincRNA expression, with peak induction of CILinc01 and CILinc02 between 4 and 6 hours (Figure 4A). Of note, stimulation with IL-1β for 4 hours also led to significant \( P < 0.001 \) induction of the expression of mRNA for the closest coding genes to PACER, CILinc01, and CILinc02, namely, PTGS2, HIVEP2, and IL-7, respectively (Figure 4B).

We then assessed whether PACER, CILinc01, and CILinc02 were also present in non-OA chondrocytes and whether stimulation of these cells with IL-1β would also induce their expression. PACER, CILinc01, and CILinc02 were expressed in both non-OA knee and non-OA hip chondrocytes (isolated from cartilage from patients with fracture of the neck of the femur (#NOF) and from postmortem (PM) cartilage and in primary OA chondrocytes isolated from hip and knee cartilage. Expression of lincRNAs and genes was determined by quantitative reverse transcriptase–polymerase chain reaction and is shown as fold change compared to control. Bars show the mean ± SEM from 3 independent experiments. \( * = P < 0.05; † = P < 0.01; ‡ = P < 0.001 \) versus unstimulated samples, by two-way analysis of variance with a least significant difference post hoc test.
led to a significant increase in expression of each of the 3 lincRNAs (Figure 4C).

**Negative regulation of the IL-1β-stimulated production of proinflammatory cytokines by CILinc01 and CILinc02.** Given the association of CILinc01 and CILinc02 with the IL-1β chondrocyte inflammatory response, and their down-regulation in OA cartilage tissue, we speculated that CILinc01 and CILinc02 might mediate the production of proinflammatory cytokines. To test this hypothesis, we examined the effect of knockdown of CILinc01 and CILinc02 expression on the human chondrocyte inflammatory response. For these experiments, we used the human chondrocyte TC28 cell line, which when incubated in low serum (0.1% FCS) without stimulation expressed type II collagen (see Supplementary Figures 3A and B, available on the Arthritis & Rheumatology website at http://onlinelibrary.wiley.com/doi/10.1002/art.39520/...
abstract). Similar to the findings in primary chondrocytes, IL-1β stimulation of TC28 cells induced a rapid release of IL-6 protein (Supplementary Figure 3C) and induction of MMPs and proinflammatory cytokines (Supplementary Figure 3D). TC28 cells were transfected with either LNA GapmeRs targeting CILinc01 or CILinc02, or a non-targeting control LNA GapmeR. Following 24 hours of transfection, cells were stimulated with IL-1β for 4 hours in order to provoke an inflammatory response. Similar to our findings in primary human chondrocytes, 4 hours of exposure of the TC28 chondrocyte cell line to IL-1β led to a significant induction of expression of CILinc01 and CILinc02. The IL-1β–induced expression of CILinc01 was significantly reduced (by 63%) in chondrocytes transfected with an anti-CILinc01 LNA GapmeR, and CILinc02 expression was significantly reduced (by 74%) in cells transfected with an anti-CILinc02 GapmeR, compared to an LNA control sequence (Figure 5A).

We then investigated the effect of CILinc01 and CILinc02 knockdown on the inflammatory response, by measuring the secretion of a panel of 17 proinflammatory cytokines in response to 4 hours of IL-1β stimulation of human chondrocytes (see Supplementary Table 5, available on the Arthritis & Rheumatology web site at http://onlinelibrary.wiley.com/doi/10.1002/art.39520/abstract). Knockdown of CILinc01 expression significantly enhanced the IL-1β–stimulated production of IL-6, IL-8, TNF, MIP-1β, and G-CSF (Figure 5B), while knockdown of CILinc02 expression significantly reduced the production of IL-6 (Figure 5C). Since previous studies have shown that NF-κB activity can regulate the expression of lncRNAs, we then also examined the effect of pharmacologic inhibition of IKK-2 on the IL-1β–stimulated production of CILinc01 and CILinc02. To this end we used TPCA-1, a known IKK-2 inhibitor (20,32,33). In cell-free enzymatic assays, TPCA-1 displays 22-fold selectivity for IKK-2 over IKK-1 and a >550-fold selectivity over other kinases, including MAP kinases and JNK kinases (32), though a recent study showed that in non–small cell lung cancer cell lines TPCA-1 also inhibited STAT-3 phosphorylation (34). Preincubation of primary chondrocytes with TPCA-1 (10 μM) significantly reduced the induction of both CILinc01 and CILinc02 that occurred after 4 hours of stimulation with IL-1β (Figure 5D).

**DISCUSSION**

This study is the first to use RNAseq to determine the profile of lncRNA expression in primary human OA chondrocytes and has resulted in the cataloging of 983 lncRNAs, including members of the lncRNA, antisense RNA, and pseudogene families. Importantly, we have identified 158 lncRNAs and 25 antisense RNAs that are absent from Gencode version 19 (35) and the HumanBodyMap lncRNA catalog (29), and might therefore be unique to chondrocytes and have a cell-specific function. In addition, this study is the first to examine the changes in lncRNA levels that are associated with the inflammatory response in human chondrocytes. In this regard, 125 lncRNAs were differentially expressed upon IL-1β stimulation of human OA chondrocytes. Of relevance, Fu et al (22) recently showed a catalog of 4,714 lncRNAs found by microarray analysis to be differentially expressed in knee OA patients compared to non-OA cartilage. In our RNAseq chondrocyte analysis, if we included lncRNAs with a P value of less than 0.05 (rather than an FDR optimized q of <0.05), which was the inclusion criterion used by Fu et al (22), 7 of these lncRNAs (namely, ENST00000419463, ENST00000369884, ENST00000419463, ENST00000421237, ENST00000412485, ENST00000455607, and ENST00000418242) were differentially expressed in chondrocytes upon IL-1β stimulation. This relatively low number of lncRNAs in common is likely due to differences in conditions (IL-1β stimulation of chondrocytes versus end-stage cartilage disease comparison) and methodologic approach (sequencing versus microarrays). As an example, the microarray studies by Fu et al (22) would not have detected changes in PACER, CILinc01, and CILinc02 since these are novel transcripts for which there are no microarray probes. Despite these differences, we speculate that these shared lncRNAs might have a function in OA, which would warrant further investigation.

Importantly, there is now evidence that lncRNAs regulate in *cis* local mRNA expression (21,36). Indeed, among those lncRNAs differentially expressed upon IL-1β stimulation was the lncRNA PACER (23), which is located adjacent to and upstream of the gene PTGS2 (COX-2) and has been shown to regulate PTGS2 production (23). As shown in the present study, PACER appears to be transcribed from the same promoter regions as PTGS2, which results in bidirectional production of both coding and noncoding RNA. Significantly, the majority of the inflammation-associated lncRNAs we identified were found to be mRNA flanking, several of which (including PACER) were located close to genes relevant to either inflammation or cartilage biology, which could be indicative of a functional role in OA.

Given these observations, we selected PACER and 7 additional inflammation-associated lncRNAs and proceeded to investigate their potential clinical relevance by comparing their expression in articular hip and
The functional significance of our finding that CILinc02 is down-regulated in human hip OA cartilage is unclear. Studies in rheumatoid arthritis suggest that IL-7 (the nearest coding gene to CILinc02) contributes to inflammation (43) and mediates the production of TNF (44), while in OA, IL-7 has been reported to induce MMP-13 and proteoglycan loss from cartilage, suggesting that it may promote cartilage degeneration (45). However, we did not detect expression of the IL-7 gene in either OA or non-OA hip cartilage samples.

Functional studies to determine the roles of CILinc01 and CILinc02 showed that knockdown of their expression in human chondrocytes significantly increased the IL-1β-stimulated production of several proinflammatory cytokines, including IL-6, suggesting that CILinc01 and CILinc02 may negatively regulate the chondrocyte inflammatory response. It is significant, therefore, that we found decreased expression of CILinc01 and CILinc02 in both knee OA and hip OA cartilage compared to normal healthy cartilage, since this could indicate that down-regulation of CILinc01 and CILinc02 in human articular cartilage leads to an inability to regulate inflammation in the joint. Of interest, the nearest coding gene to CILinc01 is HIVEP2 (also known as Schnurri-2), which has previously been reported to be a negative regulator of allergic airway inflammation via repression of NF-κB activity (46), as well as being implicated in mediating chondrocyte differentiation (47). Therefore, it is conceivable that the observed effects of CILinc01 on chondrocyte cytokine production are mediated via repression of NF-κB activity through modulation of HIVEP2 gene expression. Of note, stimulation of primary chondrocytes with an IKK-2 inhibitor blocked the IL-1β-stimulated production of both CILinc01 and CILinc02, suggesting that NF-κB activity may regulate their expression in chondrocytes.

In conclusion, these data signify that CILinc01 and CILinc02 may play an important physiologic role in regulating the pathologic response to inflammation within the OA joint, and that its down-regulation in both knee and hip OA cartilage could contribute to inflammation-driven cartilage degeneration. Clearly, future studies to determine the mode of action of CILinc01 and CILinc02 as well as other chondrocyte inflammation–associated lincRNAs in mediating OA cartilage pathology and inflammation are warranted and may lead to the identification of novel targets for the development of therapeutic agents.

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REFERENCES


