Preaxial Polydactyly in Sost/Sostdc1 Double Knockouts

C. M. Yee, N. M. Collette, G. G. Loots

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Preaxial polydactyly in Sost/Sostdc1 double knockouts

Cristal Yee
University of California, Merced

Gabriela G. Loots
Ph.D., Molecular and Cell Biology
Biomedical Scientists at LLNL

Lawrence Livermore National Laboratory, Livermore, CA

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Abstract

In the United States, ~5% are born with congenital birth defects due to abnormal function of cellular processes and interactions. Sclerosteosis, a rare autosomal recessive disease, causes hyperostosis of the axial and appendicular skeleton, and patients present radial deviation, digit syndactyly, nail dysplasia, and overall high bone mineral density. Sclerosteosis is due to a loss of function of sclerostin (Sost). Sost is a Wnt (Abbrev.) antagonist; when mutated, nonfunctional Sost results in hyperactive osteoblast activity which leads to abnormal high bone mass. Previous studies have shown that Sost overexpression in transgenic mice causes reduced bone mineral density and a variety of limb phenotypes ranging from lost, fused, and split phalanges. Consistent with clinical manifestations of Sclerosteosis, Sost knockout mice exhibit increased generalized bone mineral density and syndactyly of the digits. Sostdc1 is a paralog of Sost that has also been described as an antagonist of Wnt signaling, in developing tooth buds. Unlike Sost knockouts, Sostdc1 null mice do not display any limb abnormalities. To determine if Sost and Sostdc1 have redundant functions during limb patterning, we examined Sost\textsuperscript{-/-}; Sostdc1\textsuperscript{-/-} mice determined that they exhibit a novel preaxial polydactyly phenotype with a low penetrance. LacZ staining, skeletal preparations, and \textit{in situ} hybridization experiments were used to help characterize this novel phenotype and understand how this phenotype develops. We find Sost and Sostdc1 to have complementary expression patterns during limb development, and the loss of their expression alters the transcription of several key limb regulators, such as Fgf8, Shh and Grem.

Introduction

Sclerosteosis is a rare autosomal recessive disease that causes hyperostosis to the axial skeleton and appendicular skeleton. Patients with Sclerosteosis appear to have radial deviation, hand malformations (commonly syndactyly of digit 2 and 3), nail dysplasia, overall high bone density, and secondary neurological defects (Fig1A, B, C). Some of these secondary neurological defects, such as hearing loss and constriction of the optic nerve, can escalate to sudden death caused by increased intracranial pressure. This rare hyperostosis disease is due to loss of sclerostin (SOST), which causes hyperactive osteoblast activity.

Sost has been initially described as a BMP (Bone Morphogenic Protein) antagonist, but in the recent years Sost has emerged as a Wnt antagonist (Collette et al., 2010). Altering the function of Sost results in increased or decreased Wnt signaling. For example, using Sost transgenic mice causes osteopenia and reduced bone mineral density (BMD) in both the appendicular and axial skeleton (Loots et al., 2005) due to reduced Wnt signaling in bone. Furthermore, these transgenic mice show a variety of phenotypes ranging from fused, split, to missing phalanges due to dosage-dependent overexpression of Sost (Collette et al., 2010). Alternatively, knocking out Sost results in a growth increase of bone formation and digit syndactyly phenotype (Fig1).

Sostdc1 (Sost domain-containing protein 1; aka Sostl, USAG-1, Wise, ectodin) is a paralog of Sost and an antagonist of the LRP receptor in tooth buds (Kasai et al., 2005). Sost and Sostdc1 share 40% identity at the DNA sequence level and a 55% protein similarity and initial studies have reported Sostdc1 to influence teeth. In one instance,
Sostdc1 knockout mice generated excessive numbers of incisors as well as affected Fgf and Shh signaling in the developing tooth bud (Kassai et al., 2005) (Fig2). Since Sostdc1 is paralogous to Sost and affects the Fgf and Shh signaling when altered, these two genes may have similar functional roles during limb development. We have examined Sost^−/−; Sostdc1^−/− double knockout mice and found them to exhibit a variably penetrant preaxial polydactyly phenotype, detected as early as E12.5 (Fig3). This may suggest that Sostdc1 and Sost plays similar, redundant roles, interfering with the same signaling pathway in the limb, which leads to polydactyly.

This preaxial polydactyly phenotype in the Sost^−/−; Sostdc1^−/− double knockout mice has led to questions such as: (1) where are Sost and Sostdc1 expressed? (2) are Sost and Sostdc1 interfering with the same signaling pathway? (3) how is digit 1 determined during limb development and digit formation? To further our investigation, we have designed three sets of experiments that would uncover Sostdc1 role and contribution in limb deformity. The first aim is to locate where Sost and Sostdc1 are expressed under normal developmental conditions, via LacZ staining of organs, bone segments, and forelimb buds. Locating where these two genes are will assist in determining of where and when these two genes are expressed. Second of all, we aim to compare and analyze cartilage condensation and digit formation throughout developmental time points of wildtype, Sost knockout, Sostdc1 knockout, and Sost/Sostdc1 double knockout mice via skeletal preparations. Comparing skeletal preparations presents limb development and outlines when the polydactyly phenotype begins developmentally. Finally, the third aim is to analyze expression patterns along the anterior posterior regions via in situ experiments of different markers to determine what signals are responsible for digit 1 (thumb) formation and how other genes are effected by Sost and Sostdc1 alterations.
and radial deviation (curved lines). Sost⁺⁻; Sostdc1⁺⁻ double mutants show preaxial polydactyly, or duplication of anterior digits (asterisks and inset). Skeletal preparations of adult autopods are also shown.

Figure 2. Sostdc1 is an antagonist of the LRP receptors in developing tooth bud. (A) Sostdc1⁻⁻ mice develop a fused molar phenotype. (B) A cross-section of an embryo’s cranium shows Sostdc1 being expressed in the tooth buds. (C) Sostdc1⁻⁻ mice alters the Wnt and Shh signaling, causing supernumerary incisors.

Figure 3. The polydactyly phenotype has been categorized under the standard VII classification. To the right, displays the novel preaxial polydactyly expressed in Sost⁻⁻;Sostdc1⁻⁻ mice.

Results and Discussion

Sost and Sostdc1 Expression

Our first goal was to characterize Sost and Sostdc1 expression pattern during embryonic development. The Sost⁺⁺ and Sostdc1⁺⁺ strains available in our laboratory have been generated through a knock-in strategy, where the LacZ gene has replaced the endogenous transcript, and is transcribed in frame from the endogenous promoter, such that LacZ expression faithfully recapitulates the expression of the knocked out gene. Performing LacZ staining on a cross-section of Sost⁺⁺ and Sostdc1⁺⁺ forelimb reveals adjacent, non-overlapping expression of Sost and Sostdc1 (Fig4).
The adjacent, non-overlapping expression of Sost and Sostdc1 is not only seen in the forelimbs of mice, but as well as in the ribs, jaw, kidney, and vasculature. Comparing Sost and Sostdc1 expression discloses Sost primarily sited within the bone, whereas Sostdc1 is exposed in the surrounding tissue of the bone. Since we are largely interested in the preaxial polydactyly phenotype of Sost−/−; Sostdc1−/− mice, we concentrated on the forelimbs of the mice. Using the LacZ staining method, we performed LacZ stain on whole embryos between E9.5 to E13.5 (Fig5). Sost expression appears in the developing limb starting at E9.5 as opposed to Sostdc1, which begins to emerge at E11.5. Cross sections were made of the forelimb buds to reveal that Sost is detected in the ectoderm and digit tips of the developing limb and Sostdc1 is distinguished in the limb mesenchyme. Thus, these genes in combination are being expressed in complementary domains of the limb during limb patterning.

Figure 4. LacZ staining on Sost−/− and Sostdc−/− mice displays adjacent, non-overlapping expression patterns throughout various parts of the mouse (A). Taking a cross-section of the forelimb shows Sost is expressed in osteocytes while Sostdc1 is located within the other tissues.
Preaxial Polydactyly Phenotype

Once the pattern of both Sost and Sostdc1 was determined, skeletal preparations were used to visualize when this preaxial polydactyly phenotype occurs. With the understanding that cartilage forms before bone and acting as a skeletal frame for bone development, analyzing cartilage development prior to the double thumb phenotype will give more insight as to how this unusual development emerges. Skeletal preparations with Alcian Blue cartilage staining were prepared on E14.5 embryos (Fig 6). Based on the skeletal preparations outlining cartilage patterning defects that lead to the preaxial polydactyly phenotype of Sost<sup>-/-</sup>; Sostdc1<sup>-/-</sup> mice, it is clear that shifting Sost and Sostdc1 together triggers a butterfly affect on the underlying mechanism behind normal limb development. Altering the Sost gene or Sostdc1 gene individually does not result in any visible limb phenotype. This could imply that when one gene is nonfunctional, the other ortholog gene compensates for the other. By removing both genes at once, these two nonfunctional genes create a compound affect in a form of double thumbs.

**Figure 5.** LacZ stain comparing Sost and Sostdc1 endogenous expression in the developing embryo. Genes were knocked out and replaced by a LacZ cassette to track endogenous gene expression via tissue LacZ stain. Expression in E9.5 to E13.5 whole embryos, Sost expression in top row (A’-E’), Sostdc1 expression in bottom row. Overall, expression patterns are adjacent and relatively non-overlapping. (F) LacZ stain comparing endogenous expression in the developing limb. Whole limb dorsal view, and cut sections show Sost expression confined to the ectoderm (left panels) while Sostdc1 has significant expression in the mesenchyme (right panels), positioning the combined expression domains of these genes to affect limb patterning.
Molecular Processes Underlying Limb Development

Previous research on Sost has shown that it in the limb, it primarily functions to inhibit the Wnt signaling and alters the expression of Shh, Fgf8, and Grem1 (Collette et al., 2010). Since limb development involves a complex communication between many genes, in situ methods were used to discover what normal gene expression during limb development are being effected by Sost and Sostdc1 adjustments. We hope to determine genes that Sost and Sostdc1 modulate during limb development as well as what other genes are affected. Antisense Dig-labeled probes and colorimetric detection with BM purple on E10.5 whole mount mouse embryos were used for in situ hybridization experimentations (Fig7). In situ experiments on Sost-/- displays an increase in Shh and Fgf8 expression pattern on the posterior and AER region, respectively. In comparison, Sostdc1-/- also resulted in an expansion of Shh expression, but restricted Grem1 significantly. Outcomes from our in situ experiments convey that Sost and Sostdc1 does modify expression of genes that regulate the anterior-posterior patterning of the limb. It is important to note that based on the LacZ stains, Sostdc1 appears at a later time point (E11.5) in contrast to Sost. Thus, in situ hybridizations must be performed at E11.5 to obtain a supplementary effect upon genes in the double knockouts based on Sostdc1 contribution. Additionally, E10.5 forelimb buds do not show anterior limb markers to be altered by Sost nor Sostdc1 knockouts, but most likely illustrations the onset of limb patterning. We expect to see a difference on anterior markers due to Sost and Sostdc1 knockouts at a later time frame between E11.5 and E12.5 where digit specification and separation begins.
Conclusion

In summary, Sost and Sostdc1 are paralogous genes that both function as Wnt antagonists, influencing bone and tooth formation, respectively. Our previous research has unraveled that overexpression of Sost causes osteopenia as well as loss of digit formation, while Sost knockout mice have an increase in bone mineral density. Incorporating Sost knockout with Sostdc1 knockout produces a compound axial polydactyly phenotype with a low penetrance in mice. Uncovering the mechanism behind this compound phenotype is an important step in comprehending limb development and limb defects. Knocking in LacZ gene in Sost knockout and Sostdc1 knockout mice helped to trace gene expression throughout the body. Sost and Sostdc1 have adjacent, non-overlapping expressions evident in a cross section of the forelimb, ribs, jaw, kidney, and vasculature. Whole mount LacZ stains throughout E9.5 to E13.5 exposes this non-overlapping Sost and Sostdc1 in the forelimb bud; with Sost being located in the ectoderm and digit tips and Sostdc1 being expressed in the mesoderm. Skeletal preparations depicted the axial polydactyly phenotype due to the loss of function in both Sost and Sostdc1 gene by E14.5; which contributed to our hypothesis that Sost and Sostdc1 are engaged in limb development with distinctive roles that work cohesively for proper development of the forelimb. Our last set of experiments utilizing in situ
hybridization displays Sost and Sostdc1 individually affecting other genes involved in limb development and provides additional support that Sost\(^{-/-}\); Sosdc1\(^{-/-}\) yields double thumb phenotype through a rippling signal effect. To determine Sostdc1 impact in greater details, our next goal is to look at Sostdc1 influence on gene markers at E11.5 and E12.5 as well as to examine anterior markers.

**Methods**

*Whole-mount in situ hybridization*

Whole-mount *in situ* hybridizations was carried out using standard procedures (Khokha et al., 2003). Species-specific sequences of human and mouse Sost were cloned from the 3'UTR region of each cDNA (human Sost NM_02537.2; mouse sost NM_024449.4). Briefly, digoxigenin-labeled antisense RNA probes were generated to the desired RNA sequence and hybridized to whole-mount embryos. Expression was visualized by binding BM Purple (Roche). Antisense RNA probes for *Grem1* (*MluI-SacII* fragment of NM_011824), *Fgf8* (*PstI* 3’cDNA and UTR fragment of NM_010205; see Crossley and Martin, 1995), and *Shh* (*MscI-NarI* fragment of NM_009170; see Echelard et al., 1993) were generated as described (Hogan et al., 1994) with the following modification: proteinase K digestion was omitted for ectodermal or AER probes. A minimum of 4 embryos were used per genotype, per experiment. Wild-type controls were obtained from the same litter, whenever possible.

*LacZ stains embryos*

Genetically modified mice genes were knocked out and replaced with LacZ to track Sost and Sostdc1 expression via tissue LacZ stain. LacZ stains embryos were dissected into ice-cold 1x phosphate-buffered saline (PBS), pH 7.3. Embryos were fixed in 2% paraformaldehyde, 0.2% glutaraldehyde in 1x PBS, 2mM MgCl\(_2\) at 4°C for 30 minutes to 1 hour, followed by extensive rinsing in 1x PBS, 2mM MgCl\(_2\). Embryos were stained for 4 hours (*BatGal*) at RT or overnight (Sost\(^{-/-}\)) at 4°C in X-gal stain: 1mg/ml X-gal, MgCl\(_2\), 5mM EGTA, 0.02% Nonidet P-40, 5mM potassium ferrocyanide, 5mM potassium ferricyanide, in 1x PBS, pH 7.3. After staining, embryos were postfixed in 4% paraformaldehyde in 1x PBS, pH 7.3 at 4°C, and then cleared in glycerol for photography.

*Skeletal preparations*

Skeletal preparations were made of adult mouse using Alcian Blue 8GX for cartilage and Alizarin Red S for bone following established protocols. E14.5 mouse embryos used Alcian Blue 8GX for cartilage staining only. Embryos were dissected free of internal organs, and skin, and fixed in 95% ethanol followed by stain in 0.05% Alcian Blue 8GX, 80% ethanol, 20% glacial acetic acid. Excess stain was washed out, and embryos were then stained in 60mg/L Alizarin Red S, 1% potassium hydroxide (KOH), 25% glycerol followed by clearing in 1% KOH and 25% glycerol and storage in glycerol for photography. For skeletal preparation, whole litters were analyzed.
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