Evidences for a New Role of miR-214 in Chondrogenesis

Vânia Palma Roberto¹,²,³,⁴, Paulo Gavaia¹,⁴, Maria João Nunes⁵, Elsa Rodrigues⁵,⁶, Maria Leonor Cancela¹,³,⁴ & Daniel Martins Tiago¹

miR-214 is known to play a role in mammalian skeletal development through inhibition of osteogenesis and stimulation of osteoclastogenesis, but data regarding other vertebrates, as well as a possible role in chondrogenesis, remain unknown. Here, we show that miR-214 expression is detected in bone and cartilage of zebrafish skeleton, and is downregulated during murine ATDC5 chondrocyte differentiation. Additionally, we observed a conservation of the transcriptional regulation of miR-214 primary transcript Dnm3os in vertebrates, being regulated by Ets1 in ATDC5 chondrogenic cells. Moreover, overexpression of miR-214 in vitro and in vivo mitigated chondrocyte differentiation probably by targeting activating transcription factor 4 (Atf4). Indeed, miR-214 overexpression in vivo hampered cranial cartilage formation of zebrafish and coincided with downregulation of atf4 and of the key chondrogenic players sox9 and col2a1. We show that miR-214 overexpression exerts a negative role in chondrogenesis by impacting on chondrocyte differentiation possibly through conserved mechanisms.

Most of the elements of the vertebrate skeleton are built through endochondral bone formation from a cartilage anlage. This complex process, involving chondrocyte commitment, proliferation, differentiation and hypertrophy, is governed by tightly orchestrated genetic and epigenetic programs and its disruption leads to pathological consequences. MicroRNAs (miRNAs) are small non-coding RNAs, usually transcribed by RNA Polymerase II (Pol II), which regulate gene expression by translational repression or by messenger RNA (mRNA) degradation. miRNAs have emerged as important regulators of skeleton formation where they exert multiple levels of control from cell fate decision, to proliferation, differentiation and cellular activities. In that sense, it is not surprising that skeletal key players, such as Runx2 and Sox9, are regulated by several miRNAs. Particularly, miR-214 was shown to inhibit bone formation by regulating Atf4, and to promote osteoclastogenesis by targeting Pten. More recently, the potential use of miR-214 as a therapeutic target in skeletal disorders was evidenced when miR-214 was shown to transit from osteoclast-derived exosomes to osteoblasts, and inhibit bone formation through Atf4 blockade. Interestingly, Atf4 not only is a master regulator of osteoblast function but also modulates chondrocyte proliferation and differentiation during endochondral bone formation. Although the putative role of miR-214 in chondrogenesis remains generally unknown, it is now accepted that the Dnm3 opposite strand (Dnm3os) transcript, which encodes miR-214 and miR-199a cluster, is crucial for normal mouse skeletal development, including cartilage formation. Moreover, miR-199a was reported to act as an inhibitor of chondrogenesis by targeting Smad. These reports suggest that a role for miR-214 in chondrogenesis is still to uncover.

In this study, we explored a putative role for miR-214 on cartilage formation. We found that miR-214 is expressed in the cartilage of zebrafish, and is downregulated during differentiation of murine ATDC5 chondrogenic cells. Also, we show that Ets1 activates Dnm3os promoter in undifferentiated cells. Upon miR-214 overexpression, Atf4 expression concomitantly decreased, suggesting that this gene could also be a target of miR-214 in chondrogenesis. Supporting this notion, miR-214 gain-of-function in zebrafish was shown to impair cranial cartilage formation, and this was accompanied by a downregulation of atf4 and of crucial cartilage markers, thus unveiling a negative impact of miR-214 on chondrogenesis.

¹Centre of Marine Sciences (CCMAR/CIMAR-LA), University of Algarve, 8005–139, Faro, Portugal. ²PhD Program in Biomedical Sciences, DCBM, University of Algarve, 8005–139, Faro, Portugal. ³Algarve Biomedical Center, University of Algarve, Campus de Gambelas, 8005–139, Faro, Portugal. ⁴Department of Biomedical Sciences and Medicine, University of Algarve, 8005–139, Faro, Portugal. ⁵Instituto de Investigação do Medicamento (iMed.ULisboa), Faculty of Pharmacy, Universidade de Lisboa, Av. Prof. Gama Pinto, 1649–003, Lisbon, Portugal. ⁶Department of Biochemistry and Human Biology, Faculty of Pharmacy, Universidade de Lisboa, Av. Prof. Gama Pinto, 1649–003, Lisbon, Portugal. Correspondence and requests for materials should be addressed to M.L.C. (email: lcancela@ualg.pt) or D.M.T. (email: dtiago@ualg.pt)
Results and Discussion

**Mir-214 expression is associated with skeleton formation of zebrafish.** The cluster miR-199a-2/miR-214 is transcribed from the opposite strand of Dynamin 3 (Dnm3), in a common primary transcript called Dnm3os. In mammalian models, this transcript was shown to control both osteogenesis and osteoclastogenesis. However, the putative role of miR-214 in chondrogenesis remains to be explored. In early development in zebrafish, miR-214 was previously shown to be expressed in somites and in the mesenchyme surrounding developing skeletal elements. Although these studies pointed towards a possible role of miR-214 in skeletogenesis, they failed to demonstrate a clear association with tissue calcification.

**miR-214 temporal expression correlates with skeletogenesis time-points.** To clarify this issue, we first analysed miR-214 expression throughout zebrafish development, from blastula to adulthood, focusing on crucial stages of skeletal formation (Fig. 1A). The following pattern of expression was observed: i) low levels of expression from 18-somite stage to 36 hpf; ii) progressive increase in expression from 2–6 days post fertilization (dpf), with a peak at 6 dpf (over 30-fold change compared to 24 hours post fertilization, hpf); iii) decrease in miR-214 expression at 15 dpf; iv) progressive increase from 15 to 60 dpf (reaching 110-fold increase compared to 24 hpf); and v) a general decrease in miR-214 expression in young adults (at 81 dpf; similar levels observed in male and female) (Fig. 1A). As expected, miR-199a had a similar pattern of expression (Supplementary Fig. S1), consistent with the fact that both miRNAs are originated from the same transcript (Dnm3os). This is also in agreement with previous studies showing similar patterns of expression for miR-199 and miR-214 in different systems. Interestingly, the patterns of expression found in this study were consistent with important time-points of skeleton formation during zebrafish development. Around 24 hpf, neural crest cells populate the two anterior pharyngeal arches, which will give rise to the craniofacial skeleton. By 3 dpf, the onset of craniofacial calcification is concomitant with the appearance of major cartilaginous structures in the head, which continue to expand until 6 dpf. From 6–15 dpf the craniofacial skeleton calcification substantially increases, and the calcification of the vertebrae is undergoing. At 30 dpf, all the skeletal structures are calcified. The first mononucleated osteoclasts appear around 15 dpf, followed by multinucleated osteoclasts at 60 dpf, thus sustaining an active bone remodelling throughout adulthood. The pattern of expression found for miR-214 during zebrafish development indicates that it might regulate distinct processes of skeleton formation, and further suggests that a tight regulation of miR-214 is required for a proper skeletal development. Nevertheless, one cannot disregard the putative involvement of miR-214 in other processes occurring simultaneously with skeletogenesis, not addressed in the scope of this study.

**miR-214 spatial expression correlates with skeletal structures.** To further understand a putative involvement of miR-214 in zebrafish skeletogenesis, the spatial component of miR-214 expression was analysed by in situ hybridization (Fig. 1B) at: i) 10 dpf, corresponding to the onset of vertebra calcification; ii) 20 dpf, when vertebra calcification is completed; and iii) 90 dpf, corresponding to young adult fish with active bone remodelling. miR-214 was detected in both skeletal and non-skeletal components of zebrafish body throughout development, confirming previous studies discussed next. Regarding non-skeletal components, miR-214 was detected in zebrafish brain, muscle and kidney (Fig. 1B), consistent with Dnm3os pattern of expression found during mouse development. In zebrafish, we also detected miR-214 expression in eye lens and retina (Fig. 1B), consistent with previous data obtained in X. laevis, suggesting that miR-214 could also be associated with cell fate in zebrafish retina. Overall, this miR-214 spatial distribution suggests a functional conservation in vertebrates. Concerning skeletal elements of zebrafish, miR-214 was found at sites where new bone is being formed, and also in several cartilaginous structures. Regarding cartilage, miR-214 expression was evident in the chondrocranium, in the pharyngeal cartilage and basal region of branchial filaments, in the ceratohyal, and in the basis of pectoral fins (Fig. 1B). As for newly forming bone, miR-214 was detected in arches and in the growth zones of the vertebral centra (Fig. 1B). MiR-214 was also expressed in the notochordal sheath and in scales (Fig. 1B). Our results are in agreement with previous data showing an association of Dnm3os with mouse cartilage. However, to the best of our knowledge, miR-214 was never detected in cartilage. Thus, our results provide the first evidence indicating a possible role of miR-214 in chondrogenesis. In addition, miR-214 was detected in zebrafish vertebral column, reinforcing the idea that miR-214 is important for the onset of calcification during development.

We further investigated the distribution of miR-214 in adult zebrafish calcified tissues, and argued if its expression could be comparable to other vertebrates. Thus, we analysed the expression of miR-214 in several calcified tissues from zebrafish (branchial arches, vertebra and skull) and mouse (cartilage from the ear, vertebra, calvaria and femur); given the known role of miR-214 in myogenesis, we decided to use muscle as a positive control. We found high levels of miR-214 expression in all skeletal tissues, although the highest level was observed in the muscle of both zebrafish and mouse. This result suggested a functional conservation of miR-214 in skeletogenesis between zebrafish and mammals. The next set of experiments aimed at exploring a possible conservation of miR-214 transcriptional regulatory mechanisms.

**Dnm3os promoter contains regulatory elements associated to chondrogenesis.** In the previous section we confirmed that miR-214 and miR-199a have similar temporal expression patterns, consistent with the fact that both miRNAs derive from the same transcript. Although Dnm3os is located on the opposite strand of a Dnm3 intron, expression of both miRNAs and Dnm3 gene was shown to be distinct in both zebrafish and mouse, suggesting that Dnm3os regulatory transcription unit is independent from Dnm3. Since the transcriptional mechanisms regulating Dnm3os were poorly explored in previous studies, we decided to clarify this point.
Comparative sequence analysis of the Dnm3os putative promoter region. We searched for conserved putative skeletal transcription factor binding sites (TFBS) in a 2.5 kilobase (kb) region upstream of pre-miR-199 (putative Dnm3os promoter) of human, mouse, Xenopus, medaka, stickleback, Tetraodon, fugu and zebrafish. Our analysis evidenced that the first ~850 base pairs (bp) of Dnm3os promoter (putative proximal promoter) were considerably more conserved, and contained most of the conserved TFBS, comparing to the more distant regions (putative distal promoter). Conserved binding sites for several TFs with known roles in skeletogenesis were identified: AP2alpha (transcription factor AP-2 alpha), CEBPs (CCAAT/enhancer binding proteins), AP2beta, GATA-4 and EGR1.

Figure 1. miR-214 expression correlates with skeletal elements in zebrafish. (A) Expression of miR-214 during zebrafish development, determined by miRNA qPCR. Values were normalized using zebrafish U6 small RNA and 24 hpf as reference sample and represent the mean ± s.d. of at least 3 independent replicates. hpf hours post fertilization, dpf days post fertilization, N.D. non-detected. Gap in the y-axis separates two different scales. (B) Detection of miR-214 by in situ hybridization in zebrafish with 10 (a), 20 (b, c, d) and 90 (e, f, g, h, i) dpf. From head to tail, miR-214 was detected in eye lens (arrowhead, a, b), retina (white arrowheads, a, b), brain (arrow, a), chondrocranium (asterisk, a, b), pharyngeal cartilage (white arrows, a, c), kidney (arrows, d), scales (arrow, e), muscle myotomes (arrowheads, e), cartilage in the base of pectoral fins (arrows, f), notochordal sheath (arrow, g), osteoid of haemal arches (arrows, h) and growth zones of vertebral body (arrowheads, i). Hybridization with negative control (scrambled) probe did not produce detectable signal, as observed in 90 and 20 dpf specimens (j and k, respectively). Scale bars: 0.2 mm for a, e, f and k; 0.1 mm for b, c, d, h, i and j; and 0.05 mm for g. (C) Relative expression of miR-214 in zebrafish (left panel) and mouse (right panel) adult tissues, determined by miRNA qPCR. Values were normalized using U6 small RNA and muscle as reference sample and represent the mean ± s.d. of at least 3 independent replicates (one-way Anova, different letters indicate statistical significance, p < 0.05). B. arches, Branchial arches. dpf days post fertilization.
ETS1 (ETS proto-oncogene 1, transcription factor), SP1 (family of transcription factors), SRF (serum response factor), TCF11 (also known as NFE2L1, nuclear factor, erythroid 2-like 1) and TWIST1 (twist family bHLH transcription factor 1) (Fig. 2A). Moreover, a conserved non-canonical TATA box (TATAT), present in seven out of eight species analysed, was identified 25 bp upstream the human transcriptional start site (TSS, GenBank accession NR_038397.2), and recognized as a putative binding site for TATA box binding protein, TBP (core matrix similarity = 0.90; similarity matrix = 0.75) (Fig. 2A and Supplementary Fig. S2A). Notably, the predicted binding sites for E1A

Figure 2. Dnm3os promoter is active in skeletal cell lines. (A) Schematic representation of Dnm3os gene (not scaled) and promoter region analysed (~850 bp). Putative promoter sequences of 8 vertebrates were aligned using CHAOS/DIALIGN, and then fed to ConTra v2: core match = 0.95, similarity matrix = 0.85, TRANSFAC database. TFBS predicted are indicated. Vertical lines indicate previously validated TFBS (by ChIP-assay) according to UCSC Genome browser. TSS, Transcriptional Start Site, based on human mRNA sequence; bp, base pairs. (B) Functional analysis of zebrafish dnm3os putative promoter activity in ABSa15 and ATDC5. Cells were transfected with zebrafish full (~1926 bp/+244 bp), partial (~657 bp/+244 bp), TATA-less (~1926 bp/~140 bp) promoter constructs, or promoter-less vector. (C) Functional analysis of human Dnm3os putative promoter activity in MC3T3 and ATDC5. Cells were transfected with human full (~2299 bp/+190 bp), partial (~641 bp/+190 bp), TATA-less (~2299 bp/~30 bp) promoter constructs, or promoter-less vector. For B) and C), a schematic representation of each construct and the respective putative TFBS are indicated on the left side. Values are the mean ± s.d. of at least 5 independent experiments; one-way Anova, ***p < 0.001.
miR-214 is downregulated during ATDC5 chondrogenic differentiation. To test the functionality of human and zebrafish putative Dnm3os promoters, the corresponding genomic regions were cloned and inserted upstream the luciferase reporter gene in pGL3-Base vector. TSS (+1) for human was deduced from the 5′ end of the Dnm3os transcript variant 1 available at GenBank (NR_038397.2), whereas for zebrafish this information was not available. Therefore, we performed a 5′RACE PCR to determine the 5′ end of zebrafish Dnm3os; the longest amplified sequence contained 244 bp upstream the first nucleotide of pre-miR-199a, and was considered as the TSS. Additionally, and considering our previous in silico analysis showing two distinct regions in putative Dnm3os promoter (proximal and distal), we have cloned genomic fragments corresponding to full (~2.5 kb; proximal plus distal) and partial promoters (~850 bp; proximal), plus TATA-less promoters, for human and zebrafish. Zebrafish promoter constructs in pGL3 were transfected into ABSa15, a fish bone-derived cell line (ECACC Ref. 13112201). In these cells the activity of the full promoter was 2 times higher comparing with the partial promoter (Fig. 2B), suggesting that the sequence from -1262 bp to -657 bp in zebrafish full promoter was sensitive to regulatory elements present in fish bone-derived cells. As expected, deletion of the putative TATA box significantly decreased luciferase activity to basal levels (Fig. 2B). The human promoter constructs were tested in mouse bone-derived cell line (MC3T3), where both full and partial promoters significantly increased luciferase expression to equivalent levels, and 45 times higher than empty vector (Fig. 2C). Again, the activity of the human promoter was almost abolished when the putative TATA box was deleted (Fig. 2C), indicating that the consensus TATA box here identified is functional in both species and crucial for Dnm3os transcription. Next we investigated promoter activities in an in vitro system that was extensively used in previous studies to investigate chondrocyte differentiation, i.e. the mouse teratocarcinoma-derived ATDC5 cell line28. Interestingly, full promoters of both zebrafish and human origins significantly increased luciferase activity to a higher extent than partial promoters (Fig. 2B,C), indicating that the distal region of these promoters should contain regulatory elements that are sensitive to transcriptional regulators present in ATDC5 cells. Our data indicates that both zebrafish and human promoters are active in chondrocyte- and osteoblast-like cell lines although to different extents, suggesting that Dnm3os is differentially regulated in distinct skeletal-derived cell lineages.

miR-214 is downregulated during ATDC5 chondrogenic differentiation. To better understand miR-214 regulation in skeletal cells, in particularly in chondrogenesis, we used the ATDC5 chondrocytic cell line. First, we characterized the levels of miR-214 in critical stages of ATDC5 differentiation: i) at confluence, when cells are committed to chondrocyte lineage but are still chondroprogenitors (T0); ii) at the condensation stage / beginning of cartilaginous nodules formation (T9); iii) during nodule maturation, when chondrocytes are embedded in the matrix (T21); and iv) during mineralization, the later phase of differentiation (T36; Supplementary Fig. S3). miR-214 was highly expressed in confluent cells but strongly downregulated (over 10-fold change) during early (T9) differentiation (Fig. 3B). In subsequent stages (T21 and T36), miR-214 expression was somewhat increased, but its levels remained low comparing to T0 (6-fold lower in T36) (Fig. 3B). Not only this pattern of expression suggested that miR-214 could play a negative role on chondrogenesis, similar to what was previously shown during osteogenesis in MC3T3 cells29, but also that its levels should be tightly regulated during chondrocyte differentiation.

Ets1 stimulates Dnm3os transcription in undifferentiated ATDC5. Due to the pattern of expression of miR-214 found in ATDC5 cells, and in order to confirm our results on Dnm3os transcriptional regulation, we performed a ChIP analysis of Dnm3os promoter in undifferentiated ATDC5 cells using antibodies against ETS1, SP1, TWIST1 and RNA polymerase II (positive control). SP1 is not expressed in undifferentiated ATDC5 cells (data not shown), and thus was used as a negative control. Surprisingly, ETS1 but not TWIST1 was enriched in Dnm3os promoter (Fig. 3C), along with RNA polymerase II, thus confirming that Dnm3os, and miR-214, are expressed in undifferentiated ATDC5 cells. However, the levels expression found for ETS1 during ATDC5 differentiation (Supplementary Fig. S4) are not in line with miR-214 expression pattern (Fig. 3B). This discrepancy suggests that although ETS1 might contribute for miR-214 expression in undifferentiated ATDC5 cells, other TFs should also be determinant. Such TFs could be important for miR-214 down-regulation during ATDC5 differentiation. A good candidate for this downregulation could be AP2α, as evidenced in Fig. 3A. Nevertheless, one cannot exclude other TFs that were not yet investigated.

Still, we show for the first time that ETS1 is enriched in the Dnm3os promoter of ATDC5 cells in a region containing an evolutionary conserved ETS1 binding site (Figs 2, 3, Supplementary Fig. S2). Not only this result...
contributed to uncover the transcriptional regulation of \textit{Dnm3os} (and \textit{miR-214}) in ATDC5 cells, but also it highlighted a possible role for ETS1 in chondrogenesis, a topic that remains controversial. Ets1 is a pivotal TF in the speciation and migration of neural crest cells, which can originate cartilage. However, while \textit{in vivo} data using

\textbf{Figure 3.} Analysis of \textit{miR-214} transcriptional activity in skeletal-derived cell lines. (A) Co-transfection of human \textit{Dnm3os} promoter and skeletal TFs. ATDC5 and MC3T3 cells were co-transfected with full promoter construct and either pCMX-TWIST1, pCMX-ETS1, pCDNA3-SP1 or pCDNA3-AP2\textgreek{a}, or empty vector (pCMX-PL2 or pCDNA3) as control. Results are indicated as fold change over the respective control empty vector (mean ± s.d. of at least 3 independent experiments; Student’s t-test, **p < 0.01). (B) Relative expression of \textit{miR-214} during ATDC5 cell chondrogenic differentiation. Levels of \textit{miR-214} were determined by miRNA qPCR, normalized using U6 small RNA expression and using day 0 (T0) as reference. Equal letters between two or more columns indicate values not statistically different, whereas different letters indicate values statistically different (mean ± s.d. of at least 3 independent replicates, one-way Anova, **p < 0.05). (C) TF binding to \textit{Dnm3os} proximal promoter analysed by Chromatin ImmunoPrecipitation (ChIP). Pre-cleared chromatin from undifferentiated ATDC5 cells was immunoprecipitated with either non-specific IgGs (IgG, negative control) or specific antibodies against Ets1, Sp1, Twist1, or RNA polymerase II (as positive control). After DNA recovery, the precipitates were assessed by qPCR. Data is expressed as percentage of total input and represent the mean ± s.d. of at least 3 independent experiments (one-way Anova, *p < 0.05).
Ets1-deficient mice evidenced the formation of ectopic cartilage nodules within the heart, suggesting that Ets1 can inhibit chondrocyte differentiation of cardiac neural crest cells\(^\text{39}\), *in vitro* studies in mouse mesencephalic neural crest cells showed that Ets1 is, on the contrary, able to promote chondrogenesis\(^\text{39}\). Moreover, contradictory results obtained in *X. laevis* model showed that both the overexpression and knockdown of Ets1 impaired cranial cartilage formation\(^\text{31}\). In fish, the role of Ets1 in skeleton formation remains to be explored. Nevertheless, in gilthead sea bream this TF was shown to regulate the expression of two important regulators of cartilage formation: *bmp2* and cartilage-specific *s100*\(^\text{12,23}\). Altogether, these results indicate that ETS1 could be important to control vertebrate skeletogenesis through a putative regulation of *Dnm3os/miR-214* expression.

**Negative effects of mir-214 on chondrogenesis.** Although recent studies have demonstrated that mir-214 plays a role in skeleton formation\(^\text{40,41}\), the involvement of mir-214 in chondrogenesis remains unknown. However, data collected in our study suggests that mir-214 might have a role in chondrogenesis.

*miR-214 mitigates the expression of chondrogenic markers during ATDC5 differentiation.* Gene expression analysis of ATDC5 cells indicated that mir-214 is differentially expressed during chondrocytic differentiation. The lower levels of expression during intermediate and later stages of differentiation suggest that mir-214 might be important to maintain chondrocytes in an undifferentiated condition. To further explore this possibility, and to get further insight into the role of mir-214 in chondrogenesis, we altered the expression of mir-214 in ATDC5 cells through gain and loss of function experiments. A miRNA mimic (MmiR-214) and an antagomiR (AmiR-214) with the respective controls (NC) were used to overexpress or down-regulate mir-214. ATDC5 differentiation occurred as previously reported\(^\text{14,35}\), as demonstrated by the expression of chondrogenic markers and von Kossa staining of cells (Supplementary Figs S3 and S4). QPCR analysis confirmed the altered expression of mir-214 in both experiments compared to NC at T14 (Fig. 4A, Supplementary Fig. S5). Surprisingly, at this stage the expression levels of Col2a1, Col10a1, Tnap, Sox9 and Sp7 were not affected by mir-214 overexpression (data not shown). On the contrary, forced expression of mir-214 significantly reduced *Mgp*, *Oc* and *Atf4* levels by approximately 20%, 60% and 40% respectively (Fig. 4B), suggesting that normal cell differentiation was most likely compromised. Although the role of Mgp in chondrogenesis is not fully understood, this protein was previously associated with chondrocyte proliferation and apoptosis\(^\text{35}\), being recognized as one of the main markers of chondrogenic differentiation *in vivo* and *in vitro*\(^\text{35,36}\). Conversely, Oc is mainly synthesized in osteoblasts and associated with bone formation, though its expression was also detected in chondrocytes and vascular smooth muscle cells, especially during mineralization\(^\text{37}\). In fact, Oc overexpression in ATDC5 was previously shown to stimulate cell differentiation and mineralization\(^\text{37}\). Therefore, in our experimental conditions, down-regulation of both Oc and Mgp upon mir-214 overexpression should represent a drawback in the differentiation process, probably with consequences at mineralization. Nevertheless, *Mgp* and *Oc* were still down-regulated upon loss-of-function of miR-214 (Supplementary Fig. S5), indicating that mir-214 mode of action in chondrogenesis might be more complex. Accordingly, precisely in the later stage of differentiation, the expression of mir-214 was slightly increased in WT cells (Fig. 3B), when *Mgp* and *Oc* levels are higher. Altogether, these data suggest that a tight regulation of mir-214 levels is required for a proper control of molecules impacting on chondrogenesis, such as *Mgp* and *Oc*, and consequent normal cartilage differentiation. Since bioinformatic analysis did not indicate *Mgp* or *Oc* as direct targets of mir-214 (data not shown), down-regulation of these genes was most likely indirect. On the contrary, the repression of *Atf4* by mir-214 overexpression in ATDC5 cells is likely a direct effect, since this regulatory mechanism was previously demonstrated in osteoblasts\(^\text{10}\) and upon antagoniR transfection *Atf4* levels were restored to those found in the control (Supplementary Fig. S5). *Atf4* is a crucial player in chondrogenesis and its ablation in mouse (*Atf4*\(^\text{−/−}\)) altered both proliferative and hypertrophic growth plate zones through control of Indian hedgehog (Ihh) expression\(^\text{14}\). Moreover, *Atf4* overexpression in chondrocytes of mouse *Atf4*\(^\text{−/−}\); *Col2a1−Atf4* double mutants positively regulated osteoblast differentiation during development through a paracrine mechanism also involving Ihh\(^\text{38}\). *Atf4* is a TF essential for the regulation of osteoblast differentiation and bone development\(^\text{31}\), known to drive the expression of Sp7, through a PTH-dependent mechanism\(^\text{39}\), and Oc, by cooperative interaction with Runx2 and Satb2\(^\text{40,41}\). *Atf4* is also one of the regulators of the neural crest cells migration, in cooperation with Sox9\(^\text{42}\). Thus, based on the important role of *Atf4* on chondrogenesis, and the inverse patterns of expression of *Atf4* and *miR-214* during ATDC5 differentiation (Supplementary Fig. S4 and Fig. 3B), we propose that *Atf4* could be a target of mir-214 in ATDC5 cells, probably contributing to a mitigation of cell differentiation. However, mir-214 is known to regulate genes and pathways with important functions in chondrogenesis and alternative targets should be considered. For instance, mir-214 regulates the Hedgehog pathway in zebrafish by targeting *sufu*\(^\text{54,56}\), and was shown to regulate the WNT pathway through direct regulation of *β*-Catenin in humans\(^\text{43}\). Importantly, both targets have conserved binding sites for miR-214 (data not shown). To further explore this possibility, we tested the effect of miR-214 overexpression on Ihh pathway by assessing the levels of a universal marker for activation of this pathway, Patched 1 (*Ptc1*). Indeed, *Ptc1* levels were significantly increased in ATDC5 cells overexpressing miR-214 compared to control cells (Supplementary Fig. S6), suggesting that this pathway could also be involved in miR-214 effects in chondrogenesis.

*miR-214 overexpression impairs zebrafish cartilage formation* *in vivo.* In order to confirm the chondrogenic role of *miR-214 in vivo*, and since our previous data revealed that *miR-214* was poorly expressed or absent in early stages of zebrafish development (Fig. 1A), we decided to overexpress this miRNA in zebrafish embryos by injecting 1-cell stage zebrafish eggs with *miR-214* mimic or negative control (NC). Embryos were analysed at 3 dpf, at the onset of zebrafish craniofacial skeleton formation and when skeletal structures are mainly composed by cartilage\(^\text{25,23}\). For all groups (MmiR-214, NC and WT), the highest mortality was observed at 24 hpf, although *miR-214*-injected embryos presented twice as much mortality (approximately 30%) comparing to NC (approximately 17%) and WT (approximately 12%). In subsequent days (2 and 3 dpf), mortality decreased drastically.
in all groups (0–1%). Embryos injected with MmiR-214 had a 5-fold miR-214 up-regulation over NC at 3 dpf (Fig. 5A). Since in previous studies, Atf4 seemed to be pivotal for miR-214 mechanism of action in the mammalian skeleton, we sought to investigate atf4 expression upon miR-214 overexpression. Importantly, we found that both atf4 paralogs in zebrafish, atf4a and atf4b, were decreased upon miR-214 overexpression (Fig. 5B). This was consistent with the detection of putative binding sites for miR-214 in both transcripts (Fig. 5C), and further supported the hypothesis that both genes are miR-214 targets in zebrafish. Gross morphology analysis revealed four phenotypes in miR-214-injected embryos comparing to NC: i) the embryos were generally smaller; ii) pericardium was enlarged; iii) the eyes were smaller and less developed; and iv) larvae presented alterations in the size and number of otoliths. For the analysis of structural deformities in cartilage, the embryos were stained with alcian blue, which marks the proteoglycans present in the cartilaginous matrix produced by chondrocytes. This analysis revealed different levels of phenotype severity, which could be due to distinct levels of miR-214 mimic incorporation in each embryo. While almost half of the embryos had severe deformities in the whole body (head

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**Figure 4.** Effect of miR-214 on the expression of marker genes for chondrocyte differentiation in ATDC5 cells. ATDC5 cells were transfected with 50 nM of MmiR-214 or NC, and differentiation was induced at confluence (T0) for 14 days. (A) Levels of expression of miR-214 were determined by miRNA qPCR analysis and normalized using U6 small RNA. (B) Levels of expression of Mgp, Oc and Atf4 were determined by standard qPCR and normalized using Hprt1 housekeeping gene (similar results were obtained using Hprt6 and Gapdh housekeeping genes; data not shown). Results are presented as fold change over NC. Asterisks indicate values statistically different from NC (data are the mean ± s.d. of at least 3 independent replicates; Student's t-test, ***p < 0.001, **p < 0.01, *p < 0.05).
and trunk), two thirds presented a clear flattening of the mandible (Fig. 5D,E), a predominant observation in the miR-214-injected embryos. Some embryos also presented a shortening of the trunk due to deformities in the notochord (Fig. 5E). Notably, all miR-214-injected embryos (n = 29) presented a reduced intensity of alcian blue staining indicating that the composition of the matrix in cartilaginous structures was altered and suggesting a defective chondrocyte function (Fig. 5D,E).

Figure 5. miR-214 ectopic expression downregulates atf4 transcripts and alters cranial cartilages of zebrafish. Larvae were microinjected at 1-cell stage with MmiR-214 or NC (18 μM) and analysed at 3 dpf. (A) Expression of miR-214 was determined by miRNA qPCR and normalized using U6 small RNA. (B) Expression of atf4a and atf4b in zebrafish larvae microinjected with MmiR-214 or NC. Levels of atf4a and atf4b transcripts expressions were determined by qPCR and normalized using 18S ribosomal RNA housekeeping gene (similar results were obtained using ef1α housekeeping gene; data not shown). Results are presented as fold change over NC. Asterisks indicate values statistically different from NC (data are the mean ± s.d. of at least 3 independent replicates; Student’s t-test, ***p < 0.001). (C) Predicted miR-214 binding sites in zebrafish atf4a and atf4b 3′ UTRs. Zebrafish atf4a and atf4b transcripts sequences were collected from NCBI database and analysed using RNAhybrid. (D) Morphological alterations observed on 3 dpf embryos injected with MmiR-214 (n = 29) or NC (n = 21), and wild type (WT; n = 21), stained with alcian blue. Alcian blue intensity was classified as very weak (−), weak (+), normal (++), or intense (+++), while deformities present in the embryos were classified as absent/normal fish (−), low severity (+), severe (++) or extremely severe (+++). Different letters indicate statistically significant differences between MmiR-214, NC and WT within the same classification (Chi-square test, p < 0.05). (E) Phenotype alterations observed in miR-214-injected embryos comparing to WT and NC-injected embryos, stained with alcian blue. Arrowheads indicate head deformations resulting from flattening of the mandible; arrows indicate differences on staining intensities of main cartilaginous structures (either absent or weak in miR-214-injected embryos). cbs, cerathobranchial cartilages; ch, ceratohyal; mc, Meckel’s cartilage. (1) or (2) indicate different embryos; (‘) indicate magnifications of the same embryo. Scale bar is 0.5 mm.
In order to get further insight into the effect of miR-214 overexpression in these larvae, we performed gene expression analysis for key markers of cartilage formation: sox9, sox10, col2a1, col10a1, runx2 and mgp. Notably, all markers were down-regulated in miR-214-injected embryos (Fig. 6A) demonstrating that in fact cartilage development was hampered. The early marker for cartilage formation, Sox9, is known to be important for neural crest cell commitment to the chondrogenic lineage, morphogenesis and differentiation of cartilage, chondrocyte stacking (sox9a) and cell number (sox9b) and chondrocyte hypertrophy. As a transcription factor, Sox9 also regulates the expression of genes crucial for cartilage development, such as Col2a1, Col10a1, Runx2 and Sox10. Thus, one cannot exclude that the repression of these genes in our experiment could be mediated by sox9 genes down-regulation itself. Nevertheless, these genes play crucial roles on cartilage development and maintenance, and their down-regulation in miR-214-injected embryos further supports a cartilage hindered phenotype. Finally, to further confirm our previous observations, we performed a histological analysis of six miR-214-injected embryos using safranin-O/fast green/Mayer's haematoxylin staining, to evaluate putative effects on the deposition of proteoglycans in the cartilage matrix. Gross morphological changes mentioned earlier, regarding embryos size, pericardium enlargement and underdevelopment of the eyes, were further confirmed through this analysis (Fig. 6B and Supplementary Fig. S7). Notably, five out of the six analysed embryos exhibited an almost complete absence of safranin-O staining (Fig. 6B and Supplementary Fig. S7) indicating a severe loss of reactivity with proteoglycans, and further suggesting that the cartilaginous matrix is altered and cartilage formation is impaired. Noteworthy, in all miR-214-injected embryos, the ethmoid plate, a cartilaginous structure considered a model for the mammalian palate formation, was absent (Fig. 6B). Moreover, the few cells reactive to safranin-O in the hyosymplectic of miR-214-injected embryos were clearly delayed in differentiation and lacked an organized arrangement when compared to the NC embryos (Fig. 6B). In fact, we could not detect the presence of mature chondrocytes in miR-214-injected embryos as evidenced in NC (Fig. 6B and Supplementary Fig. S7). Nevertheless, some cartilaginous structures, e.g. pharyngeal arch cartilage (Supplementary Fig. S7A and B), seemed to be properly placed and resembled a chondrocyte packing, but lacked reactivity to safranin-O in the miR-214-injected embryos. Other structures, e.g. pectoral fins, were present in the miR-214-injected embryos, but their cells were apparently less differentiated compared to NC, and lacked safranin-O staining (Supplementary Fig. S7C and D). In general, our histological analysis confirmed that chondrogenesis is impaired when miR-214 is overexpressed, and suggested that chondrocytes lost their ability to differentiate and produce the main components of cartilage ECM. This result is consistent with the previously observed down-regulation in the levels of expression of markers of cartilage formation. Altogether, data collected in this report pointed towards a putative negative role of miR-214 on vertebrate cartilage formation.

Conclusions

Herein, we show that miR-214 has a role in chondrogenesis and an underlying mechanism for miR-214 action is proposed. The pattern of expression of miR-214 during zebrafish development and ATDC5 chondrogenic differentiation suggested that this miRNA was most likely associated to chondrogenesis. This was evidenced by the high levels of miR-214 expression found during early stages of cartilage formation in zebrafish and in undifferentiated ATDC5 cells. Since miR-214 expression strongly decreased upon ATDC5 cell differentiation, we speculated that miR-214 could be a natural inhibitor of chondrogenesis. In fact, in vitro and in vivo gain-of-function experiments revealed that miR-214 was able to inhibit chondrogenesis, as evidenced by down-regulation of chondrogenic marker genes, and by the observation of phenotypic changes in cartilage formation in zebrafish miR-214-injected embryos. Loss-of-function experiments further supported that miR-214 levels should be important for proper chondrocyte differentiation, although it also revealed that the role of miR-214 might be more complex than originally expected. We propose that at least part of miR-214 effects in chondrogenesis are Atf4-dependent since this gene was consistently decreased upon miR-214 overexpression and unaltered upon miR-214 down-regulation, thus suggesting a direct repressive effect by miR-214 (Fig. 7), as previously validated by other authors. Interestingly, miR-214 overexpression lead to an overall down-regulation of Atf4 putative transcriptional targets such as Sox9, Col2a1 and Mgp, where we have identified several conserved putative-binding sites for Atf4, and also for Runx2, one of its skeletal cooperative partners (Supplementary Fig. S8). In addition, reporter assays testing the promoter of one of these genes, the MGP human promoter, evidenced that co-expression (through co-transfection experiments in ATDC5 cells) of Atf4 with its TF partners, Runx2 and Satb2, was able to enhance its promoter activity in ATDC5 cells (Supplementary Fig. S8). Even though, one must consider that miR-214 is also known to control signalling pathways crucial for chondrogenesis, such as the Hh and WNT pathways by targeting Sufu and β-Catenin respectively. These putative regulations might also contribute for miR-214 mode of action.

Based on our results, we propose that low levels of miR-214 are most likely required in order for chondrogenesis to initiate/proceed. High levels of miR-214 during early chondrocytic cell differentiation seem to delay this process by affecting the expression of important players of chondrogenesis, as Sox9. miR-214 mode of action could be in part explained by direct repression of Atf4. Once miR-214 expression decreases, Atf4 levels also increase, and chondrocytic differentiation proceeds (Fig. 7). Taken together, our results evidence for the first time that miR-214 could have an important role on chondrogenesis and cartilage formation, similarly to what was previously observed in osteogenesis.

Material and Methods

Ethics Statement. All operators involved in animal handling and experimentation are legally accredited by the Portuguese Direcção Geral de Alimentação e Veterinária (DGAV) and all the experimental procedures involving animals followed the EU (Directive 86/609/CEE) and National (Portaria no. 1005/92 de 23 de Outubro; Portaria no. 466/95 de 17 de Maio; Portaria no. 1131/97 de 7 de Novembro) legislation for animal experimentation.
Figure 6. miR-214 ectopic expression downregulates chondrogenic markers and impairs cranial cartilage formation in zebrafish. (A) Expression of marker genes of chondrogenesis. Expression levels of sox9a, sox9b, sox10, col2a1a, col10a1a, runx2a and mgp were evaluated by standard qPCR of 3 dpf zebrafish embryos microinjected with MmiR-214 or NC (18 μM), and normalized using 18S ribosomal RNA housekeeping gene (similar results were obtained using ef1α housekeeping gene; data not shown). Results are presented as fold change over NC. Asterisks indicate values statistically different from NC (data are the mean ± s.d. of at least 3 independent replicates; one-way Anova, ***p < 0.001; **p < 0.01, *p < 0.05). (B) Histological characterization of miR-214 overexpression on zebrafish cartilage formation. 3 dpf embryos microinjected with NC (a) or MmiR-214 (b) were embedded in paraffin, sectioned and stained with the safranin-O/fast green/Mayer’s haematoxylin. Zebrafish eyes (e) are generally underdeveloped in miR-214-injected embryos when compared to NC. Cartilage associated proteoglycans (evidenced by safranin-O) are either absent or present in low amounts in the Ethmoid plate (Ep) or hyosympletic (Hs), respectively, of miR-214-injected embryos comparing to NC. Sections are in the coronal plane; (’) and (’’) indicate magnifications of the same section.
and welfare. Animal experimental procedures were performed under the authorization 0421/000/000 from the DGAV complying with the decreto de lei 113/2013 de 7 de Agosto, from the Portuguese legislation.

Biological samples. Zebrafish eggs were obtained from natural spawning of a wild-type (AB) broodstock, and fish and larvae were maintained and raised using standard methods. For total RNA, pools of up to twenty zebrafish embryos, larvae and juvenile were collected at 1 k-cell (approximately 3 hours post fertilization, hpf), 18 somites (approximately 16 hpf), 24, 36 hpf, 2, 4, 6, 15, 30, 45, 60 and 81 days post fertilization (dpf), and from adult male and female. RNA was also isolated from different adult tissues of zebrafish specimens (muscle, branchial arches, skull and vertebra) and mice specimens (muscle, ear, femur, calvaria and vertebra). Animals were euthanized before sampling with 600 ppm MS222 (Sigma-Aldrich) for zebrafish and anesthesia followed by cervical dislocation for mouse.

RNA extraction, reverse-transcription and Real-time quantitative PCR (qPCR). Samples were collected in TRI Reagent (Sigma-Aldrich) (10 volumes/weight) and the total RNA extracted according to manufacturer’s protocol. For expression analysis of miRNAs, 1 μg of total RNA was polyadenylated, reverse-transcribed and amplified using miRNA-specific primers (Supplementary Table S1) using the NCode miRNA First-Strand cDNA Synthesis and NCode SYBR miRNA qPCR kits (Invitrogen), according to manufacturer’s instructions. For gene expression analysis, 1 μg of total RNA was reverse transcribed using MMLV-RT (Invitrogen), and transcripts amplified with gene-specific primers (Supplementary Table S1) and SsoFast EvaGreen supermix (Bio-Rad), according to manufacturer’s instructions. qPCR was performed using the StepOnePlus system (Applied Biosystems), unless stated otherwise, and relative gene expression was calculated using the ΔΔCt method.

Expression of miRNAs was normalized using the expression levels of U6 small nuclear RNA (U6), while mRNAs expression was normalized using expression levels of hypoxanthine phosphoribosyltransferase 1 (Hprt1) for mice samples and 18 S ribosomal RNA housekeeping gene for zebrafish samples.

In Situ Hybridization (ISH). Zebrafish larvae and juveniles at 10, 20 and 90 dpf were euthanized with a lethal dose of MS222 (Sigma-Aldrich) and fixed for 24 hours (h) in 4% paraformaldehyde (PFA) at 4 °C. Specimens were further decalcified in a 10% ethylenediaminetetraacetic acid (EDTA)/2% PFA solution for a minimum of 2 weeks and up to 2 months depending on their size. Samples were then washed in phosphate buffered saline (PBS) and maintained in 100% methanol at −20 °C until processing. For ISH, specimens were embedded in
paraffin and sectioned (4–6 μm thick). Detection of dre-miR-214 was performed using an ISH protocol adapted from the method described by Kloosterman et al., using LNA (Locked Nucleic Acid)-modified oligonucleotide 5′-Digoxigenin (DIG) labelled probe. Briefly, sections were fixed in 4% PFA and treated with 10, 20 or 40 μg/mL of protease K for specimens with 10, 20 and 90 dpf, respectively. After 2 h in pre-hybridization solution (50% formamide, 5 × saline sodium citrate buffer (SSC), 500 μg mRNA, 50 μg Heparin, 0.1% Tween and 9.2 mM of citric acid), sections were incubated with 40 nM of LNA ISH probe (Exiqon) specific for detection of dre-miR-214 (Supplementary Table S1). As negative control, sections were hybridized with a scramble probe (Supplementary Table S1). After 16 h incubation at 55 °C in a humidified chamber (5 × SSC), sections were washed with decreasing concentrations of formamide/SSC, then with PBST (PBS with 0.1% Tween-20) and incubated again for 1 h with blocking buffer (2% blocking solution from Roche diluted in maleic acid, 2% (v/v) sheep serum and 2% (m/v) bovine serum albumin (BSA)). Anti-DIG antibody conjugated with alkaline phosphatase (Roche, diluted 1:1600) was added to each section and incubated for 16 h at 4 °C. Each section was washed 5 times with PBST, 3 times with AP buffer (100 mM Tris, 100 mM NaCl, 50 mM MgCl₂, pH 9.5) and incubated with 75 mg/mL NBT/50 mg/mL BCIP in AP buffer for signal detection. Sections were air-dried, mounted with Eukitt (Sigma-Aldrich) and imaged by microscopy (Olympus IX-81 inverted microscope).

Identification and analysis of Dnm3os promoter sequences. Pre-miR-199a-2 sequences were retrieved from mirBase database (http://www.mirbase.org/) for: Homo sapiens, Mus musculus, Xenopus tropicalis, Takifugu rubripes, Orzya latipes, Gasterosteus aculeatus, Tetraodon nigroviridis and Danio rerio, blasted at Ensembl genome browser (www.ensembl.org) and genomic sequences with 2.5 kilobase (Kb) were collected starting immediately upstream of pre-miR-199a-2 (now on called Dnm3os promoter). Multiple sequence alignment of Dnm3os promoter was performed using CLUSTALW/DIALIGN (http://www.ddbj.nig.ac.jp/blast/blast_form.html) and fed to ConTra v2 (http://www.dnbr.ugent.be/prx/biotit2-public/contrav2/) for search of putative conserved TFBS. Stringency parameters: core match = 0.95 and similarity matrix = 0.85. The human, mouse and zebrafish promoter sequences were also fed to MatInspector (Genomatix Software GmbH) and only the TF predicted by the two algorithms were considered. Additionally, we analysed Chromatin Immunoprecipitation studies where some of the TFBS here predicted were found to physically interact with DNM3OS promoter, available through Genome Browser at UCSC (http://genome.ucsc.edu/), thus supporting our in silico analysis.

Cloning of zebrafish dnm3os 5′ end. The 5′ end of zebrafish dnm3os was achieved by rapid amplification of cDNA ends (RACE) using Advantage cDNA polymerase mix (Clontech) and a zebrafish Marathon cDNA library as template previously prepared57, according to manufacturer’s instructions (Clontech). Specific reverse primers (Supplementary Table S1) were designed based on the dnm3os pre-miR-199a-2 sequence available in mirBase (http://www.mirbase.org/) and combined with universal adapter primers (AP1 and AP2 universal primers; Supplementary Table S1). Amplified PCR products were subsequently inserted into pCRII-TOPO (Invitrogen) and further analysed by standard DNA sequencing.

Plasmid Constructs. Zebrafish (2.3 kb) and human (2.4 kb) Dnm3os promoter fragments were amplified using specific primers (Dre PR Fw and Dre PR Rev, Hsa PR Fw and Hsa PR Rev, in Supplementary Table S1) and genomic DNA as template. PCR products were cloned into pCRII-TOPO vector (Invitrogen), confirmed by DNA sequencing, and further used as templates to amplify new cDNA fragments containing specific deletions of the promoter of each species. Forward and reverse primers used to amplify these fragments contained 5′ ends with Nhel and BglII restriction site sequences, respectively, which were used for cloning into pGL3-Basic luciferase reporter gene vector (Promega). Constructs generated in pGL3-basic were as follow: full, partial and TATA less human promoters (−1926 bp to +2299 bp) and −190 bp to +190 bp and −1926 bp to −140 bp, respectively) and full, partial and TATA less human promoters (−2299 bp to +190 bp, −641 bp to +190 bp and −2299 bp to −30 bp, respectively).

The pCMX-ETS1 and pCMX-TWIST1 constructs were obtained by cloning cDNA fragments of zebrafish open reading frame (ORF) of v-ets avian erythroblastosis virus E26 oncogene homolog 1 (ets1) (nucleotides 129–1427, GenBank accession KP774190) and twist1a (nucleotides 114–629, Genbank accession NM_130984.2), from 48 hpf larvae and 69 dpf juveniles respectively, into pCMX-PL2 expression vector (kindly provided by Dr. Roland Schüle, Universitäts-Frauenklinik, Klinikum der Universität Freiburg, Freiburg, Germany). BamHI and Nhel restriction site sequences were incorporated into the forward and reverse primers, respectively, and the resulting PCR products were digested and cloned into the corresponding sites in pCMX-PL2 vector. Plasmids used to express TFs in mammalian systems were kindly provided by Dr Joseph P. Stains58 University of Maryland, Baltimore, MD, for pcDNA3-SP1 and pcDNA3 control plasmid, from Dr Ann Ehrlund 59 Karolinska Institutet, Department of Medicine, Huddinge, Stockholm for pcDNA3.1-TWIST1 and from Dr José Bragança, CBME, University of Algarve60 for pcDNA3.1-AP2.α. The identity of all constructs was confirmed by DNA sequencing analysis.

Cell culture. ABA15 cells were cultured in DMEM supplemented with 10% fetal bovine serum (FBS) and 0.2% fungizone, and incubated at 33°C in 10% CO₂. MC3T3-E1 and ATDC5 cells were cultured in α-MEM (supplemented with 10% FBS) and DMEM:F12 (supplemented with 5% FBS) respectively, and incubated at 37°C in 5% CO₂. Cells were sub-cultured every 2–3 days by trypsinization. All culture media were supplemented with 1% penicillin/streptomycin and 2 mM L-Glutamine. Cell culture media and FBS were obtained from Sigma-Aldrich and all other supplements and antibiotics were obtained from Gibco.

Dual-Luciferase Reporter Assays. Cells were seeded in 12-well plates, further cultured for 14–16 h, transfected with X-tremeGENE HP (Roche) according to manufacturer’s instructions, and as follows: ABA15 (8 × 10⁴ cells/well) with 500 ng of luciferase construct and 300 ng of pRL-SV40 vector (Promega), ATDC5 (1 × 10⁵ cells/well) with 500 ng of luciferase constructs and 50 ng of pRL-null vector and MC3T3 (8 × 10⁴ cells/well) with 500 ng
of luciferase construct and 200 ng of pRL-null vector. For co-transfections 50 ng of each TF construct or empty vector (control) was added to the previous conditions. 48 h after transfection, cells were lysed and luciferase activities were measured using Dual-Luciferase Reporter Assay system (Promega) in a Synergy 4 microplate reader (Biotek). Relative luciferase activity was determined as the ratio of firefly:renilla (F-Luc/R-Luc).

**Chromatin Immunoprecipitation.** ChIP assays were performed as previously described41 with minor modifications. Briefly, ATDC5 cells were seeded in 145 mm petri dish (Greiner; 1.2 × 10^7 cells per dish) and incubated for 24 hours at 37°C and 5% CO2. After cross-linking with formaldehyde, chromatin from ATDC5 cells was immunoprecipitated with specific antibodies against TWIST1, ETS1, SP1 (H-81 × , C-20 × and E-3 × respectively, Santa Cruz Biotechnology INC.),62-64 RNA polymerase II (ab5408, Abcam)65 anti-mouse IgG antibody (whole anti-serum, Sigma-Aldrich). The recovered DNA was analysed by qPCR in an ABI7300 sequence detection system (Applied Biosystems) with SYBR green Master Mix (Fermentas), using primers covering the proximal promoter region of the Dnmt3os gene (Supplementary Table S1).

**miR-214 overexpression during ATDC5 cell differentiation.** Cells were seeded in 24-well plates (2.5 × 10^4 cells/well), incubated for 16 h and transfected with miRIDIAN microRNA mimic for mmu-miR-214 (from now on designated MmiR-214) or negative control scrambled miR 1 (NC) (Dharmacon) at a final concentration of 50 nM, using EzWay (Komita Biotech) transfection reagent. Then, cells were grown until confluence (T0) and differentiation was induced by supplementing medium with ITS mixture (10 μg/mL Insulin, 5.5 μg/mL transferring, 6.7 ng/mL sodium selenite, Gibco) and replaced every 2–3 days. A second transfection was performed 10 days after the first one, using the same procedure. At appropriate times, cells were collected for total RNA extraction.

**Microinjection of zebrafish and in vivo effect of miR-214 overexpression.** One-cell stage embryos were microinjected with 4.6 nl of MmiR-214 or NC at 18°C (whole anti-serum, Sigma-Aldrich). The recovered DNA was analysed by qPCR in an ABI7300 sequence detection system (Applied Biosystems) with SYBR green Master Mix (Fermentas), using primers covering the proximal promoter region of the Dnmt3os gene (Supplementary Table S1).
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Author Contributions
V.P.R. and D.M.T. design and performed the experiments, analyzed the data and wrote the manuscript. M.J.N.
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