MicroRNA-143 and -145 modulate the phenotype of synovial fibroblasts in rheumatoid arthritis

Bong-Ki Hong, Sungyong You, Seung-Ah Yoo, Dohyun Park, Dahee Hwang, Chul-Soo Cho and Wan-Uk Kim

Fibroblast-like synoviocytes (FLSs) constitute a major cell subset of rheumatoid arthritis (RA) synovia. Dysregulation of microRNAs (miRNAs) has been implicated in activation and proliferation of RA-FLSs. However, the functional association of various miRNAs with their targets that are characteristic of the RA-FLS phenotype has not been globally elucidated. In this study, we performed microarray analyses of miRNAs and mRNAs in RA-FLSs and osteoarthritis FLSs (OA-FLSs), simultaneously, to validate how dysregulated miRNAs may be associated with the RA-FLS phenotype. Global miRNA profiling revealed that miR-143 and miR-145 were differentially upregulated in RA-FLSs compared to OA-FLSs. miR-143 and miR-145 were highly expressed in independent RA-FLSs. The miRNA-target prediction and network model of the predicted targets identified insulin-like growth factor binding protein 5 (IGFBP5) and semaphorin 3A (SEMA3A) as potential target genes downregulated by miR-143 and miR-145, respectively. IGFBP5 level was inversely correlated with miR-143 expression, and its deficiency rendered RA-FLSs more sensitive to TNFα stimulation, promoting IL-6 production and NF-κB activity. Moreover, SEMA3A was a direct target of miR-145, as determined by a luciferase reporter assay, antagonizing VEGF165-induced increases in the survival, migration and invasion of RA-FLSs. Taken together, our data suggest that enhanced expression of miR-143 and miR-145 renders RA-FLSs susceptible to TNFα and VEGF165 stimuli by downregulating IGFBP5 and SEMA3A, respectively, and that these miRNAs could be therapeutic targets.

INTRODUCTION

Rheumatoid arthritis (RA) is an autoimmune disease that is characterized by chronic inflammatory cell infiltration and pannus formation in synovial tissues that lead to the destruction of articular cartilage and bone. In RA joints, various inflammatory cells become activated via a variety of cytokines and chemokines as well as via cell–cell contact. Among various cell types, fibroblast-like synoviocytes (FLSs) are the most abundant resident cells in the synovial membrane and play critical roles in the pathogenesis of RA. FLSs in an RA patient (RA-FLSs) have tumor-like features, including an increased proliferation rate, anti-apoptotic ability and pro-migratory and pro-invasive properties that cause pannus formation and joint destruction. Moreover, RA-FLSs produce large amounts of pro-inflammatory cytokines, including interleukin-1β (IL-1β), IL-6 and tumor necrosis factor α (TNFα), that contribute to the perpetuation of chronic inflammation.

MicroRNAs (miRNAs) have emerged as key regulators of a broad spectrum of cellular functions, such as proliferation, differentiation and apoptosis, that are associated with the pathogenesis of various autoimmune diseases. Evidence is emerging that miRNA expression is dysregulated in RA-FLSs, which may be responsible for various pathologic processes involving RA. For example, miR-126 has been identified as a regulator of phosphatidylinositol 3-kinase (PI3K)/AKT in RA-FLSs, an important signaling molecule mediating cell proliferation and apoptosis. In addition, miR-20a has been found to have two target genes in FLSs: thioredoxin interacting protein (TXNIP) and apoptosis signal-regulating kinase (ASK1). These mediate NLRP3-inflammasome and TLR4-dependent cytokine release by FLSs, respectively. miR-221 has also been identified as a regulator for increased migration and invasion of RA-FLSs. Nevertheless, global and integrated analyses of miRNA and mRNA expression levels, which can...
address pathologic features of RA-FLSs, have not been systematically performed.

In the present study, we simultaneously profiled the global expression of miRNAs and mRNAs in FLSs through microarray analysis and found previously unidentified miR-143 and miR-145 to be differentially upregulated in RA-FLSs compared to FLSs from osteoarthritis patients (OA-FLSs). We validated that miR-143 and miR-145 were highly expressed in independent RA-FLSs. Moreover, miR-143 expression in RA-FLSs was increased by transforming growth factor β (TGFβ). By using the miRNA target prediction and network model of the predicted targets, we selected two potential targets, insulin-like growth factor binding protein 5 (IGFBP5) and semaphorin 3A (SEMA3A), that could be downregulated by miR-143 and miR-145, respectively. In in vitro functional tests, IGFBP5 deficiency induced RA-FLS sensitivity to TNFα stimulation and promoted IL-6 production and NF-κB activity. Moreover, the 3′-UTR of SEMA3A mRNA was shown to be a direct target of miR-145, as determined by a luciferase reporter assay, and recombinant SEMA3A antagonized VEGF165-induced increases in survival, migration and invasion of RA-FLSs. Collectively, our data suggest that enhanced miR-143 and miR-145 expression renders RA-FLSs susceptible to TNFα and VEGF165 stimuli by downregulating IGFBP5 and SEMA3A, respectively. Thus, inhibition of miR-143 and miR-145 could lead to a reduction in RA-FLS pathogenic potential.

MATERIALS AND METHODS

Isolation and culture of FLSs

FLSs were isolated from the synovial tissues of patients with RA or OA and cultured in Dulbecco’s modified Eagle medium (DMEM, 12800-017, Gibco, ThermoFisher Scientific, Grand Island, NY, USA) supplemented with 10% fetal bovine serum (FBS), 6.0 g l⁻¹ of HEPES (0511-1KG, Amresco, Solon, OH, USA), 0.4 g l⁻¹ of l-glutamine (0374-500G, Amresco), 3.7 g l⁻¹ of sodium bicarbonate (0865-1KG, Amresco), 1 × of antibiotics (15240-062, Gibco, ThermoFisher Scientific), 1 × of MEM non-essential amino acids (11140-050, Gibco), and 1 × of antibiotics (12800-017, Gibco, ThermoFisher Scientific). Experiments were performed under 5% CO₂ at 37 °C.

Microarray analysis for miRNA and mRNA profiling

We prepared total RNA independently from two replicate samples for miRNA and three replicate samples for mRNA using miRNeasy (217004, Qiagen GmbH, Hilden, Germany) and RNeasy (74104, Qiagen) mini kits, respectively. The RNA integrity was assessed using a Bioanalyzer 2100 (Agilent Technologies, Santa Clara, CA, USA). The RNA integrity number (RIN) was close to 10 for all samples. RNA was then amplified and hybridized with an Agilent-019118 Human miRNA Microarray 2.0 G4470B (for miRNA) and an Illumina HumanHT-12 v4.0 Expression BeadChip (Illumina, San Diego, CA, USA) for mRNA, according to the manufacturer’s standard protocols. The probe intensities were normalized using the quantile normalization procedure. The data were deposited into the GEO database (GEO accession ID, miRNA: GSE91026; mRNA: GSE49604).

Integrative statistical testing

For the four comparisons of RA-FLSs versus OA-FLSs, RA-FLSs+IL1β versus RA-FLSs, OA-FLSs+IL1β versus OA-FLSs, and RA-FLSs+IL1β versus OA-FLSs+IL1β, we used the integrated statistical testing method previously reported. Briefly, for each gene, an adjusted P-value was computed by performing a two-tailed t-test and log2 median ratio test using the empirical distributions that were estimated by random permutations of the samples. The two sets of P-values from the individual tests were combined to compute the overall P-values using Stouffer’s method. Finally, differentially expressed genes (DEGs) were selected that have (1) P<0.05 and (2) absolute log2-fold changes larger than 1.34. To determine the cutoff value for log2-fold changes, we computed log2-fold changes of randomly permuted samples, fitted a Gaussian distribution to the random log2-fold changes, and then calculated the 2.5th percentile corresponding to α = 0.05 (log2-fold-change cutoff = 1.34). The DEGs were grouped into six clusters using hierarchical clustering. The cellular processes represented by the genes in each cluster were identified as the GO biological processes (GOBPs) having P<0.1 (the default cutoff value in the DAVID software).

Hierarchical clustering of DEGs

From the comparisons of RA-FLSs versus OA-FLSs and RA-FLSs+IL1β versus OA-FLSs+IL1β, we identified a total of 470 DEGs. We then categorized the DEGs into two groups of upregulated and downregulated DEGs. For the DEGs in each group, we further performed hierarchical clustering using Euclidean distance as the dissimilarity measure and the average linkage method for clusters for upregulated and downregulated DEGs in all compared groups.

Prediction of target genes

We refined lists of predicted miR-143 and miR-145 target genes through comparisons between the DEG list and network analysis. In addition, we referred to the context score from the TargetScan 6.0 database. A Venn diagram was drawn by Bioinformatics & Evolutionary genomics (http://bioinformatics.psb.ugent.be/webtools/Venn/).

Reconstruction of miR-143 and miR-145 target gene network

To reconstruct a network model for the 1081 genes (404 genes of miR-143 targets, 731 genes of miR-145 and 54 shared genes) that were the predicted targets of miR-143 and/or miR-145, we first collected all protein–protein interactions (PPIs) stored in HPRD (http://www.hprd.org/), BioGRID (https://thebiogrid.org/), STRING (http://string-db.org/) and KEGG (http://www.genome.jp/kegg/), and then combined all PPIs from the four databases into one list of PPIs. We then used Cytoscape version 3.2.0 software (http://www.cytoscape.org/) to display these PPIs for the 1081 genes as a network model.

Transfection of miRNA inhibitors and short interfering RNAs

FLSs were seeded at 1×10⁶ cells per well in a six-well cell culture plate (Nunc, Thermo Scientific, Rockford, IL, USA). After 24 h, the cells were transfected with 50 nM of miScript miRNA inhibitors (Negative control: 1027271, ina-miR-145; MIN0000437, Qiagen) or 10 nM of short interfering RNAs (siRNA) for IGFBP5 (sc-39591, Santa Cruz Biotechnology, Santa Cruz, CA, USA) with Lipofectamine 2000 (11668-027, Invitrogen, Carlsbad, CA, USA). Transfection efficiency was validated by quantitative real-time polymerase chain reaction (PCR).
Quantitative real-time PCR analysis
Total RNA samples in FLs were prepared with TRIzol Reagent (15596018, Ambion, Austin, TX, USA) according to the manufacturer’s instruction. Extracted RNA was stored at −80 °C before use. For miRNA isolation from synovial fluids, an miRNeasy Serum/Plasma Kit (217184, Qiagen) was used. To determine the miRNA and mRNA levels, 1 μg of RNA was synthesized into cDNA using the miScript II RT Kit (218161, Qiagen). The miRNA levels were assessed by real-time PCR using miScript SYBR Green PCR Kit (218073, Qiagen) with CFX96 (Bio-Rad, Hercules, CA, USA). The mRNA levels were assessed by real-time PCR using iT SYBR Green Supermix (170–8880, Bio-Rad). The gene-specific primers used are listed in Supplementary Table 1.

MTT assay
The viability of FLs was assessed by the MTT (3-[4,5-dimethylthiazol-2-yl]-2, 5 diphenyl tetrazolium bromide) assay, as described previously.10

Apoptosis assay
FLS apoptosis was determined using the APOPercentage apoptosis assay kit (Bicolor, Newtownabbey, UK). Digital images of APOPercentage dye-labeled cells, which appear bright pink against the white background under a light microscope, were used to quantify the apoptotic cell numbers.

Invasion and migration assay
For wound migration assays, FLs were seeded at 1.5 × 10^5 cells per well in a six-well plate. After 24 h, the cells were scratched with a pipette tip and then incubated in 1% FBS DMEM with 200 ng ml^-1 of recombinant human VEGF165 (293-VE, R&D Systems, Minneapolis, MN, USA) in the presence or absence of recombinant human SEMA3A (1250-S3, R&D Systems). After 12 h of incubation, FL migration was quantified by counting the cell numbers that had moved beyond a reference line.

An FL invasion assay was performed in BioCoat Matrigel Invasion Chambers (354480, Thermo Science). FLs were seeded at 1 × 10^5 cells per well in a six-well plate with 500 μl of 0.1% FBS DMEM and 200 ng ml^-1 of recombinant human VEGF165 in the presence or absence of recombinant SEMA3A (100 ng ml^-1). After 12 h, the non-invading cells were removed by scrubbing with a cotton-tipped swab, and the cells on the lower surface of the membrane were stained with Diff-Quik stain (Baxter Diagnostics, Deerfield, IL, USA). For quantification, the cells were manually counted in eight random fields. After staining, the images were captured by a slide scanner (Pannoramic MIDI, 3DHISTECH Ltd, Budapest, Hungary).

Enzyme-linked immunosorbent assay for IL-6
Cultured media from FLs were collected and stored at −80 °C. The IL-6 concentrations in the culture supernatants were measured by a Human IL-6 Duoset ELISA kit (dy206, R&D Systems) according to the manufacturer’s instructions.

Immunocytochemistry
For immunocytochemistry staining, RA-FLSs were seeded at 2 × 10^4 cells per well on eight-well glass chamber slides (154534, Nunc, Thermo Scientific). One hour after TNF-α stimulation (0.1 ng ml^-1, 210-TA, R&D Systems), the cells were fixed with chilled fresh methanol and permeabilized with PBS-T containing 0.25% Triton X-100. The slides were then treated with a blocking buffer (1% bovine serum albumin in PBS-T) and incubated overnight with anti-p65 antibody (ab7970, Abcam, Cambridge, UK) at 4 °C. After rinsing with PBS-T, the slides were incubated again with goat anti-rabbit IgG secondary antibody and then with Alexa Fluor 488 conjugate (A-11008, Invitrogen) for 30 min and mounted using mounting medium with DAPI. The mounted slides were examined using a confocal laser scanning microscope (LSM 510 Meta, Zeiss, Oberkochen, Germany).

Western blot analysis of IκBα
RA-FLSs were seeded at 1 × 10^5 cells per well in a six-well plate. After siRNA transfection or treatment with recombinant human IGF1BP5, the cells were stimulated with 0.1 ng ml^-1 of TNFα for the indicated times (0, 1, 5, 10, 15 and 30 min). The cells were then lysed with 100 μl of 1 × sodium dodecyl sulfate sample buffer and incubated at 95 °C for 3 min. Insoluble material was removed by centrifugation at 12,000 g for 15 min at 4 °C. Electrophoresis was carried out using sodium dodecyl sulfate-10% polyacrylamide gels electrophoresis, and the resolved proteins were transferred to nitrocellulose membranes. Membranes were incubated with antibodies to IκBα (sc-371, Santa Cruz Biotechnology). β-Actin was used as an internal control. Membranes were then visualized using an enhanced chemiluminescence technique. The resulting films were scanned, and the optical densities were quantified using ImageQuant LAS 4000 (GE Healthcare, Chicago, IL, USA) and ImageJ software.

Reporter vector construction
The whole SEMA3A 3′-UTR (3 kilobase pairs) was cloned as insert DNA for reporter vector construction using primers containing NotI sites (Supplementary Table 2). The insert DNA and psiCHECK-2 Dual luciferase reporter vector (C8021, Promega, Madison, MA) were digested by NotI (R0189, NEB, Ipswich, MA, USA) and then ligated. To mutate two miR-145 seed regions on the SEMA3A 3′-UTR, a Q5 Site-Directed Mutagenesis Kit (E0554S, NEB) was used. The primers for mutagenesis were designed by NEBaseChanger (http://nebasechanger.neb.com/, Supplementary Table 2). For amplification, the constructed vector was transformed into E. coli (JM109, L2005, Promega) and cultured overnight at 37 °C. The amplified vectors were harvested by the NucleoBond Xtra Midi kit (740410.10, Macherey-Nagel, Dueren, Germany) according to the manufacturer’s protocol and then purified by Spin-X (CLS8160, Sigma-Aldrich, St Louis, MO, USA).

Dual luciferase reporter assay
HEK293T cells were seeded overnight at 5 × 10^4 cells per well in a 24-well plate. The next day, the cells were transfected with 10 ng of constructed vector plus 50 nm of negative control mimic (AllStars Negative Control siRNA, Qiagen, SI03650318) or 50 nm of hsa-miR-145 mimic (Qiagen, MSY0000437) by Lipofectamine 3000 according to the manufacturer’s protocol. The cells were harvested 24 or 48 h after transfection. A Dual-Glo luciferase reporter assay kit (E2920, Promega) was used for luciferase analysis according to the manufacturer’s instruction. Luciferase activity was measured with SpectraMax L (Molecular Devices, Sunnyvale, CA, USA).

Statistical analysis
Data in the in vitro functional experiments are expressed as the mean ± s.d. Comparisons of the numerical data between groups were performed by the t-test or Mann–Whitney U-test. P-values < 0.05 were considered statistically significant.
Study approval
This study was performed with the approval of the institutional review board (CUMC09U034).

RESULTS

miR-143 and miR-145 are highly expressed in RA-FLSs
To determine the differentially expressed microRNAs (DEmiRs) in RA-FLSs, we initially performed a microarray analysis. From the four comparisons, including RA-FLSs versus OA-FLSs, RA-FLSs with IL-1β versus RA-FLSs, OA-FLSs with IL-1β versus OA-FLSs and RA-FLSs with IL-1β versus OA-FLSs with IL-1β, 16 DEmiRs were identified with significant up- or downregulation in one of those comparisons (Figure 1a). Of note, miR-143 and miR-145, out of 16 total DEmiRs, exhibited significantly higher expression in RA-FLSs compared to OA-FLSs regardless of IL-1β treatment (Figure 1a). We confirmed the higher expression levels of miR-143 and miR-145 in independent RA-FLSs by real-time PCR (Figure 1b). Notably, we found a clear positive correlation of the expression levels between miR-143 and miR-145 (Figure 1c). Given that genes encoding miR-143 and miR-145 are located in the same cluster in chromosome 5,13 this result suggests that miR-143 and miR-145 could be co-regulated in RA-FLSs. As miRNAs can be present in human biological fluids, including plasma, semen and synovial fluids,14–16 we investigated the levels of miR-143 and miR-145 in synovial fluids obtained from RA patients (RA-SF, n=4) and OA patients (OA-SF, n=5) as determined by real-time PCR. (d) Increase in miR-145 expression in FLSs by TGFβ stimulation. The OA-FLSs (n=4) and RA-FLSs (n=4) were treated with IL-1β (1 ng ml−1), TNFα (10 ng ml−1) and TGFβ (10 ng ml−1) for 48 h, and miR-145 expression was then determined by real-time PCR. The data show the mean±s.d. *P<0.05 and ***P<0.001.

Study approval
This study was performed with the approval of the institutional review board (CUMC09U034).
there was no alteration in miR-145 levels with IL-1β treatment. TNFα stimulation also failed to upregulate miR-145 expression in FLSs. However, recombinant TGFβ significantly increased miR-145 expression in both OA-FLSs and RA-FLSs, which is consistent with a previous report that TGFβ upregulates miR-145 expression in vascular smooth muscle cells.17 Altogether, we demonstrated that miR-143 and miR-145 were highly expressed in RA-FLSs and that miR-145 expression in RA-FLSs was upregulated by TGFβ stimulation.

Identification of target genes for miR-143 and miR-145 in RA-FLSs
To identify potential target genes that could be regulated by miR-143 and miR-145 in RA-FLSs, we performed global transcriptomic profiling (GSE49604) using microarray analysis in RA-FLSs, including the FLSs from which the miRNA data were produced.10 We selected a set of DEGs through the comparisons of RA-FLSs versus OA-FLSs and RA-FLSs with IL-1β versus OA-FLSs with IL-1β. As a result, 470 genes were upregulated in RA-FLSs and 135 genes were downregulated. The numbers in parenthesis refer to the number of included DEGs in each cluster. Red and blue denote high and low expression of DEGs, respectively (b) Venn diagram showing the relationship between two sets of predicted target genes for miR-143 and miR-145 and two DEG cluster sets of RA-FLSs versus OA-FLSs. The numbers of included genes are represented. (c) Network model of predicted target genes for miR-143 and miR-145 (miR-143: 404 genes, miR-145: 731 genes, shared: 54 genes, and total: 1081 genes). Node color refers to fold change of RA-FLSs versus OA-FLSs (red=up, cyan=down). Node size refers to -log10P of RA-FLSs versus OA-FLSs. Edge thickness indicates edge betweenness (upper panel). The table in the lower panel shows the fold change, probability of conserved targeting (PCT) score, and the degree of interactions of potential target gene candidates in the network model.
selected that had significant differential expression in each of the comparisons. Hierarchical clustering of 470 DEGs displayed shared and distinct expression patterns (Figure 2a). Among those expression patterns, we focused on genes with a shared expression pattern (highlighted with red or blue bars to the right of heatmap), since both miR-143 and miR-145 were upregulated irrespective of IL-1β stimulation (Figure 2a).

We then compared the two gene sets with shared up- or downregulation with predicted target gene lists of miR-143 and miR-145 that were obtained from TargetScan 6.0 (http://www.targetscan.org/vert_60/). As a result, 12 genes (ADAM19, ANTXR2, C13orf33, CREB5, EBF3, EYA4, IGFBP5, NUFBP2, PGD, SEMA3A, SOBP and TMOD1) were selected as potential candidates that could be up- or downregulated by miR-143 or miR-145 (Figure 2b).

To evaluate the target candidates, we utilized another approach to reconstruct a network model of 1081 genes that were predicted to be targets of miR-143 and miR-145 using TargetScan 6.0 (Figure 2c). We computed the degree of interaction of potential target gene candidates in the network model and identified three genes (IGFBP5, PGD and SEMA3A), which had more than five interactions with other genes in the network (Figure 2c, marked in red in the lower panel). The probability of conserved targeting (PCT) scores of the genes indicated that IGFBP5 and SEMA3A had higher PCT scores than PGD (Figure 2c, lower panel). Moreover, IGFBP5 and SEMA3A were significantly downregulated in RA-FLS with or without IL-1β stimulation, whereas PGD was upregulated.

Together, these results suggest that IGFBP5 and SEMA3A are the top two potential target candidates of miR-143 and
miR-145 in RA-FLSs. To confirm the relationship between the miRNAs and potential targets, we tested whether specific antagomirs of miR-143 and miR-145 increase IGFBP5 and SEMA3A expression levels in RA-FLSs. Mean levels of SEMA3A and IGFBP5 mRNA were significantly increased by inhibition of miR-145 and miR-143, respectively (for SEMA3A, 3.45-fold, \( P = 0.02 \); for IGFBP5, 1.31-fold, \( P = 0.02 \) (Figure 2d and e), indicating that SEMA3A and IGFBP5 expression may be regulated by miR-143 and miR-145 in RA-FLSs, respectively.

Validation of the miRNA-target relationship
It has been demonstrated that miR-143 directly binds to the 3′-UTR of IGFBP5 mRNA and thereby inhibits IGFBP5 expression in a post-transcriptional manner.18 Here we found that, in contrast to OA-FLSs, RA-FLSs showed a high degree of heterogeneity in miR-143 expression and could be divided into two subgroups according to miR-143 expression: (1) RA-FLSs with high levels of miR-143 (miR-143HIGH) and (2) ones with a low level (miR-143LOW). To investigate the relationship between miR-143 and its target IGFBP5 in RA-FLSs, we first compared IGFBP5 expression in the two subgroups of RA-FLSs. We found that IGFBP5 levels were significantly lower in the miR-143HIGH RA-FLSs than in the miR-143LOW RA-FLSs and OA-FLSs (Figure 3a). Moreover, consistent with a previous report,18 the expression of miR-143 and IGFBP5 mRNA in RA-FLSs displayed an inverse correlation (Figure 3b), suggesting that miR-143 negatively regulates IGFBP5 expression in RA-FLSs. An earlier study had described that RA-FLSs with low IGFBP5 expression displayed a myofibroblast phenotype with strong TGFβ signaling,19 which prompted us to investigate whether decreased IGFBP5 expression reflects a myofibroblast phenotype of RA-FLSs in our experimental setting. From microarray analysis, we found that IGFBP5 levels negatively correlated with expression levels of a variety of genes representing the myofibroblast phenotype with high TGFβ signaling, including ACTA2, COL3A1, COL4A2, POSTN, SPARC, CYR61, SERPINH1 and CALD1 (Figure 3c). Collectively, these results, together with a previous report19 suggest that low IGFBP5 expression linked to miR-143 is correlated with the myofibroblast phenotype of RA-FLSs.

To date, there has been no report demonstrating an interaction between SEMA3A and miR-145. Presently,
in contrast to IGFBP5 expression showing heterogeneity, a significantly lower expression of SEMA3A was consistently noted in RA-FLSs, independent of miR-145 expression (Figure 3d); this implied that other regulators might control SEMA3A levels in addition to miR-145. Therefore, it was unclear whether miR-145 directly bound to SEMA3A mRNA. To address this issue, we performed a dual-luciferase reporter assay using wild-type and mutant vector systems (Figure 3e). We found that a synthetic miR-145 mimic, but not a control mimic, significantly suppressed luciferase activity of the reporter vector expressing the wild-type 3′-UTR (Figure 3f). By contrast, a single mutation in the putative two miR-145-binding sites made luciferase activity resistant to the miR-145 mimic (Figure 3f). Moreover, this resistance was further enhanced by co-transfection of a double mutant of the two predicted miR-145-binding sites with the miR-145 mimic, which strongly suggests that miR-145 directly binds to the 3′-UTR of SEMA3A mRNA (Figure 3f).

IGFBP5 suppresses TNFα signaling in RA-FLSs

Next, we sought to determine the effect of diminished IGFBP5 expression on RA-FLS function. IGFBP5 was originally identified as a regulator of insulin-like growth factor (IGF) signaling. A recent study demonstrated that IGFBP5 also inhibited TNFα signaling by binding to TNFα receptor 1 (TNFR1) in histiocytic lymphoma cells. Therefore, we investigated whether IGFBP5 suppresses TNFα signaling in RA-FLSs. To this end, we first examined changes in TNFα-stimulated IL-6 production, a representative cytokine induced by TNFα, in FLSs following IGFBP5 siRNA transfection. We found that TNFα-induced IL-6 production was significantly higher in siIGFBP5-transfected FLSs than in control siRNA-transfected FLSs (1.8-fold change, \( P = 0.029 \) for RA-FLSs; 2.9-fold change, \( P = 0.033 \) for OA-FLSs) (Figure 4a).

It is well known that TNFα activates the NF-κB signaling pathway through IκBα degradation and p65 translocation. Thus, we tested the effect of IGFBP5 on p65 translocation and

![Figure 5](image-url)
endogenous VEGF165 secreted from RA-FLSs contributes to RA-FLSs (Figure 5a). Given our previous finding that recombinant human IGFBP5 (rhIGFBP5) significantly decreased p65 translocation to the nucleus (Figure 4c). Moreover, as shown in Figure 4d and f, the IκBα degradation rate in RA-FLSs was increased by the knockdown of IGFBP5 transcripts. Conversely, in the presence of rhIGFBP5, the degradation rate of IκBα induced by TNFα was significantly decreased (Figure 4e and f). Together, these results suggest that IGFBP5 inhibits TNFα-induced pro-inflammatory signals in RA-FLSs, suppressing NF-κB activity and IL-6 production. Therefore, the decreased IGFBP5 expression linked to high miR-143 noted in RA-FLSs may contribute to the increased downstream activation of TNFα signaling.

**SEMA3A inhibits the effect of VEGF165 in RA-FLSs**

SEMA3A acts as a chemo-repellent and a competitive inhibitor of VEGF165 through binding on its receptor, neuropilin-1. We previously demonstrated that the interaction of VEGF165 with neuropilin-1 protects RA-FLSs from apoptotic death by regulating Bcl-2 expression and Bax translocation. Moreover, SEMA3A was shown to block the VEGF165-induced increase in survival and migration of breast carcinoma cells. We therefore investigated whether SEMA3A has an inhibitory role in the survival of RA-FLSs as a VEGF165 antagonist. We found that recombinant human SEMA3A (rhSEMA3A) treatment inhibited the anti-apoptotic effects of VEGF165 in RA-FLSs (Figure 5a). Moreover, in the absence of recombinant VEGF165, rhSEMA3A significantly increased the number of apoptotic RA-FLSs (Figure 5a). Given our previous finding that endogenous VEGF165 secreted from RA-FLSs contributes to maintenance of cell survival, such an increase may be due to the inhibition of endogenous VEGF165 by SEMA3A, although it may also be hypothesized that SEMA3A itself induces FLS apoptosis independent of VEGF165. Moreover, RA-FLS death induced by sodium nitroprusside (SNP), an apoptosis inducer, was prevented by the addition of recombinant VEGF165, which was nearly completely abrogated by co-treatment with SEMA3A, as determined by MTT assay (Figure 5b). Together, these observations indicate that SEMA3A promotes RA-FLS apoptosis by inhibiting VEGF165 action.

Previously, we also have shown that VEGF165 promotes RA-FLS adhesion and migration, which are essential properties of RA-FLSs for progression of RA. As seen in Figure 5c, SEMA3A inhibited the VEGF165-induced increase in wound migration of RA-FLSs. Moreover, the VEGF165-promoted invasion of RA-FLSs in a Matrigel-plugged chamber was also almost completely blocked by co-treatment with SEMA3A (Figure 5d). These observations demonstrated that, in addition to its pro-apoptotic activity, SEMA3A suppresses the VEGF165-induced increase in migration and invasion of RA-FLSs.

Intriguingly, measuring the simultaneous levels of VEGF165 and SEMA3A mRNA in FLSs showed that the ratio of VEGF165 to SEMA3A was significantly higher in RA-FLSs than in OA-FLSs (Figure 5e), which demonstrates the relative over-expression state of VEGF165 compared to SEMA3A in RA-FLSs. Furthermore, SEMA3A mRNA levels negatively correlated with VEGF165 mRNA levels in FLSs (Figure 5f), suggesting that RA-FLSs are intrinsically more susceptible to the anti-apoptotic and pro-migratory activities of VEGF165.

Taken together, these data suggest that reduced SEMA3A linked to miR-145 overexpression in RA-FLSs leads to increased cell survival and increased migratory and invasive potential, resulting in tumor-like features for RA-FLSs.

**DISCUSSION**

Over the past decade, miRNAs have emerged as strong epigenetic regulators controlling pathological phenotypes associated with diseases involving cancer and autoimmune diseases. As miRNAs target multiple genes at the same time, it is difficult to identify the target genes responsible for miRNA-induced functional outputs. Thus, comprehensive profiling of both mRNA and miRNA expression levels is necessary to better predict target genes of miRNAs, providing greater insight into functional associations between miRNA and mRNA. In this study, we performed an integrated analysis of mRNA and miRNA profiles in RA-FLSs versus OA-FLSs, which facilitated the discovery of potential miRNA-target relationships. Moreover, network analysis of the predicted target genes provided the rationale for prioritizing the target genes. As a result, we found previously unidentified miRNAs, namely miR-143 and miR-145, that were upregulated in RA-FLSs compared to OA-FLSs. We also established a model for DEGs potentially regulated by miR-143 and miR-145 using TargetScan and successfully identified their targets, IGFBP5 and SEMA3A.

miR-143 and miR-145 are co-expressed miRNAs that are well-known potential tumor suppressors. In many cases, miR-143 and miR-145 are lower in expression levels in cancer
cells than in normal cells, as they target numerous oncogenes. Interestingly, a recent study has demonstrated that miR-143 and miR-145 are highly expressed in mesenchymal cells, including fibroblasts and smooth muscle cells, rather than epithelial cells, which suggests a cell-type-specific expression of miR-143 and miR-145 in primary cells. In the present study, we validated the high expression of miR-143 and miR-145 in independent RA-FLSs. This implies that the functional roles of miR-143 and miR-145 in RA-FLS may be different from those of various cancer cells. Moreover, miR-145 expression in RA-FLSs was increased by TGFβ stimulation. Given the elevated TGFβ in RA joints, this observation supports our results demonstrating increased miR-145 expression in RA-FLSs.

miR-143 directly binds to the 3′-UTR of IGFBP5 mRNA, directing reprogramming of fibroblasts into smooth muscle cells. A previous study of the gene expression profiles of RA-FLSs showed that IGFBP5 levels in RA-FLSs negatively correlated with high inflammatory synovitis and a myofibroblast phenotype. However, it is still unclear how IGFBP5 expression is regulated in RA-FLSs and whether it directly contributes to RA pathology. In the present study, IGFBP5 expression was heterogeneous across the RA-FLS population, as it was significantly lower in miR-143HIGH RA-FLSs than in miR-143LOW RA-FLSs and OA-FLSs. In addition, IGFBP5 levels negatively correlated with genes representing a myofibroblast phenotype. These results, together with a previous report, suggest that decreased IGFBP5 expression is epigenetically regulated as a miR-143 target and presumably is involved in the high inflammatory and myofibroblast phenotype of RA-FLSs. To address this issue, we tested the pro-inflammatory responses of RA-FLSs under IGFBP5-deficient and -excess conditions. We demonstrated that IGFBP5 deficiency resulted in upregulation of TNFα-induced NF-κB signaling in RA-FLSs, promoting IL-6 production. Conversely, treatment with recombinant IGFBP5 mitigated the TNFα downstream signaling. Collectively, these data suggest that IGFBP5-deficiency, linked to high miR-143 expression levels, renders RA-FLSs sensitive to TNFα (Figure 6). Given the high concentrations of TNFα in RA joints, an altered expression of IGFBP5 by high miR-143 levels in RA-FLSs may contribute to RA progression by promoting TNFα-induced NF-κB activation and IL-6 production, culminating in high inflammatory synovitis (Figure 6).

We have shown that VEGF165 is highly expressed in RA-SF and RA-FLSs, increasing survival and migration of RA-FLSs by binding to neuropilin-1. By contrast, SEMA3A, a well-known competitive inhibitor of VEGF165, is rarely expressed in RA synovial tissues, which raises the intriguing hypothesis that an imbalance between VEGF165 and SEMA3A expression causes synovial hyperplasia and angiogenesis, that is, the pannus formation, through a lifting of the block on VEGF165. SEMA3A was originally identified as an axonal guidance molecule, and it also suppresses cell migration and apoptosis by sharing receptors with VEGF165 in neural progenitor cells. A study has demonstrated that decreased SEMA3A expression in RA synovia negatively correlates with disease activity score 28 (DAS28) and high inflammatory synovitis. However, it remains unclear how SEMA3A expression is regulated in RA synovia and whether it can antagonize the anti-apoptotic and pro-migratory activity of VEGF165 in RA-FLSs. Here we identified for the first time that the 3′-UTR of SEMA3A mRNA is a direct target of miR-145 using a luciferase reporter assay. Moreover, we confirmed an imbalance between VEGF165 and SEMA3A expression in RA-FLSs and found that SEMA3A strongly suppressed the VEGF165-induced increase in survival, migration and invasion of RA-FLSs. Given the high concentrations of VEGF165 in RA joints, these results suggest that decreased SEMA3A linked to high miR-145 increases RA-FLS sensitivity to VEGF165 stimulation (Figure 6), adding another layer of VEGF165-mediated FLS pathology.

The pathologic hallmark of RA is the formation of an invasive pannus, which consists of synovial hyperplasia and extensive angiogenesis. RA-FLSs epigenetically acquire an aggressive phenotype and become the spearhead of the inflamed synovium, mediating chronic inflammation and destruction of the joints. Such aggressive behavior of RA-FLSs is regulated by diverse molecular mechanisms, including miRNAs. Here we provide new insights into the relationships between miR-143 and miR-145 signatures and their functional targets, IGFBP5 and SEMA3A, and define the pathological phenotypes of RA-FLSs, including pro-inflammatory, anti-apoptotic, pro-migratory and pro-invasive features. This study also offers new possibilities with regard to miR-143- and miR-145-based approaches for targeting RA-FLSs and for treating RA.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

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