

Identification of Chloride Intracellular Channel Protein 3 as a Novel Gene Affecting Human Bone Formation

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ABSTRACT

Osteoporosis is a common skeletal disorder characterized by low bone mass leading to increased bone fragility and fracture susceptibility. The bone building cells, osteoblasts, are derived from mesenchymal stromal cells (MSCs); however, with increasing age osteogenic differentiation is diminished and more adipocytes are seen in the bone marrow, suggesting a shift in MSC lineage commitment. Identification of specific factors that stimulate osteoblast differentiation from human MSCs may deliver therapeutic targets to treat osteoporosis. The aim of this study was to identify novel genes involved in osteoblast differentiation of human bone marrow-derived MSCs (hMSCs). We identified the gene chloride intracellular channel protein 3 (*CLIC3*) to be strongly upregulated during MSC-derived osteoblast differentiation. Lentiviral overexpression of *CLIC3* in hMSCs caused a 60% increase of matrix mineralization. Conversely, knockdown of *CLIC3* in hMSCs using two short-hairpin RNAs (shRNAs) against *CLIC3* resulted in a 69% to 76% reduction in *CLIC3* mRNA expression, 53% to 37% less alkaline phosphatase (ALP) activity, and 78% to 88% less matrix mineralization compared to scrambled control. Next, we used an in vivo human bone formation model in which hMSCs lentivirally transduced with the *CLIC3* overexpression construct were loaded onto a scaffold (hydroxyapatite-tricalcium-phosphate), implanted under the skin of NOD-SCID mice, and analyzed for bone formation 8 weeks later. *CLIC3* overexpression led to a 15-fold increase in bone formation (0.33% versus 5.05% bone area relative to scaffold). Using a Clc3-His-tagged pull-down assay and liquid chromatography–mass spectrometry (LS/MS)-based proteomics analysis in lysates of osteogenically differentiated hMSCs, we showed that CLIC3 interacts with NIMA-related kinase 9 (NEK9) and phosphatidylserine synthase 1 (PTDSS1) in vitro, and this finding was supported by immunofluorescent analysis. In addition, inhibition of *NEK9* or *PTDSS1* gene expression by shRNAs inhibited osteoblast differentiation and mineralization. In conclusion, we successfully identified CLIC3 to be a lineage-specific gene regulating osteoblast differentiation and bone formation through its interaction with NEK9 and PTDSS1. © 2017 The Authors. *JBMR Plus* is published by Wiley Periodicals, Inc. on behalf of the American Society for Bone and Mineral Research.

KEY WORDS: CLIC3; OSTEOBLAST; MESENCHYMAL STEM CELL; DIFFERENTIATION; BONE; LINEAGE DECISION MAKING

Introduction

Bone is a dynamic organ that throughout life undergoes constant remodeling through bone formation by osteoblasts and bone resorption by osteoclasts. Osteoporosis is a common skeletal disease characterized by reduced bone mass and increased fragility and fracture risk, occurring when bone remodeling is disrupted and bone resorption overtakes bone formation.⁽¹⁾ The vast majority of osteoporosis treatments, such as bisphosphonates, act through reduction of bone resorption, and result in modest increases in bone density; however, these treatments do not result in a true bone anabolic effect so patients do not regain the bone that has been lost at time of

diagnosis. An ideal treatment would also stimulate osteoblast differentiation and/or bone formation to help repair the already damaged bone microarchitecture. Identification of novel genes or processes that stimulate osteoblast differentiation may therefore deliver therapeutic targets for the development of novel treatments for bone diseases such as osteoporosis.

Osteoblasts are derived from the bone marrow mesenchymal stem, or stromal, cells (MSCs) and undergo terminal differentiation to form osteocytes.⁽²⁾ Besides osteoblasts, MSCs give rise to other cell types, including adipocytes and chondrocytes.⁽³⁾ Osteogenic differentiation of MSCs is a tightly regulated process that progresses through several phases. MSCs are driven to the osteogenic lineage by expression of the osteoblast-specific

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Table 1. Oligonucleotide Primers Used for Proofreading PCR for Generation of His-Tagged CLIC3 Overexpression Vector

Gene	Forward (5'–3')	Reverse (5'–3')
<i>CLIC3-His</i>	CACCATGGCGGAGACCAAGCTCCA	TCATTACTAGTGATGGTGTATGGTATGGCGGGGTGCACGGCGGGCC

Sequences of forward and reverse primers used for proofreading PCR in this study. Detected using SYBR green.

transcription factors Runx2 and Osterix.^(4–6) After initial commitment, differentiation toward mature osteoblasts is typically characterized by the expression of collagen type I, osteocalcin, osteopontin, bone sialoprotein, and alkaline phosphatase (ALP).⁽⁷⁾ However, the precise mechanism of how human bone marrow-derived MSCs (hMSCs) differentiate into osteoblasts is still largely unknown, including the exact mechanism by which glucocorticoids stimulate osteoblast differentiation in vitro.^(8,9)

With aging there is a decrease in bone mass and increased bone fragility, while at the same time there is an increase in total and bone marrow adiposity.^(10,11) As both the osteogenic and adipogenic lineages derive from the same progenitor, it appears that there is a shift in the lineage decision-making process of MSCs with increasing age. This theory is supported by previous work showing that bone marrow-derived mesenchymal progenitors from aged humans, rats, and mice preferentially differentiate toward the adipogenic, rather than osteogenic, lineage.^(10–14)

Our aim was to expand the knowledge of human osteoblastogenesis by identifying novel genes and processes involved in differentiation of hMSCs to the osteogenic lineage. Gene expression analysis of hMSCs undergoing osteogenic and adipogenic differentiation led us to study chloride intracellular channel protein 3 (*CLIC3*) as an osteogenic lineage-specific candidate. CLIC proteins are a conserved family of proteins first identified when a novel chloride ion channel was discovered in bovine kidney.^(15,16) This family of proteins can transition between a soluble globular form and an integral membrane protein that is able to mediate ion conductance and/or ion channel formation.^(17,18) However, their role as channel proteins under physiological conditions has been strongly debated. To date, six CLIC paralogues have been identified in vertebrates, all of which appear to have diverse functions.^(18,19) *CLIC3* is a largely unstudied 26.6-kDa member of the CLIC family. It has been implicated to be involved in cellular processes including integrin recycling,^(20,21) endosome trafficking,⁽²²⁾ and cell growth via interactions with ERK7.⁽²³⁾

Materials and Methods

Cell culture

hMSCs were cultured as described⁽²⁴⁾ and are detailed in the Supporting Information.

Lentiviral-mediated overexpression and knockdown

To obtain *CLIC3* overexpression, we generated full-length human *CLIC3* cDNA (Open Biosystems; GE Dharmacon, Lafayette, CO, USA) containing a His-tag stop codon into a pEntr vector and verified using proofreading PCR (Table 1). hMSCs cells were transduced with the *CLIC3* vector or empty vector (pLenti6.3 vector without *CLIC3* construct), as control. For gene knockdown of *CLIC3*, NIMA-related kinase 9 (*NEK9*), and phosphatidylserine synthase 1 (*PTDSS1*), constructs from the TRC-Hs1.0 library (Sigma-Aldrich, Zwijndrecht, the Netherlands) were used (Table 2). A nontargeting short-hairpin RNA (shRNA) vector with a scrambled sequence (Table 2) was used as a

negative control. Methods for gene overexpression and silencing have been described^(25,26) and are specified in the Supporting Information.

ALP, mineralization, and protein assays

ALP and calcium measurements were performed as described^(8,24) and are described in detail in the Supporting Information.

Western blotting

Immunodetection of *CLIC3* by Western blot was performed as described⁽²⁷⁾ and is further detailed in the Supporting Information. The membrane was incubated with a specific antibody against *CLIC3* (ab56364; Abcam, Cambridge, UK; 1:500). Membranes were probed with the secondary antibody goat anti-mouse conjugated with Alexa Fluor 680 (1:5000; Invitrogen/Fisher Scientific, Landsmeer, Netherlands; Cat. A21057).

Quantification of mRNA expression

RNA isolation, cDNA synthesis, and PCR reactions were performed as described.⁽²⁴⁾ Oligonucleotide primer pairs were designed to be either on exon boundaries or spanning at least one intron (Table 3). Gene expressions were corrected for expression of the housekeeping gene *GAPDH*. Experiments were performed at least in duplicate in a minimum of two separate experiments.

Immunocytochemistry

Detection of *CLIC3*, *PTDSS1*, and *NEK9* by immunocytochemistry was performed as described⁽²⁸⁾; specific information can be

Table 2. List of shRNAs Used for *CLIC3*, *NEK9*, and *PTDSS1*

Target gene	ID	Target sequence
<i>CLIC3</i>	sh9123	GCCTCGTTACAGGGAGTCCAA
	sh9126	GCAGGAGAAAAGAGTCAAATA
<i>NEK9</i>	sh10	GCCTTGATTATTGTTGCAGTT
	sh11	CCGAGGAATGGAAGGTTTAAT
	sh12	CCAAAGGAACTCAGACAGCAA
	sh13	GTGAAGATCGTGCAAGGAATT
	sh14	GTACATTTGGAGAGTGGCATT
	sh15	CGAGCAGGTTAAATCTCTAAT
<i>PTDSS1</i>	sh25	GCTAGATCCAAATCTTCGATA
	sh26	TGGACCTATGTTTCGATGGTTT
	sh27	GCAACAACGAAAGCCATTCTT
	sh28	GACTGAGTTGAATACCTTCTT
	sh29	GACTGAGTTGAATACCTTCTT
Nontargeting shRNA	ID	Oligonucleotide sequence
Scrambled	SHC002	CCGGCAACAAGATGAAGAGCACCAA CTCGAGTTGGTCTTTCATCTTGT GTTTTT

Target sequences of shRNAs for gene silencing of *CLIC3*, *NEK9*, and *PTDSS1*, as well as the oligonucleotide sequence for a nontargeting scrambled control.

Table 3. Oligonucleotide Primers Used in the Study

Gene	Forward (5'–3')	Reverse (5'–3')
<i>GAPDH</i>	CCGCATCTTCTTTGCGTCG	CCCAATACGACCAAATCGTTG
<i>CLIC3</i>	CTGCCCATCCTGCTCTAT	CAGCGTCTCCTCCAGAAA
<i>NEK9</i>	TCAGCAATCCAGTGAGCAG	CCAGTCGTCATATTCGCCA
<i>PTDSS1</i>	TCGCCTTACCAGGGATGAC	GAGTGAACGGACCATTGGGG

Sequences of forward and reverse primers used for qPCR in this study. All genes were detected using SYBR green.

found in the Supporting Information. Primary antibodies used were: anti-CLIC3 raised in mouse, 1:150 (ab56364; Abcam, Cambridge, UK); anti-NEK9 raised in rabbit, 1:20 (11192-1-AP; Proteintech, Manchester, UK); anti-PTDSS1 raised in rabbit, 1:50 (HPA016852; Atlas Antibodies, Bromma, Sweden).

In vivo implantation assay

All animal experiments were performed in compliance with the animal ethics board of the Erasmus Medical Center. Experiments were performed based on the protocol by Abdallah and colleagues⁽²⁹⁾ and details can be found in the Supporting Information. Five healthy, adult, female NOD.CB17-Prkdc^{scid}/NcrHsd (NOD-SCID) mice (Charles River Laboratories, 's- Hertogenbosch, Netherlands) were used for these studies; each mouse had up to four implants surgically inserted subcutaneously under anesthesia. Quantification was carried out using Image J software (NIH, Bethesda, MD, USA; <https://imagej.nih.gov/ij/>): ceramics areas and bone areas were determined by eye based on staining and morphology, the edges were hand drawn and the resulting pixel measurements were calculated back to mm². The observer assessing the pellets was blinded toward their identity.

Pull-down assay

A pull-down assay was performed to isolate proteins associating with His-tagged CLIC3 using Dynabeads (10103D; Life Technologies, Bleiswijk, Netherlands). Purification of CLIC3 protein complexes was performed by a metal-based affinity between the Dynabeads and the His-tagged CLIC3. The pull-down was performed according to the manufacturers' protocol and exact details can be found in the Supporting Information. Bait protein was obtained from hMSCs differentiated with dexamethasone for 5 days, following transduction with either His-tagged *CLIC3* or an empty vector (EV; control). Two replicate samples from each condition were included for subsequent mass spectrometry (MS) assessment.

MS

MS detection of proteins binding to CLIC3-His was performed as described⁽³⁰⁾ and is described in full in the Supporting Information.

Bioinformatic analysis

The raw MS data were analyzed by MaxQuant software (version 1.3.0.5).⁽³¹⁾ A false-discovery rate of 0.01 for proteins and peptides and a minimum peptide length of six amino acids were required. The Andromeda search engine⁽³²⁾ was used to search the MS/MS spectra against the Uniprot database (taxonomy: *Homo sapiens*, release HUMAN_2013_04) concatenated with the reversed versions of all sequences (maximum of two missed cleavages; 0.6-Da fragment mass tolerance, enzyme specificity: trypsin). The

data from the replicates were combined as averages. For selection of the most relevant interacting proteins the following criteria were set: (1) a label-free quantification (LFQ) value in CLIC3 samples of greater than 1×10^6 ; (2) number of unique peptides covering a protein equals three or more; and (3) ratio of CLIC3 samples versus control samples of 1.5 or greater. Proteins were then ranked based on the average LFQ ratio of the CLIC3 samples versus the control samples. Ingenuity Pathway Analysis (IPA) (www.ingenuity.com) and Gene Ontology analysis (GO) in DAVID (<https://david.ncifcrf.gov>) was performed using the 52 proteins identified by pull-down and MS.

Statistics

The data provided here are based on at least two independent experiments performed at least in duplicate. Values displayed are mean \pm SE. Significance was calculated using either the Student's *t* test, Mann Whitney test, or the one-way ANOVA with Tukey's post hoc test where appropriate, using GraphPad Prism 6.0 (GraphPad Software, Inc., La Jolla, CA, USA). Values of $p < 0.05$ were considered significant.

Results

CLIC3 is upregulated during human osteogenic differentiation and downregulated during adipogenesis

CLIC3 is dynamically expressed in human mesenchymal stromal cells (hMSCs) during osteogenic differentiation (Fig. 1). Figure 1 shows that expression of *CLIC3* is increased during the osteogenic differentiation of hMSCs, peaking between 1 and 3 days of culture, compared to nondifferentiating. In addition we saw that *CLIC3* expression goes down in adipogenic cells compared to nondifferentiating, showing an opposite effect to that of osteoblasts. These data led us to further scrutinize *CLIC3* as a candidate gene for human osteoblastogenesis.

Overexpression of *CLIC3* in hMSCs enhances in vitro osteogenic differentiation

To determine the role *CLIC3* has in human osteoblastogenesis we overexpressed *CLIC3* in hMSCs by lentiviral transduction and studied its effect on classical biochemical markers of osteoblast

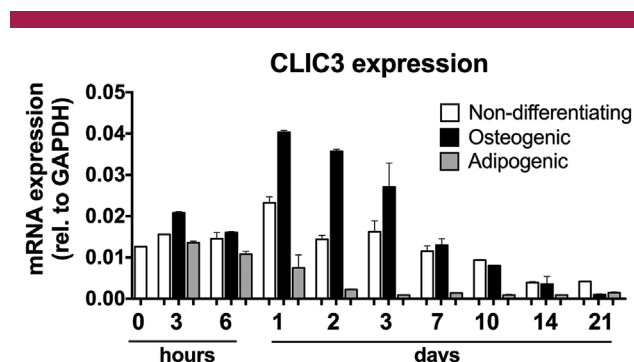


Fig. 1. *CLIC3* expression is upregulated during osteoblast differentiation and downregulated during adipocyte differentiation. mRNA expression levels of *CLIC3* in hMSCs cultured in nondifferentiating (white bars), osteogenic (black bars), and adipogenic (gray bars) conditions over 3 weeks assessed by quantitative PCR. Graph displays a representative experiment. $n = 2$.

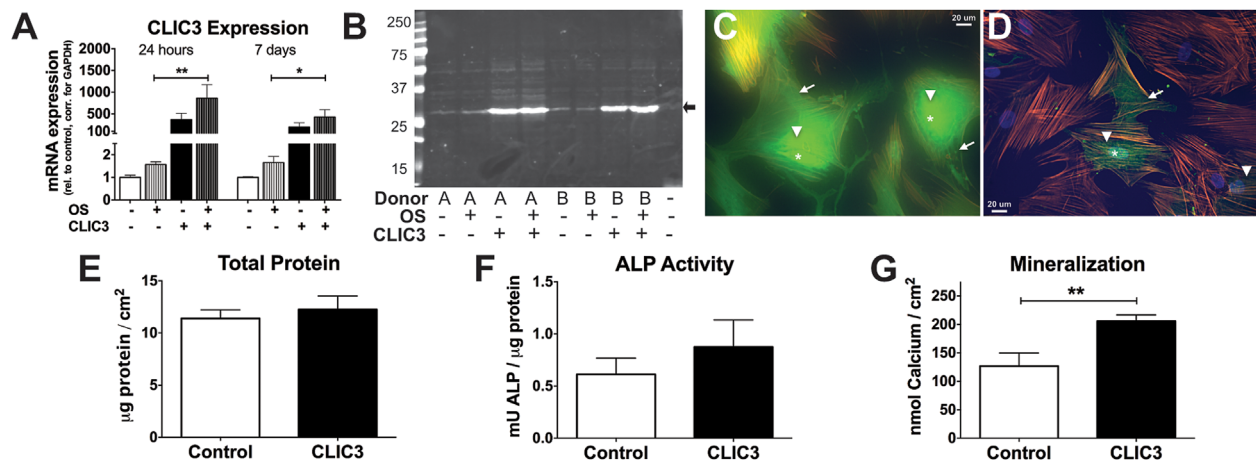


Fig. 2. *CLIC3* overexpression enhances human osteoblast differentiation. Enhancement of *CLIC3* expression was assessed by (A) quantification of *CLIC3* mRNA expression by quantitative PCR of hMSCs transduced with empty vector (white bars) or *CLIC3* overexpression vector (black bars) under nondifferentiating (solid bars) or osteogenic differentiation (patterned bars) conditions at 24 hours (left 4 bars) or 7 days (right 4 bars) after the start of differentiation, or by (B) immunoblotting in protein lysates collected 10 days postinduction of differentiation in two hMSC donors (A and B) using anti-*CLIC3* antibody. Arrow indicates the predicted size of *CLIC3* protein at 27 kDa, last lane contains loading buffer only. In additional *CLIC3* expression was visualized by immunofluorescence microscopy using *CLIC3* antibody in hMSCs differentiated to osteoblasts and transduced with *CLIC3* (C) or empty vector (D) at total magnification of $\times 400$ after 4 days of differentiation. *CLIC3*, in green, is seen in the cytoplasm (arrows), the perinuclear region (arrowheads), and some nuclear staining (asterisk); actin cytoskeleton was visualized by phalloidin antibody in orange, and nuclei by DAPI in blue. Biochemical assays are shown for total protein (day 6) (E), ALP activity (day 6) (F), and mineralization (week 3) (G) of hMSCs transduced with *CLIC3* (black bars) or empty vector (white bars) and osteogenic culture conditions. Graphs display combined results of all experiments. (A) $n = 8-12$; (E,F) $n = 18$; (G) $n = 19$. $*p < 0.05$, $**p < 0.01$. OS = osteogenic media. *CLIC3* = lentiviral transduction with *CLIC3* expression vector 1 day prior to the start of differentiation; DAPI = 4,6-diamidino-2-phenylindole.

differentiation. Efficient overexpression of *CLIC3* in hMSCs was determined by gene expression analysis and immunoblotting. Figure 2A demonstrates successfully elevated *CLIC3* mRNA expression by greater than 100-fold at both day 1 and 7 of culture in both nondifferentiating and differentiating hMSCs. Concordantly, protein levels of *CLIC3* were also strongly increased at day 10 of culture in nondifferentiating and differentiating hMSCs (Fig. 2B). Immunofluorescent *CLIC3* detection in differentiating hMSCs reveals that *CLIC3* overexpression (Fig. 2C) intensifies the level of *CLIC3* protein in most cells, compared to EV-transduced cells (Fig. 2D). *CLIC3* expression is predominately localized to the perinuclear region (arrowheads; Fig. 2C, D). In *CLIC3*-overexpressing cells (Fig. 2C) there is increased expression in the cytoplasm (arrows) and nucleus (asterisk) compared to control cells (Fig. 2D). *CLIC3*

overexpression did not affect total protein levels (Fig. 2E) or ALP activity (Fig. 2F) in osteogenic hMSCs, but it did enhance mineralization by 60% as shown by total calcium quantification after 3 weeks of culture (Fig. 2G). Overall, these results show that we have successfully overexpressed *CLIC3* in hMSCs and that *CLIC3* enhances in vitro mineralization.

CLIC3 is critical for in vitro human osteogenic differentiation

To determine if *CLIC3* is essential for human osteoblast differentiation we used shRNAs against endogenous *CLIC3* to knockdown its expression in hMSCs. Two separate shRNAs decreased *CLIC3* expression in hMSCs by 76% and 69% (Fig. 3A), respectively, compared to cells transduced with scrambled

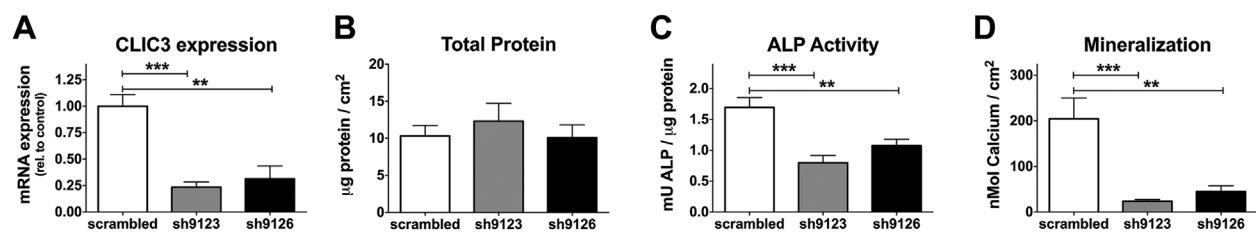


Fig. 3. Knockdown of *CLIC3* expression inhibits human osteoblast differentiation. Transduction of hMSCs with 2 shRNAs against *CLIC3* (shaded bars) reduces (A) mRNA expression of *CLIC3* as determined by quantitative PCR (day 4), (B) total protein (day 6), (C) ALP activity (day 6), and (D) mineralization (week 3) compared to scrambled control (white bar). Graphs display combined results from all experiments. (A) $n = 4-5$; (B) $n = 7$; (C) $n = 7$; (D) $n = 6$. $**p < 0.01$, $***p < 0.001$.

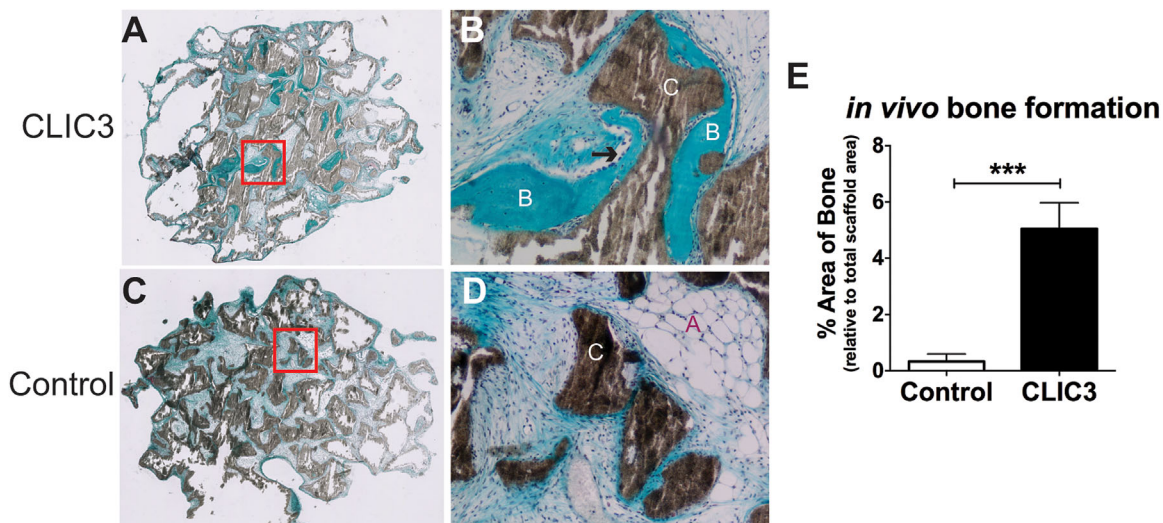


Fig. 4. *CLIC3* promotes human bone formation in vivo. Histological sections of explants of *CLIC3* overexpressing hMSCs (A, low magnification; B, high magnification) implanted under the skin of immune deficient (NOD-SCID) mice show more prevalent areas of new bone formation (labeled “B”) and osteoblasts (arrow) compared to empty vector-transduced cells (C, low magnification; D, high magnification). HA-TCP ceramic scaffold is labeled “C,” adipocytes are labeled “A.” (E) *CLIC3* overexpression in hMSCs (black bar) increases the amount of newly formed bone in explants compared to control cells (white bar) based on quantification of the area of new bone formed as a percentage of total scaffold area, using ImageJ software. $n = 9-10$. *** $p < 0.001$.

shRNA. Total protein levels were not affected by either shRNA (Fig. 3B). Knockdown of *CLIC3* mRNA expression reduced ALP activity by 53% and 37%, respectively, after 1 week of culture (Fig. 3C). After 3 weeks of culture in osteogenic medium, *CLIC3* silencing reduced mineralization by approximately 80% compared to the scrambled control (Fig. 3D).

CLIC3 enhances in vivo human bone formation

In order to test the in vivo effect of *CLIC3* manipulation on human bone formation we subcutaneously implanted hMSCs transduced with *CLIC3* or empty vector (control) loaded onto a hydroxyapatite/tricalcium phosphate (HA/TCP) scaffold in immune-deficient mice and quantified the amount of heterotopic bone formed. As shown in Fig. 4, hMSCs overexpressing *CLIC3* formed a significantly greater amount of ectopic bone compared to MSCs transduced with an empty vector. This is visualized in the representative Goldner-stained sections made from implants containing *CLIC3* overexpressing hMSCs (Fig. 4A, B) containing patches of newly formed bone and osteoblasts compared to the control-treated hMSC-implants (Fig. 4C, D) showing low level of newly formed bone. Quantification of the amount of new bone formed revealed that implants from *CLIC3*-overexpressing hMSCs contain 15 times more bone compared to implants containing control-treated cells (Fig. 4E). These results clearly show that *CLIC3* enhances in vivo bone human formation.

CLIC3 pull-down identifies 52 proteins as potential interacting with *CLIC3* in differentiating osteoblasts

In an attempt to gain insight into the molecular interactions by which *CLIC3* acts on osteoblasts, we performed a pull-down assay followed by MS analysis to identify which proteins in cell extracts of hMSCs differentiated toward osteoblasts associate with *CLIC3*. Supporting Table 1 lists all *CLIC3*-His pulled-down

proteins ranked according to the average LFQ intensity ratio of *CLIC3*-His-baited samples versus control samples, which are based on nonspecific binding. Bioinformatic analyses of these proteins were performed using Ingenuity Pathway Analysis and showed that “PRPP biosynthesis I,” “gap junction signaling,” “paxillin signaling,” “integrin signaling,” and various endocytosis and exocytosis pathways comprise the most significant canonical pathways in the list of proteins (summarized in Table 4, detailed in Supporting Table 2). Using DAVID Gene Ontology analysis, the most significant GO terms included processes related to RNA nuclear export and localization, transport, cell adhesion, and localization to the nuclear periphery and pores (summarized in Table 4, detailed in Supporting Table 3). Out of the 52 proteins identified by *CLIC3*-His pull-down, eight proteins were uniquely present in our *CLIC3* samples, as compared to control samples (Table 5), and we chose to analyze two of these proteins further: *NEK9* and *PTDSS1*.

CLIC3 interacts with *NEK9* and *PTDSS1* during osteogenic differentiation of MSCs

To confirm the colocalization of *CLIC3* with the proteins identified by the pull-down assay we performed immunofluorescent analysis. At day 5 of differentiation, *CLIC3* (Fig. 5A) and *NEK9* (Fig. 5B) colocalize in the perinuclear region (arrows, Fig. 5D) in *CLIC3* overexpressing hMSCs, with some *CLIC3* expression seen in the nucleus as well (asterisk, Fig. 5A), overlapping with DAPI nuclear staining (5C,D).

Similar to Fig. 5A, Fig. 5E exhibits the cytosolic (arrowhead), perinuclear (arrows), and nuclear distribution of *CLIC3* in osteogenically differentiated hMSCs, whereas *PTDSS1* is localized specifically to the perinuclear (arrows) and nuclear (asterisk, overlapping with DAPI nuclear staining (5G)) compartments (Fig. 5F), and colocalization is seen in the perinuclear region (Fig. 5H).

Table 4. Summary of Top of Canonical Pathways and GO Terms

Canonical pathway ^a	Biological process ^b	Molecular function ^b	Cellular component ^b
PRPP biosynthesis	Export from nucleus	RNA binding	Adherens cell junction
Gap junction signaling	RNA localization	Cadherin binding	Cytosol
Paxillin signaling	Transport	Cell adhesion: protein and molecule binding	Nuclear periphery/pore
Integrin signaling	Adhesion		
Endocytosis/exocytosis			

Top canonical pathways determined by Ingenuity pathway analysis from the 52 proteins found to pull down with Clic3-His. GO term clusters that have been identified by DAVID bioinformatics resources for the same proteins. The GO terms within the clusters had a significance of Benjamini corrected p value <0.05 . The headers "biological process," "molecular function," and "cellular components" stand for the GO category to which the GO term is annotated.

^aDetermined by Ingenuity pathway analysis (<http://www.ingenuity.com/>).

^bDetermined by DAVID Functional Annotation Analysis (<http://david.ncifcrf.gov>).

Inhibition of *NEK9* and *PTDSS1* affects osteogenic differentiation of hMSCs

To determine if *NEK9* or *PTDSS1* play a role in human osteoblast differentiation, we studied the effects of shRNA knockdown of *NEK9* and *PTDSS1* in osteogenically differentiating hMSCs. We found five shRNAs that all reduced *NEK9* mRNA expression between 60% and 82% compared to the scrambled control 4 days posttransduction (Fig. 6A). Four of the five shRNAs against *NEK9* inhibited mineralization by 70% or more after 3 weeks of osteogenic differentiation (Fig. 6B). Although the majority of *NEK9* shRNAs had no effect on total protein levels, one shRNA did decrease total proteins levels after 3 weeks of treatment (Fig. 6C). We identified five shRNAs against *PTDSS1* that all reduced mRNA expression in the range of 48% to 75% in differentiating hMSCs (Fig. 6D). We observed that three out of the five shRNAs against *PTDSS1* strongly inhibited mineralization, by 67% to 92% (Fig. 6E). The majority of shRNAs against *PTDSS1* had no effect on total protein levels; however, sh25-treated hMSCs displayed increased levels of total protein after 3 weeks of culture (Fig. 6F). Overall these results demonstrate that both *NEK9* and *PTDSS1* play a role in supporting osteoblast differentiation of hMSCs.

Discussion

CLIC3 was identified as a new gene specifically regulated in the osteogenic lineage of differentiating human MSCs. Lentiviral transduction-mediated overexpression and silencing of *CLIC3* during osteogenesis revealed a crucial function for *CLIC3* in promoting osteoblast mineralization. Overexpression of *CLIC3* in hMSCs strongly enhanced *in vivo* bone formation in a mouse model for ectopic human bone formation, further emphasizing that *CLIC3* plays an important role in human osteoblast differentiation. Bioinformatics analysis of proteins identified by *CLIC3*-His pull-down suggests *CLIC3*'s role during osteoblast differentiation may be related cytoskeletal associations and signaling, cell adhesion, and/or nuclear pore formation or transport of proteins or ions through nuclear pores. Finally, we identified that *CLIC3* interacts with *NEK9* and *PTDSS1* during osteoblast differentiation, and inhibition of the *NEK9* and *PTDSS1* expression reduces osteogenic differentiation of hMSCs.

CLICs are a diverse group of proteins, having been associated with a wide variety of biological processes and cellular functions including: angiogenesis⁽³³⁾; macrophage activation⁽³⁴⁾; DNA damage⁽³⁵⁾; maintenance of membrane structure⁽³⁶⁾; bone

Table 5. Top Proteins Identified in *CLIC3*-His Pull-Down

Protein	Gene symbol	LFQ CLIC3_A	LFQ CLIC3_B	LFQ EV_A	LFQ EV_B	Number of unique peptides	Ratio LFQ CLIC3:EV
Chloride intracellular channel protein 3	CLIC3	1.27E+09	1.19E+09	nd	nd	15	1.23E+09
Serine/threonine-protein kinase Nek9	NEK9	6.31E+06	5.12E+06	nd	nd	3	5.71E+06
1-Phosphatidylinositol 4,5-bisphosphate phosphodiesterase beta-3	PLCB3	2.84E+06	4.29E+06	nd	nd	7	3.56E+06
Aminoacyl tRNA synthase complex-interacting multifunctional protein 1	AIMP1	2.49E+06	3.29E+06	nd	nd	3	2.89E+06
Casein kinase II subunit alpha 3; alpha 1	CSNK2A1; CSNK2A3	1.70E+06	3.00E+06	nd	nd	3	2.35E+06
40S Ribosomal protein S28	RPS28	1.64E+06	1.97E+06	nd	nd	3	1.80E+06
Phosphatidylserine synthase 1	PTDSS1	1.21E+06	1.73E+06	nd	nd	3	1.47E+06
Nuclear pore complex protein Nup160	NUP160	1.15E+06	1.06E+06	nd	nd	3	1.11E+06

Proteins identified by MS from *CLIC3*-His pull-down assay in hMSCs undergoing osteogenic differentiation. Protein lysates from osteogenically differentiated hMSCs were isolated on day 5 in cultures either overexpressing His-tagged *CLIC3* or transduced with EV control, and subjected to protein pull-down for the His-tagged *CLIC3* and proteins binding to it. Proteins were determined by mass spectrometry measurements ($n = 2$). The top proteins are listed here, ranked on their ratio of *CLIC3* versus control average LFQ intensity (ie, present in *CLIC3* overexpressing condition and absent in the EV condition).

LFQ = label-free quantification; EV = empty vector; nd = not detected.

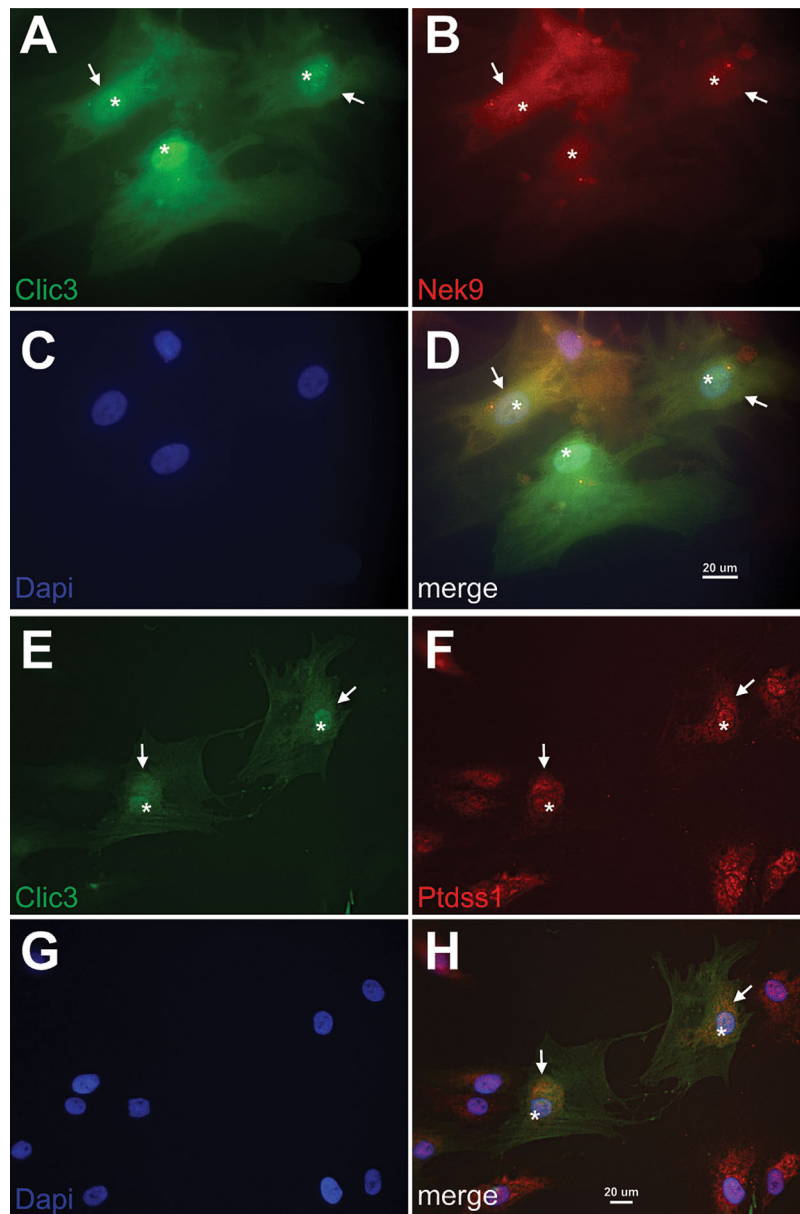


Fig. 5. CLIC3 colocalizes with PTSS1 and NEK9 in differentiating osteoblasts. Using immunofluorescence microscopy, CLIC3 (A, D, green) and NEK9 (B, D, red) were visualized together in hMSCs differentiated toward osteoblast and overexpressing CLIC3 at magnification $\times 630$. C, D show nuclear staining by DAPI in blue. Arrows indicate detection of both proteins in the perinuclear region and asterisks denote nuclear localization. Immunofluorescent detection of CLIC3 (E, H, green) and PTSS1 (F, H, red) were visualized jointly in osteogenic hMSCs overexpressing CLIC3 at magnification $\times 400$. G, H show nuclear staining by DAPI in blue. PTSS1 and CLIC3 are both seen in the perinuclear (arrows) and nuclear (asterisks) compartments. DAPI = 4,6-diamidino-2-phenylindole.

resorption⁽³⁷⁾; cell growth, cell division, and apoptosis^(23,38,39); and acidification of intracellular organelles.^(33,40) All CLIC proteins contain a ~ 240 -residue CLIC module that adopts a glutathione S-transferase (GST) superfamily fold⁽⁴¹⁾; within it there is conserved cysteine residue at what looks like an enzymatic active site.^(18,19,42) CLIC3 was first identified based on its interaction with MAPK15, at the time known as ERK7, in a yeast two-hybrid assay and subsequently confirmed by coimmunoprecipitation in COS cells; Qian and colleagues⁽²³⁾ proposed that it plays a role in controlling mammalian cell growth based on its association with MAPK15. This is the first

report of CLIC3 being involved in osteoblastic differentiation of hMSCs and human bone formation. Our in vitro results revealed the important role for CLIC3 in osteoblast differentiation and the strong induction of bone formation by hMSCs after CLIC3 overexpression seen in our model of human ectopic bone formation in a mouse supports the strong role CLIC3 has in ossification. It is worth noting that although this model provides an in vivo environment to study human bone formation, it is limited by the fact that the scaffold and cells are not within their native bone microenvironment and specific factors and vascularization may be lacking, resulting in lower bone

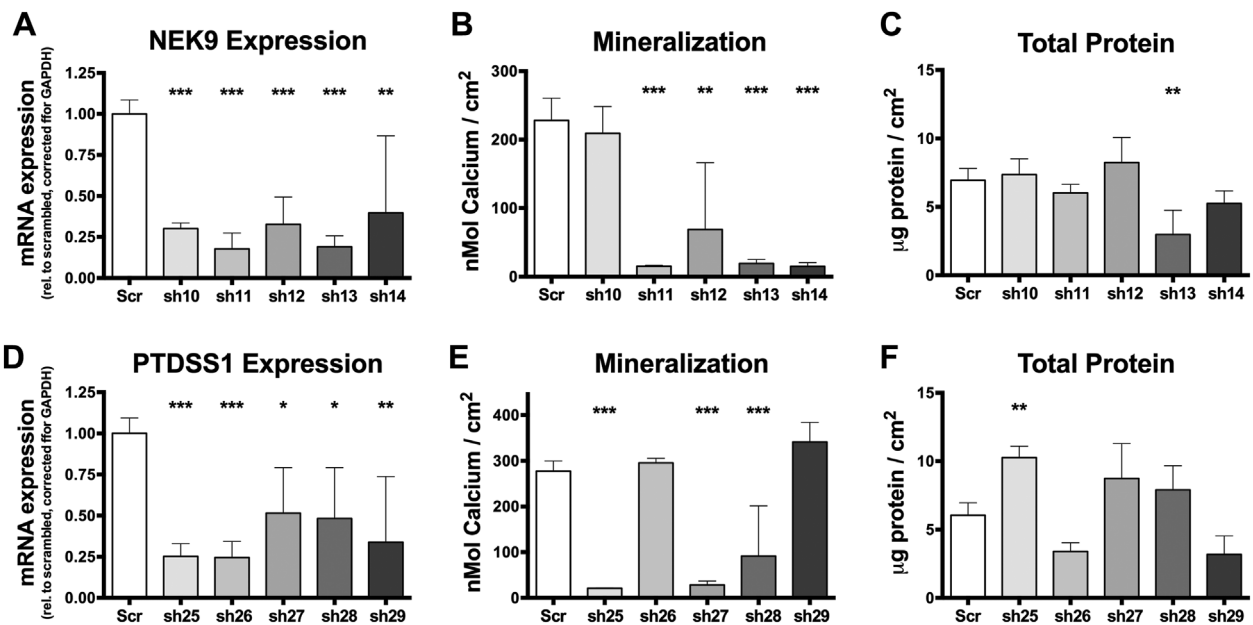


Fig. 6. Knockdown of *NEK9* and *PTDSS1* gene expression inhibits osteogenic differentiation of hMSCs. mRNA expression of *NEK9* (A) as determined by quantitative PCR (day 3), mineralization (B) (week 3), and total protein quantification (C) (week 3) after transduction of hMSCs with five shRNAs against *NEK9* (shaded bars) compared to scrambled control (white bars). mRNA expression of *PTDSS1* (D) as determined by quantitative PCR (day 3), mineralization (E) (week 3), and total protein quantification (F) (week 3) after transduction of hMSCs with five shRNAs against *PTDSS1* (shaded bars) compared to scrambled control (white bars). Graphs display combined results from all experiments. (A) $n = 4-12$; (B) $n = 6-12$; (C) $n = 6-12$; (D) $n = 4-12$; (E) $n = 5-12$; (F) $n = 6-12$. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$.

formation overall. Our findings, which are in line with previous reports,^(22,23) show that CLIC3 is predominately localized to the perinuclear and nuclear regions, supporting our bioinformatics findings suggesting a role for CLIC3 in the nuclear pore complex and/or export of RNA and ribonucleoprotein complexes from the nucleus. Like our results presented here, CLIC1, which shares a high degree of homology to CLIC3, with 48% to 49% of the protein sequences being identical,⁽²³⁾ was reported to be important for murine osteoblast differentiation through MSC lineage decision-making. CLIC1 was upregulated in response to Wnt treatment and downregulated following adipogenic treatment in murine MSCs; overexpression of *CLIC1* led to increased ALP activity and mineralization as well as increased expression of ALP, osterix, and osteocalcin.⁽⁴³⁾ The authors showed that CLIC1 overexpression caused hyperpolarization of the mitochondrial membrane potential, suggesting that its role in osteogenic differentiation is linked to supporting energy supplementation. Based on their structural similarities, it would be possible to postulate that the two proteins may have similar functions; however, we do not have any evidence thus far that CLIC3 functions in regulating mitochondrial membrane potential or energy metabolism. Although the detailed mechanism by which CLIC3 regulates human osteoblast differentiation is still far from understood, the discovery of its interaction with NEK9 and PTDSS1 during osteogenesis does give us some clues to its function.

After discovering that NEK9 associates with CLIC3 using protein pull-down and immunofluorescence, we confirmed that NEK9 is important in osteogenic differentiation by showing that *NEK9* knockdown leads to reduced mineralization. Very recently it was reported that recessive mutations in *NEK9* cause a lethal

skeletal dysplasia characterized by fetal akinesia, multiple contractures, shortened long bones, thoracic dysplasia, pulmonary hypoplasia, and protruding abdomen.⁽⁴⁴⁾ This stop-gain mutation results in a truncated protein that misses the majority of the RCC1-like domain, all of the NEK6 interaction region, and a C-terminal coiled-coil domain.⁽⁴⁴⁾ In fibroblasts from these patients, the *NEK9* mutation results in a number of cell-cycle defects including reduced proliferation capability and delayed cell-cycle progression through the G1/S boundary and S-phase. Analysis of *NEK9* in patient fibroblasts also showed reduced cilia number and length and in *Caenorhabditis elegans* the *NEK9* orthologue (*NEKL-1*) expression was restricted exclusively to a subset of ciliated cells, suggesting that NEK9 is involved in ciliary function.⁽⁴⁴⁾ Previous studies have shown the importance of primary cilia in osteoblast mechanosensing and function to enhance mineralization,^(45,46) making this a potential mechanism through which CLIC3 interacts with NEK9; however, our CLIC3 and NEK9 immunolabelings do not appear to be specific to the primary cilia. *NEK9* is one of 11 human NIMA-related kinase (NEK) genes that encode serine-threonine kinases with diverse biological roles including cell-cycle control, cilia regulation, and DNA damage sensing and repair,⁽⁴⁷⁻⁵⁰⁾ and NEK9, specifically, is known to be involved in regulating spindle organization, chromosome alignment, cytokinesis, and normal cell-cycle progression.^(51,52) Mammalian Nek9 binds γ -tubulin and localizes to the centrosomes and spindle poles during early cell division, functioning in the microtubule organizing center during mitosis.^(50,53) Our immunofluorescence results demonstrate that both CLIC3 and NEK9 are most abundant surrounding the nucleus, suggesting that CLIC3 and NEK9 are associated with the microtubule-organizing center. It has been reported that

CLIC3 and other CLIC proteins associate with the cytoskeleton or scaffolding proteins for endosome trafficking.^(20–22) In addition, recent work in our laboratory has shown that inhibition of microtubules leading to dramatic cytoskeletal modifications and an increase in focal adhesions and BMP2 activity is sufficient to cause osteogenic differentiation of hMSCs,⁽²⁸⁾ and osteogenic differentiation is associated with significant changes in cytoskeletal protein in human osteoblasts.⁽²⁷⁾ In addition, our bioinformatics analysis of CLIC3-His pulled-down proteins identified an overrepresentation of proteins involved in processes related to cytoskeletal and integrin signaling. Collectively, we suggest that NEK9 and CLIC3 may regulate osteoblast differentiation via cytoskeleton-associated signaling processes, but this requires further investigation.

Our results show that CLIC3 and PSS1 associate in differentiating hMSCs and that knockdown of *PTDSS1* significantly reduces osteoblast differentiation. *PTDSS1* encodes phosphatidylserine synthase 1 (PSS1) that, along with phosphatidylserine synthase 2 (PSS2), promotes the biosynthesis of phosphatidylserine (PS) by the exchange of L-serine with the choline moiety of phosphatidylcholine.⁽⁵⁴⁾ In 2014⁽⁵⁵⁾ it was discovered that de novo missense mutations in the *PTDSS1* gene are responsible for Lenz-Majewski hyperostotic dwarfism (LMHD), an extremely rare condition characterized by sclerosing bone dysplasia, intellectual disability, and distinct craniofacial, dental, cutaneous, and distal limb anomalies (OMIM #151050),^(56,57) which was affirmed by Whyte and colleagues,⁽⁵⁸⁾ reporting additional LMHD patients harboring mutations in *PTDSS1*. These *PTDSS1* mutations result in a gain of function phenotype leading to increased synthesis of phosphatidylserine (PS).^(55,58) Bone turnover markers in a female LMHD patient, who was confirmed to have a mutation in *PTDSS1*, revealed that her osteosclerosis was a result of accelerated bone formation along with unremarkable rates of bone resorption.⁽⁵⁸⁾ They also found elevated levels of phosphoserine in their LMHD patients' urine, which could be indicative of increased PS biosynthesis. PS has unique physical and biochemical properties that lead to its physiological importance in roles related to apoptosis, coagulation, the internalization of viruses, and Ras/Rho and protein kinase C signaling.^(54,59,60) CLIC proteins have also been shown to be heavily involved in Rho signaling in a number of cell types,^(18,42,53) providing a potential functional link between the two proteins. In bone PS is known to bind calcium within matrix vesicles leading to HA crystal formation,⁽⁶¹⁾ and enhance osteogenic differentiation of MSCs⁽⁶²⁾ and human osteoblast progenitor cells⁽⁶³⁾ to promote bone formation. In addition, PS-containing liposomes were shown to inhibit osteoclast differentiation and prevent trabecular bone loss,⁽⁶⁴⁾ making PS a potential candidate for osteoporosis treatment. Interestingly, in cultured A2780 ovarian carcinoma cells and in pancreatic and ovarian tumors that contain elevated levels of CLIC3, CLIC3 functions to mediate the return of $\alpha 5\beta 1$ from late endosomes/lysosomes to the plasma membrane.⁽²⁰⁾ We hypothesize that CLIC3 may act similarly in osteoblasts to direct the transport of PS to the plasma membrane or to matrix vesicles. These previous findings, taken together with our current work, supports the idea that CLIC3 works together with PSS1 to increase biosynthesis of PS in osteoblast differentiation and bone formation.

In conclusion, we have successfully identified *CLIC3* to be a novel gene modulating osteoblast differentiation and enhancing bone formation. Although further studies are required to determine the molecular mechanisms by which CLIC3

modulates mineralization, we postulate that CLIC3 and NEK9 both play roles in the microtubule organizing center to induce cytoskeletal changes important for osteoblast differentiation, and that CLIC3 interacts with PSS1 to enhance PS synthesis and PS translocation to the plasma membrane where it plays an important role in matrix vesicle-mediated HA formation and mineralization of the bone. The specificity of *CLIC3* to promote the osteogenic lineage while its expression decreases during adipocyte differentiation, in combination with the importance of CLIC3 during human osteoblast differentiation, could make it a potential target for future bone anabolic treatments.

Disclosures

All authors state that they have no conflicts of interest.

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Authors' roles: Study design: AB, ME, JP, JL, and BE. Study conduct: AB, CL, MSK, JV, MJ, and DD. Data collection: AB, CL, MSK, JV, MJ, and DD. Data analysis: AB and CL. Data interpretation: AB, CL, JP, JD, JL, and BE. Drafting manuscript: AB. Revising manuscript content: AB, CL, MSK, JP, JL, and BE. Approving final version of manuscript: CL, MSK, JD, ME, JP, JL, and BE. AB and BE take responsibility for the integrity of the data analysis.

References

1. Sambrook P, Cooper C. Osteoporosis. *Lancet*. 2006 Jun 17;367(9527):2010–8.
2. Matsuo K, Irie N. Osteoclast-osteoblast communication. *Arch Biochem Biophys*. 2008 May 15;473(2):201–9.
3. Chamberlain G, Fox J, Ashton B, Middleton J. Concise review: mesenchymal stem cells: their phenotype, differentiation capacity, immunological features, and potential for homing. *Stem Cells*. 2007 Nov;25(11):2739–49.
4. Ducy P, Zhang R, Geoffroy V, Ridall AL, Karsenty G. *Osf2/Cbfa1*: a transcriptional activator of osteoblast differentiation. *Cell*. 1997 May 30;89(5):747–54.
5. Komori T, Yagi H, Nomura S, et al. Targeted disruption of *Cbfa1* results in a complete lack of bone formation owing to maturational arrest of osteoblasts. *Cell*. 1997 May 30;89(5):755–64.
6. Otto F, Thornell AP, Crompton T, et al. *Cbfa1*, a candidate gene for cleidocranial dysplasia syndrome, is essential for osteoblast differentiation and bone development. *Cell*. 1997 May 30;89(5):765–71.
7. Aubin JE. Regulation of osteoblast formation and function. *Rev Endocr Metab Disord*. 2001 Jan;2(1):81–94.
8. Eijken M, Koedam M, van Driel M, Buurman CJ, Pols HAP, van Leeuwen JPTM. The essential role of glucocorticoids for proper human osteoblast differentiation and matrix mineralization. *Mol Cell Endocrinol*. 2006 Mar 27;248(1–2):87–93.
9. Iba K, Chiba H, Sawada N, Hirota S, Ishii S, Mori M. Glucocorticoids induce mineralization coupled with bone protein expression without influence on growth of a human osteoblastic cell line. *Cell Struct Funct*. 1995 Oct;20(5):319–30.
10. Perrien DS, Akel NS, Dupont-Versteegden EE, et al. Aging alters the skeletal response to disuse in the rat. *Am J Physiol Regul Integr Comp Physiol*. 2007;292:R988–96.
11. Singh L, Brennan TA, Russell E, et al. Aging alters bone-fat reciprocity by shifting in vivo mesenchymal precursor cell fate towards an adipogenic lineage. *Bone*. 2016 Jan 19;85:29–36.

12. Moerman EJ, Teng K, Lipschitz DA, Lecka-Czernik B. Aging activates adipogenic and suppresses osteogenic programs in mesenchymal marrow stroma/stem cells: the role of PPAR- γ 2 transcription factor and TGF- β /BMP signaling pathways. *Aging Cell*. 2004 Oct 20;3(6):379–89.
13. Kajkenova O, Lecka-Czernik B, Gubrij I, et al. Increased adipogenesis and myelopoiesis in the bone marrow of SAMP6, a murine model of defective osteoblastogenesis and low turnover osteopenia. *J Bone Miner Res*. 1997 Nov;12(11):1772–9.
14. Nuttall ME, Gimble JM. Is there a therapeutic opportunity to either prevent or treat osteopenic disorders by inhibiting marrow adipogenesis? *Bone*. 2000 Aug;27(2):177–84.
15. Landry D, Sullivan S, Nicolaides M, et al. Molecular cloning and characterization of p64, a chloride channel protein from kidney microsomes. *J Biol Chem*. 1993 Jul 15;268(20):14948–55.
16. Landry DW, Akabas MH, Redhead C, Edelman A, Cragoe EJ, Al-Awqati Q. Purification and reconstitution of chloride channels from kidney and trachea. *Science*. 1989 Jun 23;244(4911):1469–72.
17. Singh H. Two decades with dimorphic chloride intracellular channels (CLICs). *FEBS Lett*. 2010 May 17;584(10):2112–21.
18. Jiang L, Phang JM, Yu J, et al. CLIC proteins, ezrin, radixin, moesin and the coupling of membranes to the actin cytoskeleton: a smoking gun? *Biochim Biophys Acta*. 2014 Feb;1838(2):643–57.
19. Littler DR, Harrop SJ, Goodchild SC, et al. The enigma of the CLIC proteins: ion channels, redox proteins, enzymes, scaffolding proteins? *FEBS Lett*. 2010 May 17;584(10):2093–101.
20. Dozynkiewicz MA, Jamieson NB, Macpherson I, et al. Rab25 and CLIC3 collaborate to promote integrin recycling from late endosomes/lysosomes and drive cancer progression. *Dev Cell*. 2012;22(1):131–45.
21. Knowles LM, Zewe J, Malik G, Parwani AV, Gingrich JR, Pilch J. CLT1 targets bladder cancer through integrin α 5 β 1 and CLIC3. *Mol Cancer Res*. 2013 Feb;11(2):194–203.
22. Macpherson IR, Rainero E, Mitchell LE, et al. CLIC3 controls recycling of late endosomal MT1-MMP and dictates invasion and metastasis in breast cancer. *J Cell Sci*. 2014;127(Pt 18):3893–901.
23. Qian Z, Okuhara D, Abe MK, Rosner MR. Molecular cloning and characterization of a mitogen-activated protein kinase-associated intracellular chloride channel. *J Biol Chem*. 1999 Jan 15;274(3):1621–7.
24. Bruedigam C, van Driel M, Koedam M, et al. Basic techniques in human mesenchymal stem cell cultures: differentiation into osteogenic and adipogenic lineages, genetic perturbations, and phenotypic analyses. *Curr Protoc Stem Cell Biol*. 2011 Jun;Chapter 1: Unit1H.3.
25. van der Eerden BCJ, Weissgerber P, Fratzi-Zelman N, et al. The transient receptor potential channel TRPV6 is dynamically expressed in bone cells but is not crucial for bone mineralization in mice. *J Cell Physiol*. 2012;227(5):1951–9.
26. Drabek K, van de Peppel J, Eijken M, van Leeuwen JPTM. GPM6B regulates osteoblast function and induction of mineralization by controlling cytoskeleton and matrix vesicle release. *J Bone Miner Res*. 2011 Sep;26(9):2045–51.
27. Alves RDAM, Eijken M, Swagemakers S, et al. Proteomic analysis of human osteoblastic cells: relevant proteins and functional categories for differentiation. *J Proteome Res*. 2010 Sep 3;9(9):4688–700.
28. Brum AM, van de Peppel J, van der Leijde CS, et al. Connectivity Map-based discovery of parabendazole reveals targetable human osteogenic pathway. *Proc Natl Acad Sci U S A*. 2015 Oct 13;112(41):12711–6.
29. Abdallah BM, Ditzel N, Kassem M. Assessment of bone formation capacity using in vivo transplantation assays: procedure and tissue analysis. In: Westendorf J, editor. *Osteoporosis: methods and protocols*. New York: Humana Press; 2008. p. 89–100. *Methods in Molecular Biology (Book 455)*
30. Morhayim J, van de Peppel J, Demmers JAA, et al. Proteomic signatures of extracellular vesicles secreted by nonmineralizing and mineralizing human osteoblasts and stimulation of tumor cell growth. *FASEB J*. 2015 Jan 1;29(1):274–85.
31. Cox J, Matic I, Hilger M, et al. A practical guide to the MaxQuant computational platform for SILAC-based quantitative proteomics. *Nat Protoc*. 2009 Jan;4(5):698–705.
32. Cox J, Neuhauser N, Michalski A, Scheltema RA, Olsen JV, Mann M. Andromeda: a peptide search engine integrated into the MaxQuant environment. *J Proteome Res*. 2011 Apr 1;10(4):1794–805.
33. Ulmasov B, Bruno J, Gordon N, Hartnett ME, Edwards JC. Chloride intracellular channel protein-4 functions in angiogenesis by supporting acidification of vacuoles along the intracellular tubulogenic pathway. *Am J Pathol*. 2009 Mar;174(3):1084–96.
34. Malik M, Jividen K, Padmakumar VC, et al. Inducible NOS-induced chloride intracellular channel 4 (CLIC4) nuclear translocation regulates macrophage deactivation. *Proc Natl Acad Sci U S A*. 2012 Apr 17;109(16):6130–5.
35. Fernández-Salas E, Suh KS, Speransky V V, et al. mtCLIC/CLIC4, an organellar chloride channel protein, is increased by DNA damage and participates in the apoptotic response to p53. *Mol Cell Biol*. 2002 Jun;22(11):3610–20.
36. Berry KL, Bülow HE, Hall DH, Hobert O. A *C. elegans* CLIC-like protein required for intracellular tube formation and maintenance. *Science*. 2003 Dec 19;302(5653):2134–7.
37. Edwards JC, Cohen C, Xu W, Schlesinger PH. c-Src control of chloride channel support for osteoclast HCl transport and bone resorption. *J Biol Chem*. 2006 Sep 22;281(38):28011–22.
38. Suh KS, Mutoh M, Nagashima K, et al. The organellar chloride channel protein CLIC4/mtCLIC translocates to the nucleus in response to cellular stress and accelerates apoptosis. *J Biol Chem*. 2004 Feb 6;279(6):4632–41.
39. Xu Y, Kang J, Yuan Z, et al. Suppression of CLIC4/mtCLIC enhances hydrogen peroxide-induced apoptosis in C6 glioma cells. *Oncol Rep*. 2013 Apr;29(4):1483–91.
40. Jiang L, Salao K, Li H, et al. Intracellular chloride channel protein CLIC1 regulates macrophage function through modulation of phagosomal acidification. *J Cell Sci*. 2012 Nov 15;125(Pt 22):5479–88.
41. Dulhunty A, Gage P, Curtis S, Chelvanayagam G, Board P. The glutathione transferase structural family includes a nuclear chloride channel and a ryanodine receptor calcium release channel modulator. *J Biol Chem*. 2001 Feb 2;276(5):3319–23.
42. Ponsioen B, van Zeijl L, Langeslag M, et al. Spatiotemporal regulation of chloride intracellular channel protein CLIC4 by RhoA. *Mol Biol Cell*. 2009 Nov;20(22):4664–72.
43. Yang J-Y, Jung JY, Cho SW, et al. Chloride intracellular channel 1 regulates osteoblast differentiation. *Bone*. 2009;45(6):1175–85.
44. Casey JP, Brennan K, Scheidel N, et al. Recessive NEK9 mutation causes a lethal skeletal dysplasia with evidence of cell cycle and ciliary defects. *Hum Mol Genet*. 2016 May 1;25(9):1824–35.
45. Delaine-Smith RM, Sittichokechaiwut A, Reilly GC. Primary cilia respond to fluid shear stress and mediate flow-induced calcium deposition in osteoblasts. *FASEB J*. 2014 Jan;28(1):430–9.
46. Chen JC, Hoey DA, Chua M, Bellon R, Jacobs CR. Mechanical signals promote osteogenic fate through a primary cilia-mediated mechanism. *FASEB J*. 2016 Apr;30(4):1504–11.
47. Chen Y, Chen C-F, Riley DJ, Chen P-L. Nek1 kinase functions in DNA damage response and checkpoint control through a pathway independent of ATM and ATR. *Cell Cycle*. 2011 Feb 15;10(4):655–63.
48. O'Regan L, Blot J, Fry AM. Mitotic regulation by NIMA-related kinases. *Cell Div*. 2007;2:25.
49. Moniz L, Dutt P, Haider N, Stambolic V. Nek family of kinases in cell cycle, checkpoint control and cancer. *Cell Div*. 2011 Jan;6:18.
50. Roig J, Groen A, Caldwell J, Avruch J. Active Nerc1 protein kinase concentrates at centrosomes early in mitosis and is necessary for proper spindle assembly. *Mol Biol Cell*. 2005;16(10):4827–40.
51. Fry AM, O'Regan L, Sabir SR, Bayliss R. Cell cycle regulation by the NEK family of protein kinases. *J Cell Sci*. 2012 Oct 1;125(Pt 19):4423–33.
52. Wang Y-K, Yu X, Cohen DM, et al. Bone morphogenetic protein-2-induced signaling and osteogenesis is regulated by cell shape, RhoA/ROCK, and cytoskeletal tension. *Stem Cells Dev*. 2012;21(7):1176–86.
53. Yang S-W, Gao C, Chen L, et al. Nek9 regulates spindle organization and cell cycle progression during mouse oocyte meiosis and its

- location in early embryo mitosis. *Cell Cycle*. 2012 Dec 1;11(23):4366–77.
54. Tomohiro S, Kawaguti A, Kawabe Y, Kitada S, Kuge O. Purification and characterization of human phosphatidylserine synthases 1 and 2. *Biochem J*. 2009 Mar 1;418(2):421–9.
55. Sousa SB, Jenkins D, Chanudet E, et al. Gain-of-function mutations in the phosphatidylserine synthase 1 (PTDSS1) gene cause Lenz-Majewski syndrome. *Nat Genet*. 2014 Jan;46(1):70–6.
56. Lenz WD, Majewski F. A generalized disorders of the connective tissues with progeria, choanal atresia, symphalangism, hypoplasia of dentine and craniodiaphyseal hypostosis. *Birth Defects Orig Artic Ser*. 1974 Jan;10(12):133–6.
57. Robinow M, Johanson AJ, Smith TH. The Lenz-Majewski hyperostotic dwarfism. A syndrome of multiple congenital anomalies, mental retardation, and progressive skeletal sclerosis. *J Pediatr*. 1977 Sep;91(3):417–21.
58. Whyte MP, Blythe A, McAlister WH, Nenninger AR, Bijanki VN, Mumm S. Lenz-Majewski hyperostotic dwarfism with hyperphosphoserinuria from a novel mutation in PTDSS1 encoding phosphatidylserine synthase 1. *J Bone Miner Res*. 2015 Apr;30(4):606–14.
59. Schick PK, Kurica KB, Chacko GK. Location of phosphatidylethanolamine and phosphatidylserine in the human platelet plasma membrane. *J Clin Invest*. 1976 May;57(5):1221–6.
60. Leventis PA, Grinstein S. The distribution and function of phosphatidylserine in cellular membranes. *Annu Rev Biophys*. 2010 Jan;39:407–27.
61. Wu LNY, Genge BR, Wuthier RE. Analysis and molecular modeling of the formation, structure, and activity of the phosphatidylserine-calcium-phosphate complex associated with biomineralization. *J Biol Chem*. 2008 Feb 15;283(7):3827–38.
62. Xu C, Zheng Z, Fang L, et al. Phosphatidylserine enhances osteogenic differentiation in human mesenchymal stem cells via ERK signal pathways. *Mater Sci Eng C Mater Biol Appl*. 2013 Apr 1;33(3):1783–8.
63. Satsangi A, Satsangi N, Glover R, Satsangi RK, Ong JL. Osteoblast response to phospholipid modified titanium surface. *Biomaterials*. 2003 Nov;24(25):4585–9.
64. Wu Z, Ma HM, Kukita T, Nakanishi Y, Nakanishi H. Phosphatidylserine-containing liposomes inhibit the differentiation of osteoclasts and trabecular bone loss. *J Immunol*. 2010 Mar 15;184(6):3191–201.