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Renal agenesis in mice homozygous for a gene trap mutation in the gene encoding heparan sulfate 2-sulfotransferase

Simon L. Bullock,1 Judy M. Fletcher,2 Rosa S.P. Beddington,1,4 and Valerie A. Wilson1,3

1Laboratory of Mammalian Development, Medical Research Council (MRC) National Institute for Medical Research, The Ridgeway, London NW7 1AA, UK; 2MRC Human Genetics Unit, Western General Hospital, Edinburgh EH4 2XU, UK; 3Molecular Medicine Centre, Western General Hospital, Edinburgh EH4 2XU, UK

Heparan sulfate proteoglycans have been implicated in the presentation of a number of secreted signaling molecules to their signal-transducing receptors. We have characterized a gene trap mutation in the gene encoding a heparan sulfate biosynthetic enzyme, heparan sulfate 2-sulfotransferase (HS2ST). Transgenic mice were generated from embryonic stem cells harboring this insertion. lacZ reporter gene activity in heterozygous embryos demonstrates that the gene is expressed differentially during embryogenesis, presumably directing dynamic changes in heparan sulfate structure. Moreover, mice homozygous for the Hs2st gene trap allele die in the neonatal period, exhibiting bilateral renal agenesis and defects of the eye and the skeleton. Analysis of kidney development in Hs2st mutants reveals that the gene is not required for two early events—ureteric bud outgrowth from the Wolffian duct and initial induction of Pax-2 expression in the metanephric mesenchyme. It is required, however, for mesenchymal condensation around the ureteric bud and initiation of branching morphogenesis. Because 2-O-sulfation has been shown to influence the functional interactions of ligands with heparan sulfate in vitro, we discuss the possibility that the Hs2st mutant phenotype is a consequence of compromised interactions between growth factors and their signal-transducing receptors. These data provide the first genetic evidence that the regulated synthesis of differentially glycosylated proteoglycans can affect morphogenesis during vertebrate development.

[Key Words: Gene trap; heparan sulfate proteoglycan; sulfotransferase; kidney; induction]

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Heparan sulfate proteoglycans (HSPGs) are expressed at the surface of most animal cells, where they interact with a variety of proteins, including extracellular matrix components, growth factors, degradative enzymes, and protease inhibitors (for review, see Bernfield et al. 1992). Heparin-like molecules have been implicated in the presentation of a number of secreted signaling molecules, such as members of the fibroblast growth factor (FGF), Wingless/Wnt, transforming growth factor-β (TGF-β) and Hedgehog families (Rapraeger et al. 1991; Yayon et al. 1991; Lee et al. 1994; Reichsman et al. 1996; Ruppert et al. 1996; Binari et al. 1997), to their signal-transducing receptors. In some cell types, there is an obligate requirement for cell-surface HSPGs in the FGF-signaling pathway leading to the proposal that these molecules function as low-affinity receptors facilitating the interaction of ligands with high-affinity receptor tyrosine kinases (Klagsbrun and Baird 1991; Yayon et al. 1991; Schlessinger et al. 1995). In addition, HSPGs may bind growth factors in the extracellular matrix, altering their stability and modulating their effective concentration (Saksela and Rifkin 1990; Reichsman et al. 1996).

The specificity of HSPG–ligand interactions resides, at least in part, in the structure of the heparan sulfate glycosaminoglycan side chains, which vary in number, length, sequence composition, and sulfation pattern between cell type and developmental stage (Gallagher et al. 1986; Nurcombe et al. 1993; Kato et al. 1994). Therefore, the regulated synthesis of differentially glycosylated proteoglycans may represent an additional means to regulate cell–cell communication during development.

In a gene trap screen designed to identify genes important in mouse embryogenesis, we recovered an integration into the gene encoding heparan sulfate 2-sulfotransferase (HS2ST). This enzyme catalyzes the transfer of sulfate to position 2 of the iduronic acid component of heparan sulfate (Kobayashi et al. 1997). We show that Hs2st is expressed differentially during embryogenesis, presumably reflecting changes in the proteoglycan side chain structure. Moreover, mice homozygous for the gene trap mutation exhibit bilateral renal agenesis, resulting from a failure of ureteric bud branching and mesenchymal condensation, and defects of the eye and skeleton. These data provide the first genetic demonstration...
of heparan sulfate function in vertebrate embryonic development.

Results

ST125 represented one embryonic stem (ES) cell line selected from a gene trap screen designed to identify integrations into genes encoding secreted or transmembrane proteins expressed during early organogenesis (Skarnes et al. 1995; see Materials and Methods). Expression of the lacZ reporter was demonstrated in mid-gestation chimeras (data not shown) and subsequently adult chimeras were generated to transmit the integration through the germ line and establish a line of mice heterozygous for the gene trap allele (designated Hs2st\textsuperscript{TGNST125

Characterization of ST125 insertion site

5’ RACE was used to clone a portion of the trapped gene from the ST125 cell line, which proved to be the previously unidentified mouse homolog of hamster Hs2st. The protein product of this gene catalyzes the transfer of sulfate from 3’-phosphoadenosine 5’-phosphosulfate to position 2 of L-iduronic acid in heparan sulfate. The gene has been cloned recently from Chinese hamster ovary (CHO) cells (Kobayashi et al. 1997) and the protein was predicted to be localized to the Golgi in a type II membrane orientation.

A probe corresponding to the ST125 5’ RACE product was used to isolate a 2.1-kb mouse Hs2st clone from a 8.5-day postcoitum (d.p.c.) mouse embryo cDNA library. In addition, the same probe was used to isolate a Xenopus homolog of Hs2st from a gastrula-stage cDNA library using low-stringency screening. Sequence analysis demonstrated that the protein product of Hs2st is highly conserved in vertebrates, sharing 99% and 90% amino acid sequence identity with its hamster and Xenopus homologs, respectively (Fig. 1A). Furthermore, the mouse protein shares 58% amino acid sequence identity over a 245-amino-acid region with the predicted protein product of the Drosophila melanogaster S1 cDNA. The function of the S1 gene is unknown but it was identified as a candidate gene for the 5d mutation associated with the Segregation distorter meiotic drive system (Powers and Ganetzky 1991). Interestingly, one highly conserved block of sequence present in vertebrate HS2STs displayed significant homology with a large number of enzymes that catalyze the transfer of sulfate to a variety of molecules (Fig. 1B). This suggests that this region may be important for the catalytic mechanism although it is absent from some sulfotransferases, including heparan sulfate N-deacytlase/N-sulfotransferase (Hashimoto et al. 1992). It would, however, be present in the fusion protein generated by the gene trap insertion (Fig. 1A).

ST125 comprises a single gene trap insertion that disrupts the Hs2st transcript

FISH analysis to G-banded metaphase spreads of ST125 ES cells (Fig. 1C) revealed that Hs2st maps to a subtelomeric location on chromosome 3 (3H). Furthermore, only a single gene trap vector insertion site was evident, localized to chromosome 3, and it coincided precisely with the position of Hs2st (Fig. 1C). Molecular characterization of the mutant allele revealed a single copy of the gene trap vector and restriction mapping of genomic DNA and PCR-amplified flanking regions revealed no microdeletions or rearrangements within 10 kb on either side of the insertion site (data not shown). FISH analysis was also used to show that a genomic human HS2ST sequence hybridized to human chromosome 1p31 (Fig. 1D), a region showing synteny with mouse chromosome 3H (DeBry and Seldin 1996). The human HS2ST map position was refined further by radiation hybrid mapping to 3.87cR from D1S2167 on chromosome 1.

The junction in the 5’ RACE product of the endogenous mouse sequence and the gene trap vector splice acceptor sequence predicted that amino acid residues carboxy-terminal to position 196, including several regions with a high degree of evolutionary conservation, would not be encoded by the gene trap allele (Figs. 1A and 2A). Given the recessive nature of the Hs2st mutant phenotype (see below) the gene trap insertion is likely to cause a loss-of-function mutation; however, we cannot state categorically that it is a null allele.

Northern hybridization was performed on RNA isolated from wild-type, heterozygous, and homozygous embryos generated by intercrossing heterozygotes. Embryos were genotyped using a PCR-based assay that distinguished between the wild-type and the gene trap allele (see Materials and Methods). This analysis revealed that Hs2st transcripts were disrupted by the gene trap integration (Fig. 2B). Using a probe specific to the 5’ RACE product, a transcript of 2.4 kb was detected in RNA derived from wild-type, but not from homozygous, mutant embryos. In the latter, only a 5.6-kb band (the size predicted for the gene trap fusion mRNA) was observed and no wild-type transcript could be detected. Heterozygotes were found to possess the wild-type and fusion transcripts in equal amounts. Low levels of a 4-kb transcript, possibly representing an alternative splice product, were present in wild-type RNA but were undetectable in homozygous mutant RNA, indicating that this transcript was also disrupted by the transgene insertion.

Embryonic expression pattern of Hs2st

The lacZ expression pattern in heterozygous embryos was examined at various developmental stages (Fig. 3). At 7.5 d.p.c., all three germ layers expressed lacZ, although there appeared to be elevated expression in the embryonic ectoderm and the node (Fig. 3A). Widespread expression persisted at 8.5 d.p.c., although β-gal activity was clearly stronger in rhombomeres 2 and 4 and branchial arches 1 and 2 (which are populated by neural crest from these rhombomeres) (Fig. 3B,C). At 10.5 d.p.c., the dorsal and ventral aspects of the neural tube, brain, and midbrain-hindbrain junction showed the most intense staining (Fig. 3D,E). A day later in development, elevated
β-gal activity was found in the floor plate (Fig. 3F,G) and the sclerotome (data not shown). At 12.5 d.p.c., both the floor plate (data not shown) and the roofplate exhibited strong lacZ staining as did the mesenchyme of the limb and of the developing whisker follicles (Fig. 3H). At 13.5 d.p.c., lacZ expression predominated in embryonic mesenchyme, especially at sites of epithelial–mesenchymal interactions such as the developing teeth (data not shown) and whisker follicles (Fig. 3I). Strong staining was also apparent in the perichondria of the cartilaginous skeleton (Fig. 3I), an important site for the regulation of skeletal differentiation (Vortkamp et al. 1996). β-gal activity in Hs2st+/− embryos faithfully reflected the endogenous gene expression pattern as revealed by in situ hybridization using an antisense Hs2st probe (Fig. 3B,D,F; data not shown).

Hs2st is required for metanephric development

Genotype analysis of prenatal litters between 8.5 and 15.5 d.p.c. indicated that there was no significant death of homozygotes before the latter stages of gestation [73 homozygotes (27%) identified out of a total of 273 embryos]. Homozygous mutants, however, were either stillborn or died within 24 hr of birth and showed bilateral renal agenesis. This completely penetrant phenotype (Table 1; Fig. 4) was probably responsible for the neonatal lethality. No abnormalities were observed in other parts of the urogenital system (Fig. 4) or in other internal organs. All heterozygotes and wild types examined had overtly normal kidneys.

In amniotes, the permanent (or metanephric) kidney is formed by the interaction of two mesodermal deriv-
tives, the epithelial ureteric bud and the metanephric mesenchyme. At 10.75 d.p.c. in the mouse, the ureteric bud emerges from the Wolffian (or nephric) duct, thereby contacting the adjacent metanephric mesenchyme. The metanephric mesenchyme induces the ureteric bud to elongate and branch repeatedly, ultimately forming the collecting duct system and, at the same time, the ureteric bud tips prevent the metanephric mesenchyme from undergoing apoptosis and induce it to condense, epithelialize, and differentiate into secretory nephrons (Grobstein 1953; for review, see Bard et al. 1996; Vainio and Müller 1997).

To investigate the embryological basis of the renal defect, the developing kidneys of Hs2st−/− mutants were compared with those of normal littermates. At 11.5 d.p.c., all wild-type and heterozygous kidney rudiments (n = 85, where n = number of kidneys examined) contained a ureteric bud that had bifurcated once or, less frequently, twice, and was surrounded by a mantle of mesenchymal condensation (Fig. 4C,E). In contrast, Hs2st−/− kidney rudiments (n = 48) contained an unbranched ureteric bud that was not surrounded by condensed metanephric mesenchyme (Fig. 4D,F). In homozygous mutants at 12.5 d.p.c., no branching of the ureteric bud had occurred and there were no overt signs of mesenchymal condensation (data not shown; n = 44), demonstrating that renal development was arrested rather than delayed. Therefore, Hs2st is not required for the initial outgrowth of the ureteric bud from the Wolffian duct but rather for ureteric bud branching and mesenchymal condensation. The presence of relatively long unbranched ureters in neonatal Hs2st−/− mutants (Fig. 4B) shows that the gene is not required for the directional growth of this tissue.

Expression of Hs2st during kidney development

In heterozygotes at 10.75 d.p.c., weak β-gal activity was evident in the metanephric mesenchyme, Wolffian duct, and emergent ureteric bud (Fig. 5A). At 11.5 d.p.c., reverse section through an 11.5-d.p.c. embryo. Strong expression is exhibited by the floor plate (arrow), and limb mesenchyme (black arrowhead) express high levels of lacZ. (I) Transverse section through a 13.5-d.p.c. embryo showing vibrissa follicle expression localized to the mesenchyme (arrowheads) and the developing hairshaft (arrow). (F,G) Transverse section through an 11.5-d.p.c. embryo. Scale bar in J: 125 µm (A); 425 µm (B,C); 1000 µm (D,E); 150 µm (F,G); 1800 µm (H); 500 µm (I); 375 µm (J).
porter gene expression was almost undetectable in the ureteric bud epithelium; in contrast, the metanephric mesenchyme expressed high levels of lacZ (Fig. 5B). To follow Hs2st expression during subsequent metanephric development, 11.5-d.p.c. Hs2st<sup>+/−</sup> kidney rudiments were cultured in vitro (Fig. 5C,D). After 24 hr of culture, lacZ-expressing cells were located at some distance from the ureteric bud tip. It has been observed in vitro that, once induced, metanephric mesenchymal cells move away from the inducing tissue, thereby facilitating the induction of further mesenchyme (Saxén and Karkinen-Jääskeläinen 1975). Because mutants one day earlier in development (11.5 d.p.c.) did have Pax-2-expressing mesenchymal cells adjacent to the unbranched ureteric bud (Fig. 6B), it is likely that the paucity of cells expressing this protein at 12.5 d.p.c., and their remoteness from the ureteric bud, reflects an early transient signal between the ureteric bud and metanephric mesenchyme.

The Ret receptor tyrosine kinase and its ligand glial cell line-derived neurotrophic factor (GDNF) are components of a signaling pathway required for ureteric bud outgrowth from the Wolffian duct and collecting duct morphogenesis (Schuchardt et al. 1994, 1996; Durbec et al. 1996; Moore et al. 1996; Pichel et al. 1996; Sánchez et

| Phenotype | Bilateral renal agenesis | Cleft palate | Extra postaxial digit<sup>a</sup> of one forelimb | Extra postaxial digit<sup>a</sup> of both forelimbs |
|-----------|--------------------------|-------------|--------------------------|--------------------------|
| No. with defect/ no. examined | 29/29 | 12/29 | 16*<sup>a</sup>/29 | 5/29 |

<sup>a</sup> Of which, 13 had an extra digit on the right forelimb.

Expression of molecular markers for metanephric development in Hs2st mutants

Whole-mount in situ hybridization and immunohistochemistry was used to assay the expression of well-characterized markers of metanephric development in mutant embryos (Fig. 6). For each marker at each stage, a minimum of five and, in most cases, more than 12 mutant kidneys where compared with a similar number of heterozygous and wild-type litters. The transcription factor Pax-2, as well as being expressed in the Wolffian duct, mesonephric tubules, and the ureteric bud epithelium, is induced in the metanephric mesenchyme following interaction with the ureteric bud (Dressler et al. 1990). In Hs2st<sup>+/−</sup> urogenital systems at 11.5 and 12.5 d.p.c., Pax-2 expression in the Wolffian duct and the mesonephric tubules was identical to that seen in the normal situation. In the kidneys, however, only an unbranched Pax-2-expressing ureteric bud was present. At 12.5 d.p.c., wild-type and heterozygous kidneys contained a number of Pax-2-expressing mesenchymal aggregates surrounding branching ureteric bud tips (Fig. 6C). In contrast, in homozygous mutant mesenchyme there was only a relatively small patch of Pax-2-expressing cells, which were located at some distance from the ureteric bud tip. It has been observed in vitro
Role of Hs2st in kidney development

In homozygous, as in wild-type and heterozygous embryos, Gdnf transcripts were present in the uninduced metanephric mesenchyme at 10.75 d.p.c. (data not shown). Later, Gdnf is normally highly expressed in mesenchymal cells surrounding the ureteric bud tips (Fig. 6D,F; Durbec et al. 1996). In Hs2st−/− kidneys, Gdnf expression was attenuated at 11.5 d.p.c. (Fig. 6E) and undetectable by 12.5 d.p.c. (Fig. 6F). In homozygous, heterozygous, and wild-type embryos, c-ret was expressed along the length of the Wolffian duct and in the emergent ureteric bud at 10.75 d.p.c. (data not shown). As the ureteric bud elongated and branched in homozygous and wild-type embryos, c-ret expression resolved to the tips of branching ureteric buds (Fig. 6G). In Hs2st−/− embryos the unbranched ureteric bud tips at 11.5 d.p.c. showed only low levels of c-ret expression (Fig. 6H) and this expression had disappeared 24 hr later (Fig. 6I). Therefore, following invasion of the metanephric mesenchyme by the ureteric bud, Hs2st appears to be required for maintenance of Gdnf expression and the localization of c-ret expression to the ureteric bud tip. Likewise, expression of Wnt11 in ureteric bud tips, which is compromised in vitro by inhibitors of glycosaminoglycan synthesis (Kispert et al. 1996), was not detected within the ureteric bud tip in Hs2st−/− mutants following invasion of the metanephric mesenchyme (Fig. 6K). The maintenance of apparently normal levels of expression within the ureter of markers such as Sox9 (Fig. 6L; Kent et al. 1996) and Pax-2 (see above) in mutants indicates that Hs2st is not required for ureter identity but for maintenance of expression of genes associated specifically with branching ureteric bud tips.

Skeletal and eye defects in Hs2st mutants

As well as the kidney phenotype, other defects were apparent in homozygous mutants (Table 1; Fig. 7). There was a global increase in bone mineralization in homozygotes (Fig. 7B,C) and ectopic ossification of the sternum (six of seven mutants examined; Fig. 7E) and fusions of the cervical vertebrae (two of seven mutants examined; data not shown) occurred. In addition, a high frequency of mutants exhibited post-axial polydactyly and interestingly this predominated in the right forelimb. A cleft of the secondary palate was also a frequent occurrence (Table 1). Bilateral coloboma of the iris, resulting from a perturbation of pigmented retinal epithelium differentiation, was seen in 14/14 homozygotes examined (Fig. 7E). Histological sections revealed that mutants showed signs of retardation in eye development (data not shown). Furthermore, six of eight mutant animals had cataracts (data not shown). The expression pattern of Hs2st is consistent with a role in the development of the skeleton and the eye. The gene is highly expressed in the perichondria during skeletogenesis (Fig. 3) and at the junction of the neural and pigmented retinal epithelia during development of the eye (data not shown). No gross anatomical defects were evident in the brain and spinal cord, although a more detailed analysis may reveal subtle defects.

Discussion

Attribution of the mutant phenotype to the gene trap insertion in Hs2st

We have characterized a gene trap mutation in the gene encoding heparan sulfate 2-sulfotransferase. We find that mice homozygous for this mutation die in the neonatal period and invariably exhibit bilateral renal agenesis and defects of the eye and skeleton. Cleft palate and polydactyly also occur with incomplete penetrance.

The consistent segregation of the mutant phenotype with homozygosity for the unique gene trap insertion in Hs2st strongly suggests that it is disruption of this gene that is responsible for the lesions observed. Moreover, Hs2st is expressed in all tissues showing overt abnormalities and no intact Hs2st transcripts are de-
Figure 6. Expression of molecular markers for metanephric development in Hs2st mutants in dissected 11.5-d.p.c. kidney rudiments (A,B,D,E,G,H) and 12.5-d.p.c. urogenital regions (C,F,I,L). Heterozygous (H→) or wild-type (H+) (A,D,G,J and left-hand specimen in C,F,I,L) and homozygous mutant (→−) (B,E,H,K and right-hand specimen in C,F,I,L) specimens subjected to immunostaining with anti-Pax-2 antibody (A–C) or whole-mount in situ hybridization with probes specific to Gdnf (D–F), c-ret (G–I), Wnt11 (J,K), or Sox9 (L). Kidney rudiments are labeled with black arrowheads in C, F, I, and L. (A) At 11.5 d.p.c., Pax-2-positive cells are abundant in the metanephric mesenchyme surrounding the ureteric bud, which also expresses Pax-2 and has bifurcated twice. (B) In mutants at 11.5 d.p.c., Pax-2 expression is detected in the unbranched ureter (black arrow) and in the mesenchyme concentrated in a small population of cells at a distance from the ureteric bud tip. Note the “stream” of Pax-2-positive cells adjacent to the bud tip (white arrow). (C) At 12.5 d.p.c., Pax-2 expression is localized to the mesonephric tubules (mt), Wolffian ducts (wd), and ureteric epithelium in mutant and heterozygous or wild-type urogenital systems. Mutants have unbranched Pax-2-expressing ureters (white arrowheads) and possess few Pax-2 positive cells in the mesenchyme (black arrowheads). (D) At 11.5 d.p.c., strong Gdnf expression is detected in the mesenchyme surrounding the bifurcated ureteric bud. (E) In mutants, Gdnf expression in mesenchyme adjacent to the unbranched ureteric bud is attenuated (black arrow). (F) In 12.5-d.p.c. Hs2st mutant kidney rudiments, Gdnf transcripts are no longer detected. (G) At 11.5 d.p.c., c-ret transcripts are detectable throughout the ureteric bud (black arrow) and are abundant in the ureteric bud tips. (H) In mutants at 11.5 d.p.c., c-ret expression in the ureteric bud is comparable to wild-type or heterozygous kidneys, but strong expression is not detected in the unbranched ureteric bud tip (black arrow). (I) At 12.5 d.p.c., c-ret transcripts are abundant in the bifurcated ureteric bud tips of the heterozygous or wild-type kidney and are no longer detectable in mutant kidney rudiments. (J) At 11.5 d.p.c., Wnt11 is expressed in ureteric bud tips and in mesenchyme proximal to the Wolffian duct (arrow). (K) In mutants, Wnt11 expression is undetectable in the unbranched ureteric bud (position highlighted by black dots). (L) In heterozygous or wild-type kidneys at 12.5 d.p.c., Sox9 transcripts are detected in the ureteric system. In mutants, expression persists at apparently normal levels in the unbranched ureter (white arrowheads). Black arrows show expression of Sox9 in the male gonads (the mutant urogenital region is female and, therefore, does not express Sox9 in the gonad) (Kent et al. 1996; Morais da Silva et al. 1996). Scale bar in L: 100 µm (A,B,D,E,G,H,J,K); 800 µm (C,F,I,L).
in vitro studies have shown that 2-O-sulfation in particular is essential if heparan sulfate is to interact with certain growth factors such as FGF-2 (basic FGF; Turnbull et al. 1992; Maccarana et al. 1993; Bai and Esko 1996). This modification appears not to be required for the interaction of hepatocyte growth factor with its receptor (Lyon et al. 1994). Therefore, modulating the levels of 2-O-sulfation may confer an additional level of specificity on some, but not all, growth factor–receptor interactions. HSPG structure varies extensively between tissues during development (Bernfield et al. 1992), and one means of achieving this diversity is illustrated here by the regulated expression of Hs2st. Several embryonic signaling centres, such as the node and floor plate, show elevated expression of the gene, and there is a clear example of its apparent down-regulation on epithelialization of the metanephric mesenchyme.

There is increasing evidence that HSPGs can modulate the activity of growth factors through a number of mechanisms, including facilitating their dimerization or altering their effective concentration by acting as low-affinity receptors (Schlessinger et al. 1995). Recently, genetic studies in Drosophila have demonstrated a role for heparin-like glycosaminoglycans in the reception of the Wingless (Wg) signal (Binari et al. 1997; Hӓcker et al. 1997; Haerry et al. 1997). Embryos that are either deficient in an enzyme catalyzing the production of a glycosaminoglycan precursor or that have been treated with an enzyme that specifically degrades heparan glycosaminoglycans, exhibit phenotypes reminiscent of loss-of-Wingless signaling. It has also been suggested that changes in HSPG expression may influence the distance over which a morphogen such as Wg can act; restricting it to a short-range signal in some circumstances but allowing it to serve as a more widespread gradient morphogen in others (Hӓcker et al. 1997). Therefore, it is likely that modulating the sulfation status of HSPGs will influence the availability of critical ligands and it is tempting to speculate that the Hs2st−/− phenotype is due, at least in part, to the suboptimal presentation of signaling molecules to their receptors.

Consequences of the Hs2st mutation

Previous experiments on early kidney development in vitro showed that the somewhat indiscriminate inhibition of de novo glycosaminoglycan sulfation by chlorate reversibly blocks ureteric bud growth and branching but does not affect nephrogenesis (Davies et al. 1995). The phenotype observed in Hs2st−/− mutants, however, indicates an earlier requirement for 2-O-sulfation in the initial condensation of the metanephric mesenchyme. During collecting duct morphogenesis, Hs2st does not seem to be required for the initial outgrowth of the ureteric bud from the Wolffian duct but rather for its subsequent branching following contact with the metanephric mesenchyme. This suggests that it is required downstream of genes such as Wt1, Pax2, c-ret, and Gdnf (Kreidberg et al. 1993; Schuchardt et al. 1994, 1996; Torres et al. 1995; Moore et al. 1996; Pichel et al. 1996; Sanchez et al. 1996), which are implicated in ureteric bud outgrowth, but upstream of other genes such as Wnt4 and Bmp7 that are necessary for normal nephrogenesis (Stark et al. 1994; Dudley et al. 1995; Luo et al. 1995).

Interestingly, the metanephric phenotype of mice homozygous for a mutation in the homeobox gene Emx2, which is expressed in the ureteric bud (Miyamoto et al. 1997), is strikingly similar to that of Hs2st−/−. In both mutants, kidney development is arrested at the same stage and both show a very consistent phenotype, unlike the variability seen in Gdnf and c-ret mutants, which ranges from complete absence of ureteric bud to the development of small kidneys (Schuchardt et al. 1994; Moore et al. 1996). This suggests that there may be additional signaling mechanisms involving molecules other than Ret and GDNF that are regulated by Emx2 in the ureteric bud and require the mediation of HSPGs.

FGF-2 can substitute for the ureteric bud to induce mesenchymal condensation and prevent apoptosis in cultured metanephric mesenchyme (Perantoni et al. 1995). Interestingly, FGF-2 stimulates growth and inhibits differentiation of chondrocytes (Suzuki 1992) and this effect is likely to depend on the presence of sulfated glycosaminoglycans as their removal by chlorate promotes chondrocyte differentiation (Chintala et al. 1995). Therefore, impaired FGF-2 signaling could explain many of the defects observed in Hs2st−/− mutants. Mice homozygous for a null mutation in Fg2, however, show no overt defects in kidney morphogenesis or skeletogenesis (R.
Although it appears to be required later than Hs2st, the metanephric mesenchyme of Wnt4 served in developing kidney (Finch et al. 1995), the phenotype observed in Hs2st−/− embryos may be attributable to perturbation of more than one FGF-signaling pathway. Cells expressing a vertebrate wg homolog Wnt1 can induce tubulogenesis in isolated metanephric mesenchyme (Herzlinger et al. 1994). Wnt1, however, is not itself expressed in the embryonic kidney, indicating that other Wnt family members may normally be responsible for early inductive events in metanephric development. The expression of Wnt4 is consistent with such a role although it appears to be required later than Hs2st. The metanephric mesenchymal of Wnt4−/− mutants does condense but fails to transform into an epithelium (Stark et al. 1994). Wnt11 is expressed at the ureteric bud tips and therefore it may contribute to mesenchymal induction (Kispert et al. 1996). If so, its signaling, like that of Wg in Drosophila, may be compromised in the absence of appropriately sulfated HSPGs.

It is also possible that failure to undergo changes in adhesion could contribute to the Hs2st−/− phenotype. HSPGs can interact with a variety of matrix components via their heparan-sulfate chains, including certain collagens, tenascin, thrombospondin, and fibronectin and can be concentrated in focal adhesions (for review, see Bernfield et al. 1992; Woods and Couchman 1994). Clearly metanephric mesenchymal condensation requires alterations in adhesive properties and defective sulfation might prevent such changes. The transmembrane HSPG Syndecan-1 can regulate cell adhesion and cell morphology (Leppä et al. 1992) and is up-regulated dramatically in the metanephric mesenchyme on induction (Vainio et al. 1989). Therefore, it is possible that its function is compromised by the Hs2st mutation. At present, the potential roles of Hs2st in modulating signaling and adhesive properties cannot be distinguished and it is possible that both are important in kidney development.

Materials and methods

Production of gene trap ES cells

Electroporation and selection of CGR8 (129/Ola) ES cells was performed as described (Skarnes et al. 1995). The vector pGTm6 was a gift from W. Skarnes and forms fusion proteins in a different open reading frame (ORF) to the parent plasmid pGTL8TM (Skarnes et al. 1995). The vector contains a splice acceptor sequence upstream of CD4 transmembrane domain sequence, a lacZ-neomycin phosphotransferase fusion (β-gal) and a polyadenylation signal such that integrations into introns of transcriptionally active genes will produce a β-galactosidase (β-gal) fusion protein that contains endogenous sequence amino-terminal to the site of insertion. The inclusion of a transmembrane domain-encoding region in the vector enriches for insertions downstream of secretory signal sequences or type II transmembrane domains because β-gal activity is only preserved when the enzyme is retained in the cytoplasm (Skarnes et al. 1995).

The vector was electroporated into 1 × 10^8 CGR8 ES cells. Colonies (149) resistant to G418 were recovered and 39 (26%) of these exhibited β-gal activity. In 16 of these clones, β-gal activity showed a subcellular localization typical of fusion with a secreted or transmembrane protein (Skarnes et al. 1995). Chimeras were generated from these cells and the ES cell line ST125 was one integration selected for further analysis on the basis that lacZ reporter gene expression was evident in 8.5- and 9.5-d.p.c. chimeric embryos.

Rapid amplification of cDNA ends

Total RNA was isolated from the ST125 cell line using the guanidine thiocyanate method (Chomczynski and Sacchi 1987). cDNA was synthesized from 5 µg of total RNA and 5′ rapid amplification of cDNA ends (RACE) performed using Marathon cDNA amplification kit (Clontech) according to manufacturer's instructions. Following adaptor ligation, 35 cycles of PCR were conducted using Marathon adaptor primer 1 (5′-CCATCCTAATACGACCTCACTATAGGG-3′) and a primer specific to the CD4 transmembrane domain region (5′-TGAGGATTGACGCTGAGCGGC-3′) and a primer specific to the CD4 transmembrane domain region (5′-TGAGGATTGACGCTGAGCGGC-3′) and a primer specific to the CD4 transmembrane domain region (5′-TGAGGATTGACGCTGAGCGGC-3′) and a primer specific to the CD4 transmembrane domain region (5′-TGAGGATTGACGCTGAGCGGC-3′). A 1/50 dilution of the PCR products was subjected to 30 cycles of nested PCR (adapter primer 2 (5′-ACTCCTATAGGGCTGAGCGGC-3′), CD4 primer 2 (5′-AGAAGATCCCTCTGCAGACACCC-3′)) and gene trap RACE products verified using Southern blotting and hybridization with an end-labeled oligonucleotide probe specific for the En-2 exon sequence in the gene trap vector (5′-GTCCAGTGCTCCGAAAAACCAAGAAGAAGAAGC-3′) using standard procedures. Following size selection of hybridizing products (Quiex II Gel Extraction Kit) and subcloning, 10-20 differentiated ES cells were isolated per vector. RT-PCR using standard procedures.

Isolation of cDNA and genomic clones

PCR employing primers specific to the ST125 5′ RACE product (5′-GAGATGGAAGGAAATATGGG-3′ and 5′-GAGAAGGAAATATGGG-3′) was used to amplify a 319-bp probe, termed ST125A, which was used to screen a 8.5-d.p.c. mouse embryo cDNA λGT10 library (Fahrner et al. 1987) using standard procedures. Two independent clones representing full-length Hs2st cDNAs were isolated. The same probe was used to
isolate bacteriophage λ clones harboring genomic DNA inserts corresponding to the mouse Hs2st gene from a CGR8 ES cell λKO genomic library (Nehls et al. 1994) and the human Hs2st gene from a human fibrosarcoma cell line genomic λDASHII library [Stratagene; hybridization: 0.5 M phosphate buffer (pH 7.2), 7% SDS, 1 mM EDTA at 50°C for 16 hr; Washes: 2× SSC/0.1% SDS at 50°C, 3 times]. Xenopus laevis Hs2st cDNAs were isolated from an amplified gastrula cDNA library in Uni- ZAP XR (Cho et al. 1991) using low-stringency screening. Filters were hybridized with the mouse ST125A probe in 30% formamide, 5× SSC, 5× Denhardt’s solution, 0.02% NaPP, 0.1% SDS, 0.1 mg/ml of yeast tRNA, 0.03 M phosphate buffer (pH 6.5) at 37°C and washed in 2× SSC/0.1% SDS 3 times at 37°C.

**FISH and karyotyping**

Metaphase chromosome spreads were made from ST125 ES cells or human blood lymphocyte preparations using standard procedures (Robertson 1987). To determine the localization of Hs2st and the gene trap insertion, Giemsa banding was carried out (Kleven et al. 1991) and 15 spreads from each of two slides were recorded as digital images using Quips Capture software (Vysis) along with their position on the slide. One slide was hybridized to a 12-kb mouse genomic Hs2st probe, and the other to a gene trap vector-specific probe. Chromosomes showing hybridization on the captured spreads were compared with the corresponding Giemsa-banded image. The chromosome was identified by karyotyping using Quips Karyotyper software (Vysis). This analysis additionally confirmed the absence of cytogenetically visible chromosome anomalies. A similar procedure was used to identify the chromosomal localization of human HS2ST, using a 16-kb human genomic probe. To show colocalization of the gene trap vector insertion with mouse Hs2st in ST125 cells, double FISH was carried out (Fantes et al. 1995) using a digoxigenin-labeled Hs2st probe and a biotinylated gene trap vector probe.

**Radiation hybrid mapping**

An intron-specific portion of human HS2ST was subcloned and the sequence used to design primers that specifically amplified a 101-bp PCR fragment from human, but not hamster DNA (5’-TGTGAACCATGTTAGCCA-3’, 5’-TGAATACTAAAAAGA-GCTG-3’). These were used to screen the GeneBridge 4 human–hamster radiation hybrid panel (Gyapay et al. 1996). Data were submitted to the Human Genome Mapping Project and the human–hamster radiation hybrid panel (Gyapay et al. 1996). These were used to screen the GeneBridge 4 human–hamster radiation hybrid panel (Gyapay et al. 1996). Data were submitted to the Human Genome Mapping Project and mapped using the Whitehead Institute statistical program RH-MAPPER.

**ES cell chimeras and transgenic mice**

Mice were maintained on a 10-hr light, 14-hr dark cycle. Noon on the day of finding a vaginal plug was designated 0.5 d.p.c. Germ-line mice were generated by injection of ST125 ES cells into C57BL6 blastocysts followed by transfer to a foster mother (Hogan et al. 1994). Chimeric male mice were mated to C57BL6 females and F1 progeny analyzed for the presence of the transgene. Heterozygous animals were backcrossed onto a C57/BL6 background and heterozygous siblings intercrossed.

**Recovery of embryos and lacZ expression pattern analysis**

Embryos were dissected from the uterus in M2 medium containing 10% fetal bovine serum (Advanced Protein Products) as described (Hogan et al. 1994) and X-gal-stained overnight according to the method of Beddington et al. (1989). In all cases, transgenic samples were incubated with stage-matched wild-type tissues to control against endogenous β-gal activity. Ex vivo kidney cultures were performed as described before staining (Kispert et al. 1996).

**Genotyping procedures**

Hs2st KO mice could be distinguished from wild-type littermates by X-gal staining of tail biopsies. To establish an allele-specific genotyping assay, Hs2st genomic sequences flanking the vector insertion site were amplified from lacZ-positive genomic DNA (High-fidelity PCR system; Boehringer Mannheim), cloned and sequenced. Subsequently, PCR primers were designed such that a common primer, specific to sequences 5’ to the insertion (5’-ATCAAGTAAATTGCCTAGGC-3’) was used in conjunction with a primer specific to sequences 3’ to the insertion (5’-GGGAAGAATTCACCCCAACA-3’) and a primer that recognizes sequences in the gene trap vector (5’-TACCTAGTCAGTGACTCA-3’) to generate 344- and 174-bp products specific to the mutant and wild-type alleles, respectively. Genomic DNA (500 ng) was denatured at 94°C for 5 min and immediately transferred to ice before amplification in a 20-µl reaction volume (0.6 µM each primer, 50 mM KCl, 10 mM Tris-HCl (pH 8.3), 1 mM MgCl2, 100 µM dNTPs, 1 unit of AmpliTaq (Perkin-Elmer)), to 30 cycles (94°C for 20 sec, 60°C for 30 sec, 72°C for 30 sec) and a single elongation step (72°C for 5 min).

**Northern hybridization**

Total RNA was isolated from individual 15.5-d.p.c. embryos recovered from an intercross of heterozygous Hs2st KO parents using the method of Chomczynski and Sacchi (1987), following harvesting of yolk sacs for DNA genotyping. Total RNA (10 µg) from embryos of each genotype was electrophoresed in a 1.5% formaldehyde-agarose gel as described (Sambrook et al. 1989) and transferred to Hybond N membrane (Amersham) according to manufacturer’s instructions. The membrane was hybridized at 65°C with the ST125A probe in 5× Denhardt’s, 4× SSC, 50 mM NaH2PO4, 10% dextran sulfate and washed in 0.5× SSC/0.1% SDS and then 0.2× SSC/0.1% SDS at 65°C before autoradiography. The membrane was reprobed with a β-actin cDNA probe (Harrison et al. 1995).

**Whole-mount in situ hybridization and immunohistochemistry**

Whole-mount in situ hybridization was performed as described (Wilkinson 1992) with two modifications. Urogenital ridges were treated for 30 min in 20 µg/ml proteinase K (as opposed to 15 min in 10 µg/ml for embryos and dissected kidneys) and, in all cases, embryo powder was omitted from the procedure. Digoxigenin-labeled antisense riboprobes were generated as described (Sambrook et al. 1989) and a primer that recognizes sequences in the gene trap vector (5’-ACCTAGTCAGTGACTCA-3’) to generate 344- and 174-bp products specific to the mutant and wild-type alleles, respectively. Genomic DNA (500 ng) was denatured at 94°C for 5 min and immediately transferred to ice before amplification in a 20-µl reaction volume (0.6 µM each primer, 50 mM KCl, 10 mM Tris-HCl (pH 8.3), 1 mM MgCl2, 100 µM dNTPs, 1 unit of AmpliTaq (Perkin-Elmer)), to 30 cycles (94°C for 20 sec, 60°C for 30 sec, 72°C for 30 sec) and a single elongation step (72°C for 5 min).

Whole-mount in situ hybridization and immunohistochemistry were performed with Wnt11 (Durbec et al. 1996), and c-ret (Durbec et al. 1996), and Sox9 (Morais da Silva et al. 1996). The Hs2st antisense riboprobe was transcribed from a cDNA clone representing amino acids 8–330.

For whole-mount immunohistochemistry, urogenital ridges were treated for 4× PFA for 15 min, washed once in PBS containing 1% Triton X-100 and several times in PBS containing 0.1% Triton X-100. Following an overnight incubation at 4°C in a 1,400 dilution of anti-Pax-2 IgG (Dressler and Douglass 1992) in PBS, Pax-2 expression was detected using a horseradish peroxidase (HRP)-conjugated goat anti-rabbit IgG secondary antibody (Sigma Immunochemicals) as described (Kispert and Herman 1994).
Skeletal preparations and histochemistry

Skeletal preparations of newborn mice were made using an alcian blue-alizarin red method as described (Kessel and Gruss 1991) and hematoxylin and eosin staining was performed according to Kaufman (1992).

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