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Functional study of promoter gene polymorphisms of sclerostin

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Date of receipt: 16/07/2016
Date of acceptance: 25/09/2016

Work rewarded with the scholarship of Molecular Bone Biology FEIOMM 2013.

Summary
Sclerostin, encoded by the SOST gene, inhibits the Wnt pathway and, consequently, tends to decrease bone mass. Some polymorphisms of the SOST promoter have been associated with bone mineral density (BMD), but the molecular mechanisms involved are unknown. The aim of this study was to study the functional role of one polymorphism in vitro. We cloned the proximal promoter region of SOST gene, containing different alleles at the rs851054 SNP, in luciferase reporter vectors and transfected them into the cell lines HEK-293T, SAOS-2 and HOS-TE85. We did not find significant differences in the transcriptional activity of vectors with either the A or the G allele of the SNP. The co-transfection of vectors expressing RUNX2 and OSX markedly increased the transcriptional activity of the SOST promoter constructs (A allele, 2.5±0.9 fold, p<0.05; G allele, 1.9±0.8 fold, p<0.05), without significant differences between the rs851054 alleles. Moreover, no allele differences were detected in EMSAs.

In conclusion, the DNA region upstream of the TSS of the SOST gene has a strong promoter activity that is enhanced by RUNX2 and OSX. Frequent allelic variants in this region have been associated with BMD, but the mechanisms involved remain to be elucidated because no functional differences between alleles were detected in vitro.

Key words: sclerostin, gene regulation, polymorphisms, transfection.
Introduction
Long-term bone mass development is determined by the balance between resorption and bone formation. So, when bone forming activity is insufficient to replace the part destroyed during resorption, it will inevitably lead to decreased bone tissue, characteristic of osteoporosis. Osteoblasts, responsible for bone formation, derived from mesenchymal stem cells, can also lead to other cell types such as chondrocytes or adipocytes. The proliferation and differentiation of osteoblast precursors are controlled by different intracrine, paracrine and endocrine factors. Thus, some transcription factors, such as RUNX2 and Osterix (OSX) are considered essential in the early stages of osteoblast differentiation. The Wnt pathway also plays an important role. Wnt ligands join the cell membrane receptors together and, through various intracellular mediators, modify the expression of target genes that generally promote bone formation and increasing osteoprotegerin (OPG) expression, which secondarily is a decrease in resorption. Wnt receptor ligands are molecular complexes which include at least two proteins; Frizzled and LRP, of which there are several forms. As with many other regulatory systems, there are also inhibitors of the Wnt pathway. One of the most widely studied is the sclerostin, encoded by the SOST gene. This gene is expressed preferentially in the osteocytes. Being an inhibitor of the Wnt pathway, sclerostin has a negative influence on bone mass. Its role in skeletal biology seems important. Indeed, neutralizing sclerostin antibodies exert a potent anabolic effect on the skeleton, and the SOST gene inactivating mutations cause an exaggerated increase in bone mass.

In line with the biological role of sclerostin, allelic variants of the SOST gene appear to influence bone mass. Thus, in some studies genome scan (GWAS) and other genetic association studies, we found a link between some common SOST gene polymorphisms and bone mineral density (BMD). Our group also found an association between some polymorphisms located in the promoter region of SOST and BMD in postmenopausal women. The aim of this study was to explore the regulatory capacity of one of these polymorphisms with functional in vitro studies. We used vectors in which the promoter of SOST is inserted in front of a reporter gene encoding a protein (luciferase) whose activity can be easily measured. Experiments were carried out to test the binding capacity of nuclear proteins to these regions assessing delay electrophoretic mobility.

Material and methods
Construction of reporter vectors
SOST promoter (1-1440) was cloned from genomic DNA of two individuals with known genotype homozygous for alleles A and G, respectively, of the rs851054 polymorphism (Figure 1). Both fragments were otherwise identical and equal to the reference of the human genome sequences. DNA extraction occurred after informed consent, within a study of genetic factors involved in osteoporosis, authorized by the Ethics Committee on Clinical Research. For amplification of the corresponding genomic fragments, PCR primers designed with the Primer3 program were used which included sequences for the restriction enzymes Smal and Xh01, to facilitate further cloning. PGL2 vector and ampiclon were cut with these enzymes and the fragments were ligated using T4 DNA liga-

Transfection and analysis of transcriptional activity
HEK-293T cells (a line derived from human kidney) and human osteoblastic lines Saos-2 and HOS-TE85 were transfected with vectors containing the promoter sequences of SOST and a RSV-βGAL control vector, constitutively expressing the LacZ gene, encoding β-galactosidase, in order to normalize the results which depend on transfection efficiency. As a negative control and parallel transfection with empty vector was also performed. For transfection HEK-293T cells 125,000 (50,000 or Saos-2 or HOS-TE85) were seeded into each well of a 24 well plate. After reaching 80% confluency were transfected 500ng vector using Lipofectamine 3000 according to the manufacturer's recommendations (Invitrogen). At 48 hours, the medium was aspirated and the cells with 70 mu1 buffer, after which galactosidase activity (Galacto-Light Plus™ β-Galactosidase Reporter Gene Assay System, Applied Biosystems) was measured and luciferase (Luciferase lysed Assay System, Promega) by luminometry. The co-transfection with expression vectors and OSX5 RUNX2 was performed following a similar procedure, but ensuring that the total amount of exogenous DNA to be transfected was held constant in all wells trays.

From each transfection, double and triple techniques were performed. Furthermore, each experiment was repeated at least three different times to obtain biological triplicates. The results were expressed as the ratio between luciferase and galactosidase activity in cell lysates.

Analysis of nuclear protein binding
Protein extracts were obtained from 50 x 106 HOS-TE85 cells. To do this, in a lysis buffer was used containing protease inhibitors (50 mM KCl,
0.5% NP-40, 25 mM HEPES, 1.5 pM leupeptin, aprotinin 46.88 µM, 125 µM DTT, 1 mM PMSF). These were centrifuged for 1 minute at 10,000 rpm at 4°C. After washing, the cells were re-suspended in 100 µl of extraction buffer (500 mM KCl, 25 mM HEPES, 10% glycerol, 1.5 pM leupeptin, aprotinin 46.88 µM, 125 µM DTT, 1 mM PMSF) and centrifuged for 5 minutes at 14,000 rpm at 4°C. The supernatant was recovered, the protein concentration was quantified and adjusted to 5 µg/µl.

To test the delay of electrophoretic mobility shift assay (EMSA), oligonucleotide pairs were used, including IR dye labeling 700 in the 5' end of the "forward" component of each pair (Biolegio). The sequences were as follows:

rs851054 allele A Fwd: AACAGAAACACCTTGGGCCATGGCCCA
rs851054 allele A Rev: TGGCCCAAGGCGTTTCTGTT
rs851054 allele G Fwd: AACAGAAACGCCTTGGGCCATGGCCCA
rs851054 allele G Rev: TGGCCCAAGGCGTTTCTGTT

Prefabricated gels 6% polyacrylamide (Invitrogen) and the Odyssey Infrared EMSA kit were used for retardation. After annealing, the oligonucleotides were incubated with protein extract for 20 minutes, loaded on the gel and were run at 70 V for 60 minutes, after which the images of the gels were captured. In these experiments, if the probes bind nuclear proteins occurs a delay in electrophoretic mobility compared to that experienced by the isolated probe migration.

**Results**

Constructions SOST promoter sequence showed high transcription activating capacity by increasing levels of luciferase expression up to 1,000 times compared with empty vectors. The activity was apparently higher in HEK-293T cells than in the other lines (Figure 2).

However, no significant differences between the two alleles SOST promoter in any of the cell lines tested. In fact, the ratio between the transcriptional activity of alleles A and G (Ratios A/G) in Saos-2 and HOS TE85-293T cells were 1.3±0.7, 1.0±0.4 and 0.8±0.7, respectively. None of the three was significantly different from the unit (Figure 2).

The co-transfection of vectors expressing constitutively RUNX2 and OSX increased transcriptional activity of the SOST promoter (on average 2.5±0.9 times above baseline for allele A and 1.9±0.8 times to the G allele; both p<0.05). This increase was also be independent of which allele was present in the SOST promoter (Figure 3).

Furthermore, in experiments delay electrophoretic mobility see that the region where the polymorphism lies was able to bind nuclear proteins, presumably with regulatory function without us to observe differences between alleles (Figure 4). Therefore, no further studies were performed to identify the nature of these proteins.

**Discussion**

The role of sclerostin in bone biology is undeniable, as revealed by the sharp BMD increase observed after administration of romosozumab, a monoclonal antibody that blocks sclerostin action, and excessive bone mass experienced by patients with sclerosteosis or Van Buchem syndrome or carriers of rare mutations that induce a rare disease such as loss of SOST gene function14. Although the evidence is limited, several studies suggest that frequent allelic variants of the gene can also influence, to a lesser extent, BMD and osteoporotic fracture risk. In fact, in a study of genomic sweep with a large sample size of rs479299 polymorphism association was found with BMD15. This polymorphism is located in an intergenic region, being the closest SOST gene, located at about 40 kb. On the other hand, several groups, including ours, have analyzed the relationship of some common polymorphisms in the promoter region of SOST with BMD. So, we found a significant association of rs851054 and rs851056 polymorphisms, only separated by 560 bp and members of the same haplotype block with BMD of the spine in women12. Some authors have confirmed the association of the polymorphism in the SOST promoter with BMD in other population groups16-20. On the other hand, in a recent study involving a small group of patients, an association of rs851054 alleles with the expression of SOST in bone tissue was found21.

Figure 1. Schematic of the promoter region of SOST gene and location of some common polymorphisms (minor allele frequency greater than 5%)
The main objective of this study was to explore the potential functional impact of one of these polymorphisms and to analyze their influence on the transcriptional activity of the SOST gene promoter. Transfection experiments with reporter vectors and delay in electrophoretic mobility did not reveal differences between the allelic variants studied. There are several limitations of the models used and numerous reasons that could explain these negative results. First, it could be argued that the in vitro model does not adequately reflect the in vivo situation. For example, it cannot be excluded that the regulatory activity requires genomic regions with activity enhancer, promoter and far not included therefore in the region cloned into the vectors. On the other hand, at least theoretically it might be thought that the difference in allele activity is only expressed in response to regulatory factors in certain cell types such as osteocytes, but are absent in the cell types used in our experiments. Second, it appears that the association with BMD might rely on other common polymorphisms in linkage disequilibrium with those studied here. If so, they should be in distant, remote regions of the promoter, since in this region there is a strong linkage. In this regard, it should be noted that there is a regulatory region of SOST located at about 50 kb, containing some polymorphisms linked inconsistently with BMD. This region is capable of fixing MEF2C transcription factor and plays an important role in regulating SOST because their lack causes the Van Buchem disease phenotype in humans and increased bone mass in murine models. A third possibility is that the association with BMD depends on some low frequency polymorphisms located in the promoter region. Finally, polymorphisms might act through epigenetic mediators that cannot be adequately recapitulated in in vitro models. Further studies are thus required including the systematic analysis of the variants of the 5' and 3' of the gene and others which are associated with considering the tridimensional structure of chromatin and mutagenesis studies in vivo to clarify the mechanisms by which these allelic variants may modulate the activity of sclerostin and associate with differences in bone mass.

Our study confirmed previous results indicating that RUNX2 and OSX increase the expression of SOST in humans. This indicates that these transcription factors have a complex role in the osteoblastic line. On the one hand, they have a well-established role as determinants of the early stages of differentiation of mesenchymal cells into osteoblasts and enable an adequate number of bone-forming osteoblasts. They also promote SOST promoter activity. Therefore, in cells capable of expressing this gene, such as osteocytes, the secretion of sclerostin could promote and thus help prevent exaggerated bone formation. In any case, the SOST promoter variants do not appear to influence the response to these factors.

In conclusion, in this study we have confirmed that the region located before the start of translation of the gene SOST has potent promoter activity, which also induced transcription factors RUNX2 and OSX. Frequent variants of this region have been associated with bone mass, but the mechanisms involved are still unknown, since the alleles show no differences in in vitro transcriptional activity.

Acknowledgements: This research was carried out with the support of a grant from the FEIOMM 2013 "Interaction between Osterix, Runx2 and sclerostin: molecular mechanisms and impact on bone mass" and the Institute of Health Carlos III (PI12-615), co-financed with the European Union’s FEDER funds. The authors would also like to express their grateful to Dr. Svante Paabo for generously supplying Runx2 expression plasmid.

Figure 2. Transcriptional activity of the promoter sequence of SOST after transfection into osteoblast-like cells (HOS-TE85 OSAS) and HEK-293T line. In the left slide all the experiments (black bars, vectors with promoter sequences SOST white bars, empty vector) are represented; in the right, the ratio of the activity of the constructs with the A and G alleles in each of the experiments. the average values of 5-6 independent experiments performed in duplicate or triplicate are shown. The vertical lines indicate the standard error.
Declaration of interest: The authors declare no conflicts of interest.

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Figure 3. Comparison of the transcriptional activity of the vectors A and G alleles in HEK-293T cells in basal conditions and after co-transfection of expression vectors Runx2 and OSX

Figure 4. Test of delay in electrophoretic mobility. The onset of delay band is observed when the nuclear protein extract, whose intensity decreases by adding unlabeled probe is added. There is some difference in band intensity between alleles A and G, parallel to the signal strength of the undelayed probe. However, in several experiments no consistent difference between the allele probe and the probe was observed with the G allele

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Allele A Allele G