Targeted disruption of Hoxd-10 affects mouse hindlimb development

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SUMMARY

Targeted disruption of the Hoxd-10 gene, a 5′ member of the mouse HoxD linkage group, produces mice with hindlimb-specific defects in gait and adduction. To determine the underlying causes of this locomotor defect, mutant mice were examined for skeletal, muscular and neural abnormalities. Mutant mice exhibit alterations in the vertebral column and in the bones of the hindlimb. Sacral vertebrae beginning at the level of S2 exhibit homeotic transformations to adopt the morphology of the next most anterior vertebra. In the hindlimb, there is an anterior shift in the position of the patella, an occasional production of an anterior sesamoid bone, and an outward rotation of the lower part of the leg, all of which contribute to the defects in locomotion. No major alterations in hindlimb musculature were observed, but defects in the nervous system were evident. There was a decrease in the number of spinal segments projecting nerve fibers through the sacral plexus to innervate the musculature of the hindlimb. Deletion of a hindlimb nerve was seen in some animals, and a shift was evident in the position of the lumbar lateral motor column. These observations suggest a role for the Hoxd-10 gene in establishing regional identity within the spinal cord and imply that patterning of the spinal cord may have intrinsic components and is not completely imposed by the surrounding mesoderm.

Key words: Hoxd-10, gene targeting, limb defects, homeosis, neuronal defects, spinal cord defects

INTRODUCTION

The mouse Hox complex contains 39 genes distributed in four linkage groups designated HoxA, B, C and D. Each gene encodes a transcription factor belonging to the Antennapedia homeodomain class. Based on DNA sequence similarities and on the position of the genes on their respective chromosome, individual members of the four linkage groups have been classified into thirteen paralogous families (Scott, 1992). Members of a paralogous family often share common patterns of gene expression and function. Mutational analysis in the mouse has demonstrated that Hox genes, alone and in concert with other Hox genes, are used to regionalize the embryo along its major axes (reviewed by Krumlauf, 1994). Regionalization of the embryo by Hox genes appears to be accomplished by the controlled temporal and spatial activation of these genes, such that a 3′ Hox gene is activated prior to and in a more anterior region of the embryo than its 5′ neighbors (Duboule and Dollé, 1989; Graham et al., 1989; Duboule, 1994). However, Hox genes normally function not as individual entities, but as members of highly integrated circuits, such that paralogous genes, adjacent genes on the same linkage groups, and even nonparalogous genes in separate linkage groups interact positively, negatively, and in parallel with each other to orchestrate the morphological regionalization of the embryo (Condie and Capecchi, 1994; Rancourt et al., 1995; Davis et al., 1995; Horan et al., 1995a,b; Davis and Capecchi, 1996; Favier et al., 1996; Fromental-Ramain et al., 1996a,b; van der Hoeven et al., 1996; Zákány and Duboule, 1996; Zákány and Duboule, 1997).

The sixteen 5′ Hox genes belonging to the paralogous families 9 through 13 all show DNA sequence similarities to the Drosophila AbdB gene, which specifies the identity of the most posterior segments of the larva and adult fly. The homologous mouse genes are also expressed in the posterior region of the mouse embryo and in the developing limbs (Dollé and Duboule, 1989; Dollé et al., 1989). Targeted disruptions of some of these genes have produced defects in the formation of the axial skeleton, in the genesis of posterior internal and external organs, in the specification of motor nerves, and in the patterning of limb bones (Dollé et al., 1993; Small and Potter, 1993; Davis and Capecchi, 1994; Davis et al., 1995; Satokata et al., 1995; Rijli et al., 1995; Davis and Capecchi, 1996; Favier et al., 1996; Fromental-Ramain et al., 1996a,b; Kondo et al., 1996; van der Hoeven et al., 1996; Zákány and Duboule, 1996). In this paper we describe the defects associated with the disruption of Hoxd-10. The 10th paralogous gene family contains three members, Hoxa-10, Hoxc-10 and Hoxd-10. The consequences of mutating the Hoxc-10 gene have not been reported, but description of an ES cell line with a mutation in the Hoxd-10 gene has been published (Rijli et al., 1994). However, generation of ES cells with this mutant allele appears to have compromised the ES cell line, since mouse chimeras made from this cell line die in utero. Mice homozygous for the Hoxa-10 mutation show anterior homeotic transformations of the lumbar
vertebrae, male and female sterility, anterior transformations of thoracic and lumbar motor neurons, and defects in the proximal hindlimbs (Rijli et al., 1995; Satokata et al., 1995; Favier et al., 1996). Except for the defects in the hindlimbs, there is remarkably little overlap between the reported mutant phenotypes resulting from disruption of Hoxa-10 and those we describe here resulting from Hoxd-10 disruption. In Hoxd-10 mutants, anterior transformations of the axial skeleton are restricted to the sacral and first caudal vertebrae. Similarly, the neuronal defects seen in Hoxd-10 mutant homozygotes are posterior to those observed in Hoxa-10 mutants.

MATERIALS AND METHODS

Generation of Hoxd-10 mutant mice and genotype analysis

A replacement-type gene targeting vector composed of 14.8 kilobases (kb) of genomic DNA containing the Hoxd-10 gene and flanked by two thymidine kinase genes was constructed in a pUC plasmid (Fig. 1; Thomas and Capecchi, 1987; Deng et al., 1993). To disrupt the Hoxd-10 gene, the neomycin resistance (neo) expression cassette KT3NP4 (Deng et al., 1993) was inserted at the HpaI site in exon 2 in the same transcriptional orientation as the Hoxd-10 gene. The cleavage site corresponds to amino acid 38 in the second alpha-helix of the homeodomain.

The targeting vector was linearized and electroporated into R1 ES cells (Nagy et al., 1993), and cell selection and DNA extraction were performed as described by Mansour et al. (1988); Davis and Capecchi (1994). A primary screen for the cell lines containing targeted disruption of the Hoxd-10 locus was done by Southern blot analysis of SphI-digested genomic DNA using a 2.6 kb Xhol/SphI 3' flanking probe (Fig. 1). This probe hybridizes to a large fragment (greater than 23 kb) in wild-type DNA and to a 12.1 kb DNA fragment in the Hoxd-10 mutant allele. Additional diagnostic digests and probe hybridizations (including a neo-specific probe) confirmed the integrity of the targeting event (data not shown). One out of 123 cell lines analyzed tested positive for a gene targeting event. This ES cell line was injected into C57Bl/6J (B6) blastocysts to produce chimeric mice. Chimeras were mated to C57Bl/6J mice and the resulting offspring were mated to C57Bl/6J mice to obtain homozygous Hoxd-10 mutant bands.

Histology and skeletal preparations

Newborn mice were fixed in Bouin’s solution at room temperature for 1 week, dehydrated through graded ethanol and xylene and embedded in paraffin. 10 µm serial sections were collected and stained with hematoxylin and eosin (Chisaka and Capecchi, 1991). Age and sex-matched littersmates were collected and used for skeleton preparations as described (Mansour et al., 1993).

Whole-mount immunohistochemistry

Embryos were fixed and embedded (Carpenter et al., 1993) and then eviscerated to expose the ventral side of the spinal column and the nerves and plexus at the base of the hindlimb. Embryos were reflected in 4:1 methanol (MeOH):dimethylsulfoxide (DMSO) overnight at room temperature, bleached with 6% H2O2 in 4:1 MeOH:DMSO, and rehydrated through graded MeOH:PBS with addition of 0.5% Tween 20 (PBST) to 100% PBST. Embryos were blocked with PBST, 2% skim milk powder, 1% DMSO, and then incubated overnight at 4°C in a 1:1 dilution (in blocking solution) of the 2H3 antibody (Developmental Studies Hybridoma Bank) directed against a subunit of the neurofilament protein. Horseradish peroxidase-conjugated goat anti-mouse secondary antibody (Jackson Immunoresearch) was used at a dilution of 1:100 and detected with 0.0015% H2O2 in 1 mg/ml diaminobenzidine (DAB)/PBST. Embryos were rinsed, then stained overnight with 0.02% alcan blue 8GX as described by Rancourt et al. (1995). The skin was gently peeled away from the hindlimbs and the embryos photographed under a dissecting microscope.

Whole-mount in situ hybridization

Embryos were collected at E12.5 and E13.5, fixed overnight in 4% paraformaldehyde/PBS and processed for whole-mount in situ hybridization with Hoxd-9 and Hoxd-11 antisense RNA probes as previously described (Carpenter et al., 1993). The Hoxd-9 probe was made from a 770 bp PstI-EcorI fragment in the 3' untranslated region of the Hoxd-9 gene (Renucci et al., 1992); the Hoxd-11 probe was made from a 300 bp AccI-BamHI fragment from the Hoxd-11 gene including part of the homeodomain and 3' untranslated sequence (Duboule, 1991). Probes were labeled with digoxigenin-11-UTP (Boehringer Mannheim) and detected with alkaline phosphatase-conjugated antibodies directed against digoxigenin (Boehringer Mannheim). Alkaline phosphatase reaction product was visualized with NBT and BCIP (Boehringer Mannheim).

RESULTS

Targeted mutagenesis of Hoxd-10

A targeting vector designed to disrupt the Hoxd-10 locus (Fig. 1) was electroporated into R1 ES cells. One cell line, shown by Southern blot analysis to carry the Hoxd-10 mutant allele, was used to generate chimeric mice that transmitted the mutant allele through the germline (see Materials and Methods). Genotypes at birth and at weaning segregated in the expected Mendelian ratios showing no preferential loss of mutant homozygotes.

Alterations in gait, adduction, and male fertility

Adult Hoxd-10 mutant mice exhibited hindlimb defects manifested in two locomotor phenotypes: an abnormal gait and adduction, the inability to bring their hindlegs towards the ventral midline. The gait phenotype varied in severity, with 24% of the mice acting outwardly normal; the remainder demonstrated mild to severe deficits (Table 1). In mildly affected mice, the gait was jerky, with the hindlimbs being stiff, but not spastic; the hindfeet were splayed out in an abnormal
posture. Severely affected mice had very stiff hindlimbs, walked with their hindlimbs outstretched behind them, and swung their hindfeet forward with effort in order to support their weight. In some cases, the animal put its weight on the top of the foot rather than the bottom. Affected hindlimbs retracted in response to touch. One heterozygous animal also had a mild gait defect, but others were unaffected. These defects were present in mutants as they first began to walk and were not progressive.

To test adduction, a mouse was positioned on a pencil along its ventral midline (Fig. 2). When support was withdrawn, wild-type and heterozygous Hoxd-10 mice responded by gripping the pencil with both of their forefeet and hindfeet and maintaining balance (Fig. 2A). Mutant homozygotes were unable to bring their hindfeet together to grip the pencil (Fig. 2B). Most mutant animals tested showed this adduction deficit, even those in which the gait defects were minimal (Table 1). A single heterozygous animal was also unable to grip the pencil.

An additional defect apparent in adults was a marked decrease in fertility in mutant homozygous males. Six mutant males were tested in three trials for their ability either to mate (as judged by the presence of copulation plugs) or to induce a pregnancy. One of these six males impregnated a female on one mating trial; all other males were unable to copulate. Histological sections through the male reproductive system revealed no differences relative to control animals (data not shown). The testes had descended, all of the reproductive organs appeared normal, and normal quantities of sperm were present in the epididymis and vas deferens. The decreased fertility may be due to the hindlimb locomotor problems interfering with mounting.

Six homozygous mutant female mice were tested for fertility in matings with heterozygous or wild-type males. All females were able to produce offspring, but there was significant mortality in the first two litters from four of these females. All pups died during the first postnatal day with an absence of milk in their stomachs. Pups taken from such crosses and fostered with lactating Swiss Webster females survived to adulthood. However, in subsequent litters, these mutant females were able to successfully raise their pups.

**Alterations in the axial and appendicular skeleton**

Mice normally possess four sacral vertebrae. The most anterior sacral vertebra (S1) fuses with the pelvis to anchor the pelvic girdle. Both S1 and S2 vertebrae have winged transverse processes, while S3 and S4 have anteriorly projecting, club-shaped processes. In wild-type animals, S1, S2, and S3 are fused with each other at the tips of their transverse processes, while S3 and S4 have anteriorly projecting, club-shaped processes. In wild-type animals, S1, S2, and S3 are fused with each other at the tips of their transverse processes, while S4 remains unfused (Fig. 3A). Fifteen mutant skeletons were examined for changes in the sacral and caudal vertebrae. Mutant animals showed an increased number of fused sacral vertebrae and alterations in the shape of the transverse processes (Fig. 3B,C). Most had three or four sacral vertebrae with winged transverse processes compared with only two seen in wild-type animals. Half of the mutant animals also had four and occasionally five fused vertebrae (Fig. 3). Two heterozygous animals also had unilateral fusions of four transverse processes. Additionally, in mutants, both the S1 and S2 vertebrae articulated with the pelvis. This is suggestive of an anterior transformation of the S2 vertebra into an additional S1 with more posterior sacral and anterior caudal vertebrae assuming a more anterior phenotype.

Skeletal alterations were also observed in the hindlimbs of mutant animals. The position of the patella was shifted upward to the forward surface of the femur in 13 of 16 mutant hindlimbs examined (Fig. 4A-D). In addition, an ectopic sesamoid bone was observed embedded in the patellar tendon at the proximal edge of the patella in 5 of 16 mutant animals.
Based on similarities in shape and position to the lateral sesamoid bones, this bone was termed an anterior sesamoid. Ectopic sesamoids are often generated as a secondary remodeling response to changes in tension at a joint (e.g., Davis and Capecchi, 1994; Favier et al., 1996). Changes were also noted in the articulation between femur and tibia which were shifted out of alignment with the tibia rotated outward (Fig. 4E,F). Rotations were observed in 10 of 16 mutant hindlimbs. These rotations result in an abnormally positioned foot.

**Alterations in the nervous system**

We examined the developing nervous system by carbocyanine dye tracing, immunohistochemistry and histological analysis. Dye tracing was used to examine limb innervation in five wild-type and seven mutant animals during development. Injections of the lipophilic tracer DiI were made into the base of the developing limbs at E12.5. DiI retrogradely labeled spinal motor neuron somata (Fig. 5A,B) and anterogradely labeled the distal projections of these neurons through the lumbosacral plexus into the limb (Fig. 5C-F). In humans and mice, the lumbosacral plexus has two components: the rostral lumbar plexus and the caudal sacral plexus. Due to the positioning of the embryos during the injections and the position of the plexus elements within the developing hindlimb at E12.5, the DiI injections primarily labeled motor neurons whose fibers pass through the sacral plexus. These fibers travel along the posterior margin of the femur and diverge to form the major nerve trunks providing distal limb innervation. The major nerve arising from this plexus is the sciatic nerve, which branches at the level of the knee to form the tibial and peroneal nerves that innervate the musculature of the lower leg and foot (Marieb, 1995). In nine out of ten limbs examined in five control animals these injections intensely labeled four spinal nerves. The two central nerves contained the largest complement of labeled fibers. These four nerves emerge from lumbar segment L3 through L6. In a single control embryo, fibers emerging from only three spinal segments were observed on one side of the animal. Fibers from all labeled nerves coalesced to pass through the pelvic girdle and the sacral plexus, then diverged into multiple branches in the limb.

Seven mutant embryos were analyzed by carbocyanine dye tracing. One of these embryos had four labeled spinal nerves providing sacral plexus fibers on both sides of the animal. Four animals had four spinal segments providing motor innervation on one side and three spinal segments on the contralateral side (Fig. 5B). The remaining two animals had three segments providing motor innervation bilaterally. In mutant animals having four labeled spinal nerves, the contributions of the most anterior and posterior segments were reduced compared with

**Fig. 2.** Hindlimb adduction in wild-type (A) and Hoxd-10 mutant (B) mice. Wild-type animals can grip a pencil with both forelimbs and hindlimbs, while mutant mice are unable to bring their hindlimbs together in order to grip the pencil with their hindfeet (arrow).

**Fig. 3.** Sacral vertebrae in wild-type and Hoxd-10 mutant adult mouse skeletons. (A) Wild-type animals have two sacral vertebrae with wing-shaped transverse processes (S1, S2), a transitional third sacral vertebra (S3), and one unfused fourth sacral vertebra (S4). (B) In mutants, the first three sacral vertebrae and sometimes the fourth (C) all have wing-shaped transverse processes. In wild-type animals, the first three sacral vertebrae are fused, while in mutants the fourth and sometimes fifth sacral vertebrae are fused. In wild-type animals, only the first sacral vertebra articulates with the ilium, while in mutants, the second sacral vertebra articulates as well (arrows in B and C). Scale bar, 2 mm.

**Fig. 4.** Knee joints and alignment of the lower limb in wild-type (A,C,E) and mutant (B,D,F) animals. A and B illustrate the shift in articulation between femur and tibia following a rotation of the lower leg (white arrows) and an upward shift of the patella (black arrows) in mutant animals. C and D illustrate a shift in the patella (p) and the presence of an additional bone, the anterior sesamoid, anterior to the patella (dotted arrow in D) in mutant animals. E and F show a frontal view of the right knee joint and lower limb in wild-type (E) and Hoxd-10 mutant (F) mice. In F, the tibia has rotated off axis resulting in a lateral twisting of the foot (arrow). The underlying fibula is now exposed. Scale bar, 2 mm.
The contributions from the same segments in control animals. In animals with only three labeled segments, the most anterior segment was missing.

Anterograde tracing with DiI also showed alterations in terminal nerve arborizations of the tibial nerve in mutant animals (Fig. 5C-F). This nerve was tipped with a flattened, hand-shaped terminal arbor of axons, which were bundled into many large fascicles with smaller branching fascicles growing toward their targets. Terminal arbors of this nerve in mutant animals showed a decrease in the size and density of the fascicles (Fig. 5C-F). Labeling of E12.5 embryos with the 2H3 neurofilament antibody showed a similar reduction in the fascicles (Fig. 5C-F). Labeling of E12.5 embryos with the 2H3 neurofilament antibody showed a similar reduction in the density of these terminal arbors (data not shown).

To visualize the complete set of nerves as it innervates the developing hindlimb, animals at E13.5 were examined using whole-mount labeling with the 2H3 antibody (Fig. 6). As the spinal nerves enter the developing hindlimb, they diverge into the lumbar and sacral plexuses which are positioned respectively anteriorly and posteriorly around the head of the femur. Muscle nerves branch off distally from these plexuses to provide the innervation to the hindlimb musculature. In 3 of 10 mutant hindlimbs examined, one of these muscle nerves, the peroneal nerve, was missing (Fig. 6). This nerve was present in all 21 heterozygotes and 6 wild-type control hindlimbs examined.

Transverse sections of newborn animals showed that the spinal cord, dorsal root ganglia, and spinal nerves were grossly normal in mutant animals (Fig. 7). Spinal nerves emerged from the appropriate intervertebral level accompanied by Schwann cells and followed normal trajectories to innervate the limb. Dorsal root ganglia were located in the expected places and appeared normal in size and shape. The spinal cord appeared to be the same size as in littermate controls, and its laminar organization appeared normal. The carbocyanine dye injections suggested that there might be some alterations in the population of motor neurons innervating the hindlimb, so the morphology and position of spinal motor neurons and the lateral motor column (LMC) were examined. The LMC containing the motor neurons that provide hindlimb innervation is present in the spinal cord in five lumbar segments, L1 through L5 in newborn mice (Lance-Jones, 1982). A population of large, ventrolaterally positioned cells resembling spinal motor neurons was readily visible in transverse sections through wild-type and mutant newborn animals (Fig. 7). In our wild-type animals, however, these cells extended more rostrally than reported, beginning opposite the rostral portion of thoracic (T) vertebra 13 and continuing posteriorly into L5. The majority of LMC motor neurons were adjacent to the L1 and L2 vertebrae, with smaller populations present as far caudally as L5. This change in position may reflect differences between the mouse strain used for previous motor neuron mapping studies (CD1) and the strains used in construction of Hoxd-10 mutant mice (B6 and 129). In mutant animals, the LMC neurons were displaced posteriorly by half a segment, with the neurons first appearing near the middle of the T13 vertebra. The largest populations of these neurons were adjacent to the L2 and L3 vertebrae.

**Gross anatomy of hindlimb musculature and innervation**

Hindlimbs from six mutant and four wild-type adult animals were examined for alterations in the organization of musculature. Muscles from the anterior, medial and lateral compartments of the distal part of the leg were dissected and scored.
for the presence and the appearance of the musculature (Table 2). Mice have five muscles in the lateral compartment, in contrast to humans which have only three muscles in comparable locations (Marieb, 1995). Three of these five muscles, peroneus brevis, peroneus longus, and peroneus tertius, were correlated with their human counterparts. Two additional muscles provide fine motor control of the fourth and fifth digits. We have termed these additional muscles the fourth digit dorsolateral and the fifth digit lateral, based on the position of their tendon insertion sites into the fourth and fifth digits. In general, the musculature of the mutant animals appeared grossly normal. Most muscles were present in their expected locations; the only differences noted with some frequency were the misrouting of tendons to their insertion points (Table 2). This may be a response to alterations in the position of the leg imparted by the skeletal abnormalities described above. Cross sectional area and muscle position were also examined on sections taken through hindlimbs of newborn animals. Muscles and their positions in mutant animals appeared normal in comparison to wild-type littermates (data not shown).

Muscle nerves were also examined in dissected hindlimbs and the presence and position of six nerves were scored (Table 2). With the exception of a slight reduction in the size of the common peroneus in two mutant animals, there were no differences in the position and trajectory of the hindlimb nerves examined.

Hoxd-9 and Hoxd-11 expression appears normal in Hoxd-10 mutant homozygotes

The expression of Hoxd-9 and Hoxd-11 was examined in Hoxd-10 mutant embryos (E12.5 to E13.5) to determine whether the disruption of Hoxd-10 affected the expression patterns of neighboring genes. Expression of Hoxd-9 in wild-type, Hoxd-10 heterozygous, and Hoxd-10 homozygous mutant embryos was found in forelimb, hindlimb, spinal cord, and prevertebrae adjacent to the hindlimbs as previously reported (Dolle and Duboule, 1989). In wild-type embryos at E12.5, Hoxd-9 expression had an approximate anterior border in the spinal cord at the level of prevertebra 19. High levels of expression were maintained posteriorly through anterior lumbar segments and then decreased at more caudal levels. Labeling in the prevertebrae was observed beginning at prevertebra 22 and continued caudally for approximately 8 segments. The patterns of Hoxd-9 expression were not altered in Hoxd-10 heterozygous and mutant homozygous embryos (data not shown). Like Hoxd-9, Hoxd-11 is normally expressed in the spinal cord, prevertebrae and developing limb buds (Izpisúa-Belmonte et al., 1991). The patterns of Hoxd-11 expression in wild-type, Hoxd-10 heterozygous and Hoxd-10 homozygous embryos were indistinguishable (data not shown).

**DISCUSSION**

Targeted disruption of the Hoxd-10 gene results in mice with adduction and gait defects affecting the hindlimbs. In these mutants, both the axial and appendicular skeletons are affected, as is the organization and robustness of the hindlimb motor innervations. These alterations may provide a cause for the observed defects in locomotion and reproductive behavior.

Disruption of the Hoxd-10 gene results in anterior transformation of sacral vertebrae, with S2 transformed into a second S1 and three to four more posterior vertebrae also adopting a more anterior morphology. This results in a duplication of S1 and a secondary articulation to the ilium. Disruption of the paralogous gene Hoxa-10 also results in homeosis, though at a more anterior level, with L1 transformed into T14 and L2, L3 and L4 acquiring characteristics normally associated with their immediate anterior vertebrae, respectively (Rijli et al., 1995). Curiously, disruption of the more 5’ Hoxa gene, Hoxa-11, also results in a slightly more anterior homeotic transformation in the lumbosacral region, with S1 being transformed to a 7th lumbar vertebra, S2 to S1 and so on (Davis and Capecchi, 1994; Favier et al., 1995). In this context the transformations resulting from the Hoxd-10 mutation appear to be more posterior than anticipated either from the behavior of mutations in paralogous family members and adjacent 5’ genes, or from the anterior limit of Hoxd-10 expression in the prevertebrae. Clearly it is the coordinated temporal and spatial expression of multiple Hox genes in any given region that specifies the identity of groups of vertebrae. Disruption of any given Hox gene results in a break in this coordinated pattern. The mutant phenotype is exposed in the region of minimal functional overlap. Often this occurs at the anterior limit of expression of the mutant gene; however, in the case of Hoxd-10, it occurs at a more posterior level. It is important to remember, however, that the expression patterns of these genes are dynamic and that the quantitative output, i.e., how many Hox genes are expressed in a given region, is as important as which specific Hox genes are expressed in the region. Thus, the point of minimal functional overlap could occur early or late during the formation of the affected vertebrae.

**Table 2. Number of Hoxd-10 mutant mice examined for hindlimb muscles and nerves**

<table>
<thead>
<tr>
<th>Genotype</th>
<th>+/+</th>
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<tr>
<td><strong>Muscles</strong></td>
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<tr>
<td>Anterior</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Extensor digitorum longus</td>
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<td>12*/12</td>
</tr>
<tr>
<td>Tibialis anterior</td>
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<tr>
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<td>10/12</td>
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<tr>
<td>Medial</td>
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<td></td>
</tr>
<tr>
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<td>8/8</td>
<td>12/12</td>
</tr>
<tr>
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<td>8/8</td>
<td>12‡/12</td>
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<td>7/10</td>
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<tr>
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<tr>
<td>Peroneus longus</td>
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<td>9**/10</td>
</tr>
<tr>
<td>4th digit dorsolateral</td>
<td>5/5</td>
<td>8/8</td>
</tr>
<tr>
<td>5th digit lateral</td>
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<tr>
<td>Posterior femoral cutaneous</td>
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<td>10/10</td>
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</table>

*1 abnormal (shifted laterally); †2 abnormal (reduced or shifted); §1 abnormal (tendon displaced); ¶1 abnormal (duplicated tendon); ¶¶1 abnormal (bifurcated tendon); **1 reduced, 1 potentially absent (or 5th digit lateral); ††1 missing (or p. tertius); ‡‡ reduced in 2 cases.
Skeletal alterations are also observed in the hindlimbs of Hoxd-10 mutants, including a shift in the position of the patella, and the presence of an ectopic sesamoid. These changes are likely to involve the patellar tendon, in which both the patella and the anterior sesamoid are embedded. Altering the tension produced by this tendon through shortening and/or improper alignment might cause the anterior shift of the patella with a simultaneous shift forward of the lower part of the hindlimb skeleton, affecting the articulation between femur andibia. This would destabilize the knee joint and produce the outward rotation in the lower part of the leg. There were no alterations in the shape of the distal part of the femur and no extra ligaments that might provide an alternate explanation for the reshaping and destabilization of the knee joint, as is observed in the Hoxa-10 mutant hindlimbs (Favier et al., 1996). The defects in the formation of the knee are consistent with strong expression of Hoxd-10 in this joint (Dollé and Duboule, 1989; Favier et al., 1996). The destabilization of the knee joint and rotation of the lower limb could be sufficient to explain the defects in gait and adduction.

Unexpectedly, mutations in Hoxa-10, Hoxd-10 and even in Hoxc-10 (S.L. Hostikka and M. Capecchi, unpublished results) primarily cause defects in the formation of the knee joint. Mutations in other 5' Hox genes primarily affect the formation of forelimbs (Hoxa-11, Hoxd-11 and Hoxd-12) or both fore- and hindlimbs (Hoxa-13 and Hoxd-13). However, the elbow and knee joints are particularly disparate structures and their differences need to be genetically programmed. It appears that one of the roles of Hox-10 paralogs is to mediate the formation of these structural and functional differences.

Disruption of Hoxa-10 affects male fertility due to a malformation in the gubernaculum (Rijli et al., 1995; Satokata et al., 1995). Though male fertility was affected in Hoxd-10 mutants, the testes were present in their normal locations and histological analysis suggested that the testes, epididymis, vas deferens and sperm production were normal. The effect of Hox-10 on male fertility, therefore, is not likely to stem from abnormalities in the male reproductive system, but is likely to be operational instead. Thus, the hindlimb locomotor defects in mutant males may interfere with their ability to mount and impregnate a female.

There are no major alterations in the hindlimb musculature of Hoxd-10 mutants since all expected distal hindlimb muscles were present in both newborn and adult mutants. Tendons derived from these muscles showed occasional abnormalities such as duplications or misrouting, and in one case there was a deletion of one of the peroneal group muscles. Preliminary observations on myosin expression indicate that the normal complement of muscle protein is produced in these muscles (Craig Neville and Nadia Rosenthal, personal communication).

Hox genes have been implicated in pattern formation of the central nervous system at the level of the hindbrain. Hox gene activity is required for the production and/or maintenance of rhombomeres (Carpenter et al., 1993; Mark et al., 1993) and for the production of facial motor neurons (Barrow and Capecchi, 1996; Goddard et al., 1996; Studer et al., 1996). Hox genes are also expressed in the developing spinal cord, where they may act as regulators of neuronal identity. Embryological studies have suggested that the segmented appearance of spinal nerves and dorsal root ganglia is generated by the surrounding paraxial mesoderm (Detweiler, 1934; Keynes and Stern, 1984; Phelan and Hollyday, 1990). This implies that there is no intrinsic segmental organization in the spinal cord. However, there are some intrinsic features of spinal cord patterning which may result from activity of Hox genes or other transcriptional regulators. One feature is the position of the lateral motor columns (LMC) adjacent to limbs. While signals from the lateral plate mesoderm of the developing limb bud may establish the position of the LMCs, evidence from limbless chicken mutants (Lanser et al., 1986) and from limb ablation experiments (Hamburger, 1934; Hamburger, 1958; Oppenheim et al., 1978; Lamb, 1981; reviewed by Hamburger and Oppenheim, 1982) suggests that LMCs are produced even in the absence of a limb bud, but that limb buds are required for continued survival of LMC neurons. Our observations on Hoxd-10 mutants, as well as published observations on other Hox gene mutants, suggest that these genes may contribute to the establishment of intrinsic patterning within the spinal cord (LeMouellic et al., 1992; Rijli et al., 1995). Hoxd-10 mutant animals have a decrease in the spinal cord domain providing motor innervation via the sacral plexus as well as a shift in the segmental position of LMC neurons. Additionally, there is a decrease in fiber number in the tibial nerve and an occasional loss of the peroneal nerve. Hoxa-10 mutant animals also show alterations in peripheral nerves (Rijli et al., 1995), which may reflect concomitant alterations in the spinal cord. Hox-8 mutants have defects in autopod clenching which may also result from motor neuron defects at the brachial levels (LeMouellic et al., 1992).

Alterations in the peripheral nerves in Hoxa-10 mutants are particularly interesting in light of our current observations on Hoxd-10 mutants. Rijli et al. (1995) interpret their peripheral nerve alterations as anterior transformations of the spinal nerves emanating from segments T13 through a duplicated L5 (corresponding to wild-type T13 through L6). This suggests that a homeotic transformation of central nervous system structures (e.g., the point of origin of the spinal nerves) is occurring and that it is anterior to the transformation observed in the vertebral column. Hoxd-10 mutants suggest similar transformations in the spinal cord and vertebral column. We observed a reduction in the number of spinal segments providing motor innervation through the sacral plexus during embryogenesis and a shift in the position of the major populations of neurons in the LMC of newborn animals. These two observations could be explained by proposing an anterior transformation in the spinal cord such that the spinal L3 segment is transformed anteriorly to L2 with no additional posterior segments undergoing transformation (Fig. 8). Thus, there would be no overall change in the number of spinal cord segments. Lateral motor column motor neuron histogenesis occurs in L1 through L6 (Lance-Jones, 1982) with the embryonic motor neurons projecting through the sacral plexus located in L3 through L6. An L3 to L2 transformation would reduce the number of motor neurons projecting through the sacral plexus. The reduction in spinal nerves passing through the sacral plexus supports this hypothesis. Additionally, part of the tibial and peroneal motor pools are derived from L3 (McHanwell and Bisoeoe, 1981), so the reduction in fiber number in the developing tibial nerve and occasional deletion of the peroneal nerve might reflect the deletion or respecification of those motor neurons which would normally be generated in L3.

The neuronal defects observed in Hoxd-10 mutants are more
pronounced in embryos and newborn mice than in adults. This difference may reflect the plasticity of the nervous system. Motor neurons are, as most neurons, produced in excess and then reduced through apoptosis during refinement of the neuron-target connections. Since the Hoxd-10 mutation appears to reduce but not eliminate the allocation of nerves to the sacral plexus, gross neuroanatomical defects are not apparent in the adult. This may reflect a rescue of cells normally destined for apoptosis or a recruitment of ectopic neurons to provide distal limb innervations. It will be of interest, however, to determine whether fine motor control in the hindfeet has been compromised in these mutants.

A possibility exists that the changes in prevertebral identity observed in Hoxa-10 and Hoxd-10 mutant mice influence the patterning of the nervous system in these mutant mice. This appears unlikely since in both Hoxa-10 and Hoxd-10 mutant homozygotes the alterations in the nervous system occur anteriorly to those observed in the vertebral column (Rijli et al., 1995; and this report). In the Hoxd-10 mutants the spinal cord defects occur at the lumbar level and influence the generation of the lumbar LMC, while the vertebral transformations occur at the sacral level. This is consistent with Hox gene expression domains in the neural tube being maintained at levels anterior to those in the prevertebral column (Dollé and Duboule, 1989; Haack and Gruss, 1993). However, at E9 the separation of Hoxd-10 expression in the neural tube and paraxial mesoderm is not as great (Zákány et al., 1997), allowing for possible overlap at these early time points. Unambiguous allocation of intrinsic or extrinsic specification of the spinal motor column may require the use of conditional mutations directed at selective inactivation of these genes in the neural tube or surrounding mesoderm.

The phenotypic consequences of targeted disruption of the Hoxd-10 locus appear to be limited to inactivation of this gene alone (i.e., there do not appear to be contributions from an altered function of adjacent genes). Two observations support this idea. First, there are no apparent overlaps between the reported mutant phenotypes of Hoxd-9, Hoxd-10 and Hoxd-11 mutant homozygotes (Davis and Capecchi, 1994; Favier et al., 1995; Favier et al., 1996). Second, no significant changes in Hoxd-9 and Hoxd-11 expression patterns were evident in Hoxd-10 homozygous mutants relative to wild-type embryos.

In summary, Hoxd-10 mutant mice show deficits in locomotion and adduction in the hindlimbs. The locomotor defects appear to be due to a malformed knee joint and a rotation of the tibia and fibula relative to the femur. However, defects are also apparent in the formation of the motor component of the nervous system in the lumbar region. The incomplete penetrance and variable expressivity suggest that other Hox genes, possibly Hoxa-10 and Hoxc-10, contribute to the formation and organization of these nerves. This hypothesis can be tested by analysis of mice carrying multiple Hox gene mutations.

Fig. 7. Transverse sections through the spinal column of wild-type (A), and Hoxd-10 mutant (B) newborn mice. The position of the lateral motor column is outlined. Higher magnification views of the same sections (C,D) show large motor neurons in the LMC in both animals (arrowheads). d, dorsal root ganglion. Scale bar for A and B, 100 µm and for C and D, 20 µm. Dorsal is up.

Fig. 8. Schematic representation of spinal segmental identity in wild-type (A) and Hoxd-10 mutant (B) animals. The L3 spinal segment normally projects nerve fibers to both the lumbar and sacral plexuses. Transformation of this segment to an L2 phenotype in mutant animals (B) results in its projection only to the lumbar plexus and a deletion of the L3 projection to the sacral plexus.
Deficits in the nervous system were apparent both in terms of robustness and organization. These deficits may reflect a common cause, since changes in specification of the L3 LMC would concomitantly reduce the allocation of nerves to the sacral plexus, or they may have separate causes, in which Hox genes would participate in decisions affecting both cell proliferation and cell specification. Irrespective of the mechanism, these studies suggest that Hox genes are intimately involved in the patterning of the nervous system, starting with the hindbrain and proceeding caudally along the entire axial column of the mouse.

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