**Name of Journal:** *World Journal of Stem Cells*

**Manuscript NO:** 76694

**Manuscript Type:** LETTER TO THE EDITOR

**Long noncoding RNAs in mesenchymal stromal/stem cells osteogenic differentiation: Implications in osteoarthritis pathogenesis**

Quintero D *et al*. lncRNAs in MSCs osteogenic differentiation

Daniel Quintero, Hugo C Rodriguez, Anish G Potty, Dimitrios Kouroupis, Ashim Gupta

**Daniel Quintero,** Department of Orthopaedics, Division of Sports Medicine, University of Miami, Miller School of Medicine, Miami, FL 33136, United States

**Hugo C Rodriguez,** Holy Cross Orthopedic Institute: Fort Lauderdale Practice, Oakland Park, FL 33334, United States

**Anish G Potty, Ashim Gupta,** South Texas Orthopedic Research Institute, Laredo, TX 78045 United States

**Dimitrios Kouroupis,** Diabetes Research Institute, Cell Transplant Center, University of Miami, Miller School of Medicine, Miami, FL 33136, United States

**Ashim Gupta,** BioIntegrate, Lawrenceville, GA 30043, United States

**Ashim Gupta,** Future Biologics, Lawrenceville, GA 30043, United States

**Author contributions:** Gupta A and Kouroupis D conceptualized the study; Quintero D, Rodriguez HC, Potty AG, Kouroupis D, and Gupta A outlined and designed the manuscript; Quintero D, Rodriguez HC, Kouroupis D and Gupta A drafted the manuscript; Potty AG, Kouroupis D and Gupta A critically reviewed and edited the manuscript; all authors approved the final version of the article for publication.

**Corresponding author: Ashim Gupta, MS, PhD, Director,** Future Biologics, 2505 Newpoint Pkwy, Suite 100, Lawrenceville, GA 30043, United States. [ashim6786@gmail.com](mailto:ashim6786@gmail.com)

**Received:** March 26, 2022

**Revised:** April 27, 2022

**Accepted: May 21, 2022**

**Published online:**

**Abstract**

This letter focuses on a recently published article that provided an exceptional description of the effect of epigenetic modifications on gene expression patterns related to skeletal system remodeling. Specifically, it discusses a novel modality of epigenetic regulation, the long noncoding RNAs (lncRNAs), and provides evidence of their involvement in mesenchymal stromal/stem cells osteo-/adipo-genic differentiation balance. Despite focus on lncRNAs, there is an emerging cross talk between lncRNAs and miRNAs interaction as a novel mechanism in the regulation of the function of the musculoskeletal system, by controlling bone homeostasis and bone regeneration, as well as the osteogenic differentiation of stem cells. Thus, we touched on some examples to demonstrate this interaction. In addition, we believe there is still much to discover from the effects of lncRNAs on progenitor and non-progenitor cell differentiation. We incorporated data from other published articles to review lncRNAs in normal progenitor cell osteogenic differentiation, determined lncRNAs involved in osteoarthritis pathogenesis in progenitor cells, and provided a review of lncRNAs in non-progenitor cells that are differentially regulated in osteoarthritis. In conclusion, we really enjoyed reading this article and with this information we hope to further our understanding of lncRNAs and mesenchymal stromal/stem cells regulation.

**Key Words:** Long noncoding RNAs; Epigenetics; Mesenchymal stromal/stem cells; Degenerative bone diseases; Osteoarthritis; Osteoporosis

Quintero D, Rodriguez HC, Potty AG, Kouroupis D, Gupta A. Long noncoding RNAs in mesenchymal stromal/stem cells osteogenic differentiation: Implications in osteoarthritis pathogenesis. *World J Stem Cells* 2022; In press

**Core Tip:** This letter summarizes that long noncoding RNAs (lncRNAs) are involved in mesenchymal stromal/stem cells (MSCs) osteo-/adipo-genic differentiation balance. We added that the interaction between lncRNAs and miRNAs is strongly involved in the regulation of the function of the musculoskeletal system, by controlling bone homeostasis and bone regeneration, as well as the osteogenic differentiation of stem cells. Additionally, MSCs/progenitor cells lncRNAs are involved in osteogenic differentiation, osteoarthritis pathogenesis, and lncRNAs in non-progenitor cells are differentially regulated in osteoarthritis.

**TO THE EDITOR**

We read with great interest the review article by Xia *et al*[1], titled “Epigenetic regulation by long noncoding RNAs in osteo-/adipo-genic differentiation of mesenchymal stromal cells and degenerative bone diseases”. We believe the article provides an exceptional description of the effect of epigenetic modifications on gene expression patterns related to skeletal system remodeling. Specifically, it discusses a novel modality of epigenetic regulation, the long noncoding RNAs (lncRNAs), and provides evidence of their involvement in mesenchymal stromal/stem cells (MSCs) osteo-/adipo-genic differentiation balance. We agree with the authors’ insight that lncRNAs are relevant to clinical practice as altered MSCs differentiation status can be implicated in the initiation/progression of various musculoskeletal pathologies such as osteoarthritis and osteoporosis. We do, however, have several clarifications we wish to provide.

In the introduction, MSCs are defined as “a heterogenous population of cells which include fibroblast, myofibroblast and progenitor cells”[1]. Even though this definition was previously introduced by International Society for Cell & Gene Therapy Mesenchymal Stromal Cell Committee[2], it can be mis-leading within the present article as authors evaluate the effect of lncRNAs on cells that possess differentiation capacity and not fully differentiated cells (such as fibroblasts). Instead, authors could introduce MSCs as mesenchymal stromal/stem cells are fibroblast-like cells capable of multilineage differentiation at least *in vitro* that possess strong paracrine and immunomodulatory properties *in vivo*. Additionally, even though MSCs are originated from a single cell population during embryogenesis, authors should acknowledge that MSCs show intrinsic propensities to osteo-/adipo-genic differentiation strongly related to their tissue of origin and functional MSC subset heterogeneity[3]. This may significantly affect the role of specific lncRNAs on the overall epigenetic regulation of MSCs differentiation.

In the present article authors have nicely presented the interactions between lncRNAs and epigenetic modifiers during osteo-/adipo-genic MSCs’ differentiation. However, in recent years the crosstalk between lncRNAs and miRNAs interaction has emerged as a novel mechanism in the regulation of the function of the musculoskeletal system, by controlling bone homeostasis and bone regeneration, as well as the osteogenic differentiation of stem cells[4]. We totally acknowledge that the topic of the present article is not miRNAs, however authors could elaborate more on this significant interaction. For example, ANRIL lncRNA was correlated with increased MSCs osteogenic differentiation in the present article. According to recent studies, the molecular mechanism of ANRIL lncRNA effects is based on its direct binding to circulating miR-7a involved in activating the NFKB signaling pathway[5]. Other lncRNAs that exert their osteoinductive activities on progenitor cells *via* binding to miRNAs are MALAT1 and PGC1β-OT1[6,7]. Similarly, HOTAIR lncRNA *via* miR-17-5p interaction inhibits osteogenic differentiation in individuals with a traumatic osteonecrosis of the femoral head. This is in relation to a variable activation of SMAD7 which directly influences osteoblastic differentiation[8]. On this basis of lncRNAs and miRNAs interactions, it seems that H19 lncRNA is a major regulator of MSCs osteogenic differentiation. Specifically, H19 lncRNA act *via* three modes of action: (1) up-regulate miR-675 expression and inhibit the phosphorylation of TGF-β1 and Smad3; (2) inhibit the expression of miR-141 and miR-22 and promote Wnt/β-catenin signal transduction pathway; and (3) inhibit the expression of miR-107, miR-27b, miR-106b, miR-125a, and miR-17 resulting in Notch signaling pathway regulation[9-11].

Pathological mechanisms of osteoarthritis (OA) development involve the interplay of different OA symptoms, including inflammatory and degenerative changes that lead to destruction of articular cartilage, deranged chondrocyte regeneration, osteophyte formation, subchondral sclerosis and hyperplasia of synovial tissue. Yet, we must make a distinction between lncRNAs expression in progenitor cells and lncRNAs expression changes in terminally differentiated cells such as chondrocytes as their implication on cell differentiation and protein expression are remarkably different. Herein, in addition to the present article data we incorporated data from other literature to: (1) review MSCs/progenitor cells lncRNAs involved in osteogenic differentiation; (2) determine MSCs/progenitor cells lncRNAs involved in OA pathogenesis; and (3) provide a review of lncRNAs in non-progenitor cells that are differentially regulated in OA.

On this basis, we identified four lncRNAs that are upregulated in MSCs/progenitor cells: DANCR, MALAT1, THRIL and LINC0051**;** and five lncRNAs are downregulated in MSCs/progenitor cells, specifically chondrogenic cell line ATDC5: XIST, NR024118, HULC, LncRNA-ATB, OIP5-AS1**.** A summary of these findings is featured in Figure 1 and Table 1[12-20].

lncRNAs strongly regulate chondrocytes expression patterns in both physiological and pathological conditions. Twelve different lncRNAs were upregulated in terminally differentiated chondrocytes. We summarize these findings in Figure 2 and Table 2[21-32].

In conclusion, we believe there is still much to discover from the effects of lncRNAs on progenitor and non-progenitor cell differentiation. We incorporated data from a recent review article by Ghafouri-Fard *et al*[33] among other articles to (1) review lncRNAs in normal progenitor cell osteogenic differentiation; (2) determine lncRNAs involved in OA pathogenesis in progenitor cells; and (3) provide a review of lncRNAs in non-progenitor cells that are differentially regulated in OA. We provided a superficial review of lncRNAs expression and osteoarthritis to clarify what was mentioned and separated the regulation in progenitor and non-progenitor cells, which was not previously published. Again, we really enjoyed the reading by Xia *et al*[1] and with this information we hope to further our understanding of lncRNAs and mesenchymal stromal/stem cells regulation.

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**Footnotes**

**Conflict-of-interest statement:** All the authors report no relevant conflicts of interest for this article.

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**Provenance and peer review:** Invited article; Externally peer reviewed.

**Peer-review model:** Single blind

**Corresponding Author's Membership in Professional Societies:** American Academy of Regenerative Medicine; American College of Sports Medicine; International Society for Extracellular Vesicles; American Society of Regional Anesthesia and Pain Medicine; North American Neuromodulation Society; and Orthopedic Research Society.

**Peer-review started:** March 26, 2022

**First decision:** April 25, 2022

**Article in press:**

**Specialty type:** Orthopedics

**Country/Territory of origin:** United States

**Peer-review report’s scientific quality classification**

Grade A (Excellent): 0

Grade B (Very good): B

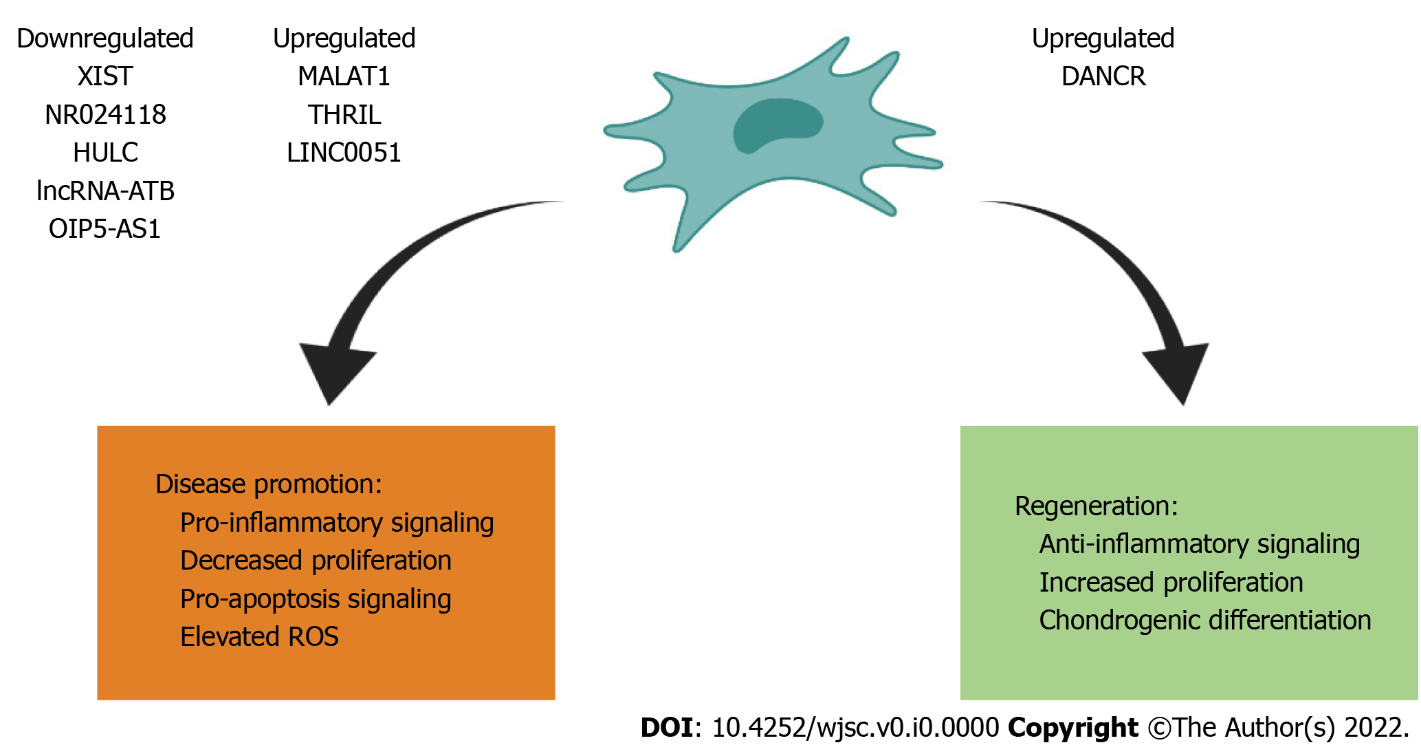
Grade C (Good): C, C

Grade D (Fair): 0

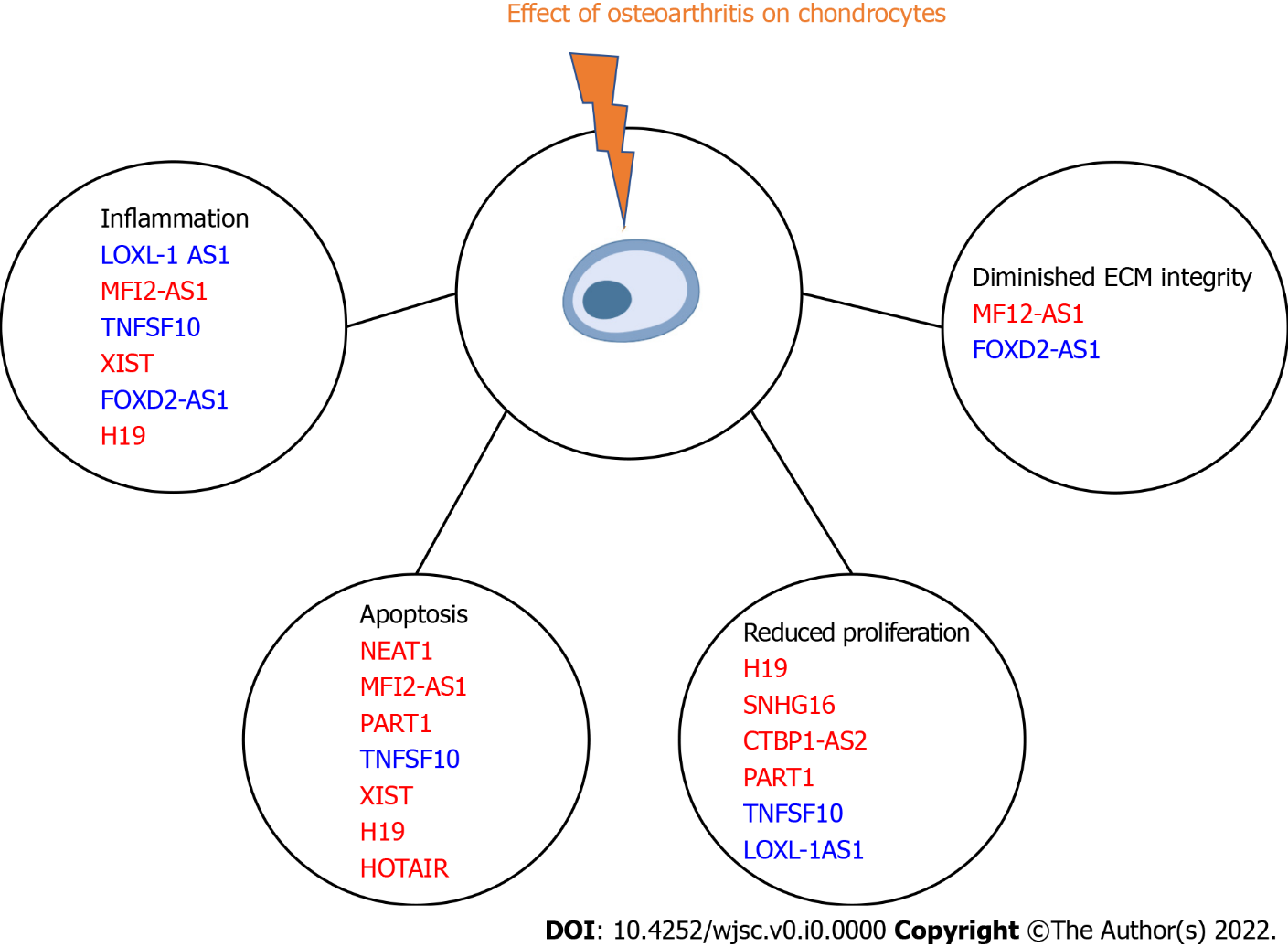
Grade E (Poor): 0

**P-Reviewer:** He BC, China; Niu ZS, China; Yao J, China **A-Editor:** Soriano-Ursúa MA, Mexico **S-Editor:** Gong ZM **L-Editor:** A **P-Editor:** Gong ZM

**Figure Legends**

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**Figure 1 Effects of various long noncoding RNAs on mesenchymal stromal/stem cells/progenitor cells for disease promotion and regeneration.**



**Figure 2 Effects of various long noncoding RNAs on chondrocytes in osteoarthritis.** Red text indicates promotion of pathogenesis, while blue text indicated regeneration by opposing pathogenic signaling. ECM: Extracellular matrix.

**Table 1 Supplementary information to Figure 1 detailing source and mechanism of activity associated with modified long noncoding RNAs**

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Upregulated** | | | **Downregulated** | | |
| **lncRNAs** | **Function** | **Ref.** | **lncRNAs** | **Function** | **Ref.** |
| DANCR | Increased proliferation and chondrogenesis | Wang *et al*[12], 2020 | XIST | Increased inflammation and apoptotic rate | Lian *et al*[13], 2020 |
| MALAT1 | Decreased rate of synovial fibroblast proliferation | Nanus *et al*[14], 2020 | NR024118 | Inflammation, apoptosis, and ROS elevation | Mei *et al*[15], 2019 |
| THRIL | Upregulated inflammatory injury and apoptosis | Liu *et al*[16], 2019 | HULC | Increased inflammation | Chu *et al*[17], 2019 |
| LINC0051 | Results in anti-proliferative actions | Zhang *et al*[18], 2020 | lncRNA-ATB | Increased inflammation | Ying *et al*[19], 2019 |
|  |  |  | OIP5-AS1 | Decreased cell proliferation and migration, decreased cell anti-inflammatory mediator secretion | Zhi *et al*[20], 2020 |

lncRNAs: Long noncoding RNAs.

**Table 2 Supplementary information to Figure 2 detailing source and mechanism of activity associated with modified long noncoding RNAs**

|  |  |  |
| --- | --- | --- |
| **lncRNAs** | **Function** | **Ref.** |
| ARFRP1 | Increased apoptosis related proteins | Zhang *et al*[21], 2020 |
| LOXL-1 AS1 | Improved inflammation and proliferation rate | Chen *et al*[22], 2020 |
| NEAT 1 | Increases apoptosis, decreases autophagy, decreases viability | Liu *et al*[23], 2020 |
| MFI2-AS1 | Increases inflammation, ECM degradation, and apoptosis | Luo *et al*[24], 2020 |
| PART1 | Low cell proliferation and increased cellular apoptosis | Zhu *et al*[25], 2019 |
| TNFSF10 | Improves cellular proliferation, anti-apoptotic, and anti-inflammatory actions | Huang *et al*[26], 2019 |
| XIST | Increases inflammation and apoptosis | Wang *et al*[27], 2019 |
| FOXD2-AS1 | Decreases inflammation, decreases ECM degradation | Wang *et al*[28], 2019 |
| H19 | Decreases proliferation, increases apoptosis, increases inflammation | Hu *et al*[29], 2019 |
| SNHG16 | Decreases proliferation | Fan *et al*[30], 2020 |
| CTBP1-AS2 | Decreases proliferation | Zhang *et al*[31], 2020 |
| HOTAIR | Increases apoptosis | He *et al*[32], 2020 |

ECM: Extracellular matrix; lncRNAs: Long noncoding RNAs.