

The fickle finger of fate

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Commentary

In this issue of the *JCI*, Niedermaier and colleagues demonstrate that a chromosomal inversion in mice results in dysregulation of Sonic hedgehog (Shh), such that *Shh* is ectopically expressed in a skeletogenic domain typically occupied by Indian hedgehog (Ihh). This molecular reversal eliminates phalangeal joint spaces, and consequently, *Short digits* (*Dsh*) heterozygotes (*Dsh/+*) have brachydactyly (shortened digits). *Ihh* is normally downregulated in regions that will become the joint space, but in *Dsh/+* mice, Shh bypasses this regulatory control and persists; accordingly, cells maintain their chondrogenic fate and the developed digits are shorter than normal. The significance of these data extends far beyond the field of skeletal biology: they hint at the very real possibility that the endogenous *Shh* regulatory region contains a repressor designed to segregate the activity of Shh from Ihh. The existence of such a repressor provides a window into the distant past, revealing that Shh and Ihh must once have shared responsibilities in establishing tissue boundaries and orchestrating vertebrate tissue morphogenesis.

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The fickle finger of fate

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In this issue of the JCI, Niedermaier and colleagues demonstrate that a chromosomal inversion in mice results in dysregulation of Sonic hedgehog (Shh), such that *Shh* is ectopically expressed in a skeletogenic domain typically occupied by Indian hedgehog (Ihh) (see the related article beginning on page 900). This molecular reversal eliminates phalangeal joint spaces, and consequently, *Short digits* (*Dsh*) heterozygotes (*Dsh/+*) have brachydactyly (shortened digits). *Ihh* is normally downregulated in regions that will become the joint space, but in *Dsh/+* mice, *Shh* bypasses this regulatory control and persists; accordingly, cells maintain their chondrogenic fate and the developed digits are shorter than normal. The significance of these data extends far beyond the field of skeletal biology: they hint at the very real possibility that the endogenous *Shh* regulatory region contains a repressor designed to segregate the activity of *Shh* from *Ihh*. The existence of such a repressor provides a window into the distant past, revealing that *Shh* and *Ihh* must once have shared responsibilities in establishing tissue boundaries and orchestrating vertebrate tissue morphogenesis.

Ancient physicians viewed the skeleton as “the foundation of the rest of the parts of the body and all the members rest upon them and are supported, as proceeding from a primary base” (1). Defects in this structural foundation also serve as portholes through which the process of

fetal skeletogenesis can be analyzed. For example, studies of genetic perturbations that result in distal limb truncations have shown that the morphogen Sonic hedgehog (*Shh*) establishes precisely where skeletogenic condensations will form in the tips of the hand- and footplates (2, 3). This spatial patterning information is further refined by bone morphogenetic protein (*Bmp*) signaling, as shown by the fact that disruptions in the *Bmp* signaling pathway lead to fusions, or syndactyly, of the digits (4). Once the spatial pattern of the skeletogenic condensations is achieved, a closely

related cousin, Indian hedgehog (*Ihh*), takes over and plays an instrumental role in segregating inner chondrogenic cells from the flattened, elongated perichondrial cells at the periphery (5). *Ihh* secreted from chondrocytes stimulates the differentiation of perichondrial cells into osteoblasts (5–7), mesodermal cells that give rise to bone.

Another critical feature of limb skeletogenesis is the creation of the articulations, or joint spaces, between the skeletal elements. In the fingers, joint spaces are created when a single, larger skeletogenic condensation cleaves into 2 or 3 smaller segments, each of which will give rise to a phalange (8). *Wnt14* is critical in determining where a joint space will form (9), but precisely how the cleavage event is controlled remains uncertain. One thing is clear, however: when a separation fails to take place, the phenotypic consequence is brachydactyly (shortened digits). Thus, while we have a fairly complete picture of the range of skeletal malformations that can occur, how these disruptions are related to one another and to the basic program of skeletogenesis remains unknown.

Reading the bones

In this issue of the *JCI*, Niedermaier et al. provide new insights into the process of

Nonstandard abbreviations used: *Bmp*, bone morphogenetic protein; *Dsh*, *Short digits*; *Dsh/+*, *Dsh* heterozygote; *Ihh*, Indian hedgehog; *Shh*, Sonic hedgehog.

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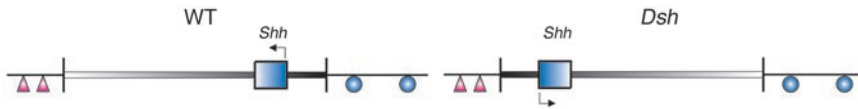


Figure 1
A chromosomal inversion causes *Shh* relocation. Schematic representation of the inversion event leading to development of *Dsh* mutant mice; the *Shh* gene, normally influenced by distal enhancer sequences (blue circles), is moved nearer to other regulatory sequences (pink triangles).

skeletal morphogenesis, describing the phenotypic consequences of a chromosomal inversion that disrupts both temporal and spatial regulation of *Shh* (10). Chromosomal inversions are genetic mishaps that take place when a region of the chromosome breaks and rejoins in a new orientation (Figure 1). This structural aberration can serve as a window into embryogenesis, especially when genes within the inverted region are subjected to new regulation as a result of enhancers or promoters located on the other side of the breakpoint.

In the current study Niedermaier and colleagues examined heterozygous mice in which an inversion caused foreshortening of the distal limb skeletal elements, hence their nom de plume, *Short digits* (*Dsh*) (10). In the homozygous state, the *Dsh* phenotype is far more severe; in fact, *Dsh/Dsh* embryos bear a striking resemblance to *Shh*^{-/-} embryos in that both exhibit cyclopia and incomplete cleavage of the embryonic forebrain. But the emphasis in this study was on the skeletal anomalies of the limb; *Dsh/Dsh* embryos showed an absence of all distal limb skeletal elements. Because *Dsh*^{+/+}*Shh*^{-/-} mice had a phenotype nearly identical to that of *Shh*^{-/-} and *Dsh/Dsh* mice, the authors concluded that *Dsh* and *Shh* were allelic.

inversion functionally removed *Shh* activity from a number of its endogenous sites of activity, but also directed ectopic *Shh* expression to – of all places – the region where *Ihh* is normally expressed, the limb skeletogenic condensations (Figure 2) (10). Ectopic expression of *Shh* occurred simultaneously with the loss of *Ihh* in the digit condensations. While the 2 events – induction of *Shh* and loss of *Ihh* – might be controlled by separate mechanisms, a more plausible explanation is that the loss of normal *Ihh* expression is a secondary consequence of ectopic *Shh* expression, since previous investigators have shown that a negative feedback loop controls Hedgehog signaling in the skeleton (11).

Another phenotypic peculiarity was that the molecular substitution of *Shh* for *Ihh* only took place in the digits, while more proximal skeletal elements (i.e., humerus, radius, and ulna) were spared. Here, the answer may lie in the timing: the limb skeleton develops in a proximal-to-distal direction, and the inversion disrupted *Shh* expression relatively late in gestation, perhaps after the proximal skeletal elements were specified. But what remains a puzzle – and a very exciting one at that – is just how this inversion resulted in ectopic *Shh*

expression in the digits. In other words, where did the instructions driving *Shh* expression in the digits originate? The chromosomal rearrangement appears to have created a new, chimeric regulatory domain for the *Shh* gene, composed partially of resident enhancer sequences and partially of enhancer sequences that were translocated along with the *Shh* gene.

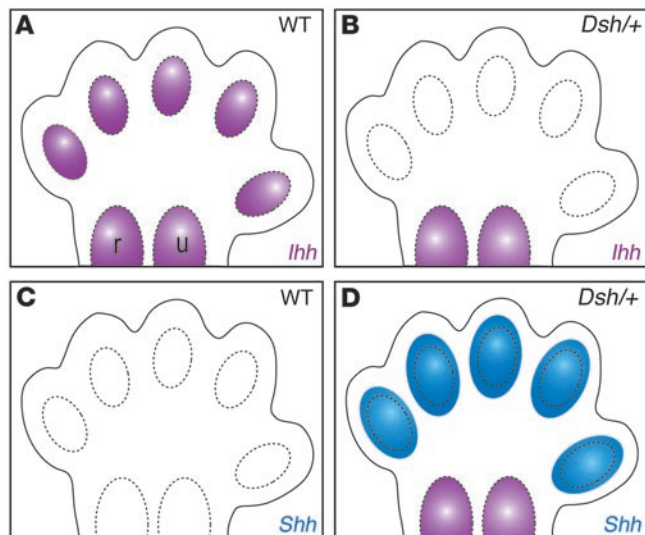
Another perplexing question arises: how was the information in this chimeric regulatory domain derived? One possibility is that the inversion positioned the *Shh* coding sequence so that it was adjacent to a new enhancer that drives the expression of one or more genes in the digits. This possibility seems unlikely, however, as none of the genes near the insert point were expressed in the digits (10). A more plausible and attention-grabbing possibility is that the instructions for driving spatio-temporal expression of *Shh* are contained within the normal *Shh* regulatory region but are somehow masked. If this is the case, then the inversion event may have resulted in the loss of a repressor activity that normally restricts *Shh* from being expressed in skeletogenic condensations.

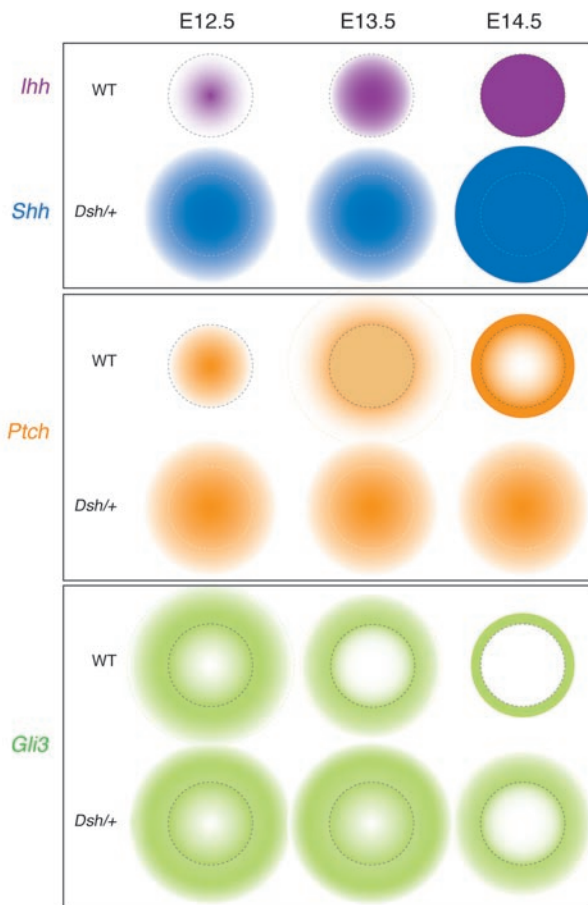
The implications of this finding are far reaching: if a repressor sequence was disrupted by chromosomal rearrangement, then the ectopic *Shh* represents part of a more ancient pattern of expression. Furthermore, these data suggest that the normal *Shh* regulatory sequence may have buried within it instructions for targeting *Shh* expression to skeletogenic condensations. That *Shh* and *Ihh* might have shared expression patterns sometime in the distant past is not a large stretch, since gene duplication

It’s about time; it’s about space

Perhaps the most unexpected finding came from studying the *Dsh*^{+/+} phenotype: here, the authors found that the chromosomal

Figure 2
The chromosomal inversion causes *Shh* to be expressed ectopically in the phalanges. (A and B) In WT embryos (A), *Ihh* is expressed in mesenchymal aggregates of cells called cartilage condensations, up to the cartilage-perichondrium boundary (dotted lines); whereas in *Dsh*^{+/+} embryos (B), *Ihh* is lost from the phalangeal condensations. (C and D) Whereas in WT embryos (C), *Shh* is never expressed in skeletal condensations, in *Dsh*^{+/+} embryos (D), *Shh* expression replaces *Ihh*, albeit in a larger domain that extends past the cartilage-perichondrium boundary. u, ulna; r, radius.



**Figure 3**

Disrupted expression boundaries in *Dsh/+* animals lead to perichondrial and joint defects (1). A transverse schematic of the phalangeal skeletal condensation, showing expression domains corresponding to a specific Hedgehog gene (*Ihh* in WT, *Shh* in *Dsh/+*), the Hedgehog target *Ptch*, and the Hedgehog repressor *Gli3*. Dotted lines indicate the normal cartilage-perichondrium boundary. In WT phalanges, *Ihh* (purple) is initially expressed (E12.5) in a small domain that expands over time to encompass the skeletal condensation. At E12.5 in *Dsh/+* phalanges, *Shh* expression (blue) encompasses the normal *Ihh* domain and over time extends beyond what would normally be the cartilage-perichondrium boundary. *Ptch* (orange), an indicator of Hedgehog responsiveness, is first expressed in WT phalanges coincident with *Ihh*; eventually, *Ptch* is downregulated in cells expressing *Ihh* and upregulated in cells adjacent to the *Ihh* domain, where it functions to limit the spread of Hedgehog expression. Note the absence of color within the element, which shows the relative lack of *Ptch* in this domain (10). In *Dsh/+* phalanges, *Ptch* is expressed coincident with the broader domain of *Shh*; however, in contrast to what occurs in WT tissues, *Ptch* does not become downregulated in chondrocytes. *Gli3* (green), a repressor of Hedgehog signaling, is initially expressed in WT phalanges in a domain overlapping with *Ihh*; over time, this domain is restricted to the perichondrium, where *Gli3* limits the activity of the Hedgehog protein. In *Dsh/+* phalanges the *Gli3* expression domain is initially broader, and only after an extended period of time does it become restricted to the perichondrium.

is thought to be one of the chief mechanisms responsible for the diversification of gene function.

Limbs and faces

The precise regulatory region perturbed by the chromosomal translocation in *Dsh/+* mice is unknown, but it appears to be responsible for controlling ectopic *Shh* expression in both the limbs and the facial prominences. Niedermaier and colleagues report that *Shh* is ectopically expressed in the *Dsh/+* frontonasal prominence, which leads to flaws in craniofacial patterning (10). In this anatomical locale, *Shh* regulates proximodistal outgrowth and mediolateral expansion of the neural crest cell population that resides in the facial midline (12). Eventually these cells will produce the skeletal elements of the middle and upper face, a developmental step that is also dependent upon Hedgehog signaling (13, 14). Clinicians and researchers have long been aware of the connection between facial and limb malformations, and a surprisingly large number of syndromes are characterized by anomalies in both craniofacial and

be affected by any perturbations, be they genetic or environmental, in that regulatory domain.

A molecular switcheroo

Niedermaier et al.'s molecular dissection of the *Dsh/+* phenotype has provided additional insights into the molecular underpinning of brachydactyly. Specifically, the investigators show that *Shh* does not simply replace *Ihh* in *Dsh/+* chondrocytes; rather, *Shh* is more widely distributed in the cartilage and perichondrium of the digits, and it is active in this larger domain, as illustrated by the expansion of Hedgehog target genes *Gli1* and *Ptch* (Figure 3) (10).

The consequences of ectopic Hedgehog expression are delayed chondrocyte differentiation, retardation of the maturation of perichondrium into periosteum, and loss of distal joint spaces. Given that all 3 defects occur simultaneously, one cannot help but wonder if they are interrelated features, all initially dependent upon a single precipitating event. We speculate that this event is the establishment of a boundary that segregates chondrocytes and osteoblasts into

adjacent but immiscible cell populations. This boundary is, to some extent, a consequence of local Hedgehog signaling.

Around the same time that the cartilage-perichondrium boundary is being specified, some cells within the *Ihh*-positive skeletal condensation are faced with another decision: whether to progress along a chondrogenic pathway and undergo hypertrophy or to differentiate into a specialized type of perichondrial cell that will line the joint space (16). This cell fate decision, along with formation of a proper perichondrium, is disrupted in the digits of *Dsh/+* mice as a consequence of ectopic Hedgehog activity (10). Just how does this imbalance between Hedgehog responsiveness and repression cause cells to alter their fate? Organisms use boundaries, the juxtaposition of like and unlike cells, in a myriad of ways: to define compartments, convey positional information, and even establish new tissues and signaling centers that can only form at the interface between 2 populations. In a wide variety of other developmental contexts, Hedgehog proteins play a critical role in specifying boundaries and thus defining cell and tissue compartments (17–19); the same may be true during skeletogenesis, but data to support this hypothesis are still sparse. What the Niedermaier et al. study reveals, however, is that removal or expansion of one of the factors that contributes to the establishment of a boundary can cause a multitude of processes, including those that shape and control development of the skeleton, to go awry.



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Altered regulation of IL-2 production in systemic lupus erythematosus: an evolving paradigm

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In systemic lupus erythematosus (SLE), IL-2 production by T lymphocytes in vitro is impaired. Deficient IL-2 production may be an outcome of a primary SLE T cell disorder that is due to impaired signal transduction. In this issue of the JCI, evidence is presented that an anti-TCR/CD3 complex autoantibody present in SLE sera can bind to T cells and activate the Ca²⁺-calmodulin kinase IV (CaMKIV) signaling cascade, resulting in downregulation of IL-2 transcription and IL-2 production (see the related article beginning on page 996). Because IL-2 may contribute to the maintenance of T cell tolerance, deficient IL-2 production could promote a breach of T cell tolerance that results in autoantibody production in SLE.

Recently, it has been recognized that diverse autoantibodies directed against intra- and extracellular autoantigens exist in patients with systemic lupus erythematosus (SLE) for years before the clinical diagnosis is made (1); this suggests that physiologic mechanisms

that maintain tolerance to self antigens have been breached. Tolerance to self antigens is established and preserved by a subpopulation of T lymphocytes known as Tregs (2), and the loss of tolerance is a pathologic process giving rise to autoimmunity. This circumstance raises the possibility of the existence of abnormal T cell clones that mediate defective helper and suppressor effector functions, which result in autoantibody generation by forbidden B cell clones. In SLE, defective signaling cascades are believed to give rise to a primary T cell disorder that is characterized by impaired effector functions (3). These effector dysfunctions are, at least in part, a result of skewed expression of

various effector molecules, including CD40 ligand (e.g., CD154) and multiple cytokines, and may reflect an imbalance of gene expression. An extracellular factor(s) in the microenvironment that interacts with T cells and exacerbates these dysfunctions has not been previously identified.

Tregs, skewed cytokine production, and loss of tolerance

Impaired effector T cell functions due to skewed cytokine production may create a microenvironment that promotes a strong Th2 immune response relative to Th1 and Treg activity. Relative overproduction of IL-4, IL-6, and IL-10 by Th2 cells and underproduction of IL-2, IL-12, TGF- β , and IFN- γ by Th1 cells and Tregs can result in imbalanced autocrine and paracrine effects on T and B cells in the microenvironment. Because of the reduced numbers of CD4⁺CD25⁺ Tregs (4) as well as the diminished generation of IL-2 and TGF- β , there may be insufficient suppressor activity in SLE to counterbalance the enhanced Th2 effect on B cell antibody production. Taken together, these conditions create a microenvironment that promotes a

Nonstandard abbreviations used: CaMKIV, Ca²⁺-calmodulin kinase IV; CRE, cAMP response element; CREB, CRE-binding protein; CREM, CRE modulator; IL-2R, IL-2 receptor; pCREB, phosphorylated CREB; pCREM, phosphorylated CREM; PKA-II β , type II β PKA; pRII β , phosphorylated RII β ; RII β , β type II regulatory subunit; SLE, systemic lupus erythematosus.

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