Specific ablation of the nidogen-binding site in the laminin γ1 chain interferes with kidney and lung development

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SUMMARY
Basement membrane assembly is of crucial importance in the development and function of tissues and during embryogenesis. Nidogen 1 was thought to be central in the assembly processes, connecting the networks formed by collagen type IV and laminins, however, targeted inactivation of nidogen 1 resulted in no obvious phenotype. We have now selectively deleted the sequence coding for the 56 amino acid nidogen-binding site, γIII114, within the Lamc1 gene by gene targeting. Here, we show that mice homozygous for the deletion die immediately after birth, showing renal agenesis and impaired lung development. These developmental defects were attributed to locally restricted ruptures in the basement membrane of the elongating Wolffian duct and of alveolar sacculi. These data demonstrate that an interaction between two basement membrane proteins is required for early kidney morphogenesis in vivo.

Key words: Basement membrane, Wolffian duct, Kidney, Lung, Morphogenesis, Mouse

INTRODUCTION
Basement membranes are specialized extracellular matrices that play fundamental roles in tissue development and function. These thin sheet-like structures are produced through complex interactions between the major components laminins, collagen IV, the heparan sulfate proteoglycan perlecan and nidogen (Mayer and Timpl, 1993). Laminins are heterotrimeric cross-shaped multidomain molecules composed of three genetically distinct chains (α, β and γ). To date, 14 different laminin isoforms have been identified and the diversity is mainly due to the existence of five different α chains (Miner and Patton, 1999). Conversely, among the three known γ chains, the laminin γ1 chain shows a ubiquitous expression pattern and is found in all laminin isoforms, with the exception of laminin 5 and laminins 12-14 containing the γ2 and γ3 chain, respectively (Miner and Patton, 1999; Koch et al., 1999; Libby et al., 2000). Purification of laminin 1 under non-denaturing conditions demonstrated that nidogen 1 forms a stable complex with laminin (Paulsson et al., 1987). In addition, nidogen 1 has been shown to bind to most of the currently known basement membrane proteins, including perlecan and collagen IV (Timpl and Brown, 1996). The finding, however, that nidogen 1 can mediate the formation of ternary complexes between laminin and collagen IV in vitro (Fox et al., 1991), led to the hypothesis that it is crucial for basement membrane assembly, by connecting the major networks formed by laminins and collagen IV.

The precise mechanisms how basement membranes are assembled and how the biological function of the proteins is maintained within the basement membrane in vivo, are still unclear. In particular, the differential sites of synthesis for nidogen and laminin chains in mesenchymal and epithelial tissues, respectively, (Thomas and Dziadek, 1993; Ekblom et al., 1994) argues for the presence of other factors involved in the assembly processes. Recent results obtained by targeted deletion of cell-surface receptors, such as β1 integrins and dystroglycan showed disruption of basement membrane structures (Kreidberg et al., 1996; Williamson et al., 1997; DiPersio et al., 1997; Sasaki et al., 1998; Henry and Campbell, 1998) and emphasize a role for transmembrane complexes in coordinating the spatial and temporal local concentrations of proteins at the sites of basement membrane formation.

It is widely believed that basement membranes serve as both structural barriers and as a substrate for cellular interactions. The genetic inactivation of most of the major components has demonstrated that each of the proteins serves specific functions. Though all mutations interfere at specific stages with basement membrane integrity, the underlying mechanisms most probably differ. Mice deficient for perlecan develop normally before they die of heart failure at 10.5 day post coitum (dpc), because of basement membrane instability caused by mechanical stress.
was achieved within a unique 400 bp Apal/NheI fragment using 5'-GATCGGGCGTCGAGCTGCAAAAGTGTAGTCAGGGCA GACTCGCTACG and 5'-ACAGAGAGCTGTCATGCAATAAAA- CACATTGCC, and verified by sequencing. After subcloning the mutated Apal/NheI fragment into the HindIII/BamHI genomic clone, the HSV-TKNeo cassette flanked by two loxP sites was inserted into the NheI site in the following intron. To identify Cre recombination, an additional EcoRV restriction site was introduced after the second loxP site. The mutated clone was then extended by a 3’-following 5.2 kb BamHI genomic fragment.

Generation of hetero- and homozygous ES cells

R1 cells were cultured and transfected with the NovI linearized targeting vector as described (Mayer et al., 1997). Genomic DNA from G418 resistant clones were screened by Southern hybridization after EcoRV restriction with a 0.5 kb HindIII/EcoRV fragment located upstream of the targeting vector. Positive clones were analyzed for additional random insertion using the neomycin gene as probe. Two clones were expanded and transiently transfected with Cre recombinase under the control of the PGK promoter. Deletion of the selection cassette was verified after EcoRV restriction using the same probe as described above. To obtain homozygous mutant ES cells, one of the positive clones was transfected once again with the initial targeting vector to obtain one allele still carrying the selection cassette (4.5 kb fragment) and one devoid of (2.7 kb).

Generation of mice lacking the nidogen-binding module γ1IIl4

Two independent heterozygous clones lacking the selection cassette, CIA2 and CII13C6, were injected into C57/B6 blastocystos and transferred into pseudopregnant CD1 foster mothers. Highly chimeric male founder mice were obtained which were crossed with C57/B6 and 129Sv females to obtain heterozygous F1 offspring. Heterozygous mice were mated to obtain time-staged homozygous embryos. All F1 and F2 progeny were genotyped by Southern blotting or by PCR using 5'-AGAGTGAACTCTGATGAC as forward and 5'-TGCA- GAAGTGTTCAACCGCCATTCT as reverse primer.

Preparation of embryoid bodies

Undifferentiated ES cells were trypsinized, diluted in ES medium without LIF (EB medium) as described (Wobus et al., 1991) and plated onto cell culture dishes for 45 minutes to allow residual feeder cells to attach. The supernatant was washed twice with EB medium and ES cells were finally diluted at a concentration of 32×10^3/ml. The cells were then placed in hanging drops of 25 μl (Wobus et al., 1991). After 2 days, cell aggregates were transferred into EB medium filled bacterial dishes. Medium was changed every 48 hours. After 10 days in culture, intact embryoid bodies were collected, washed twice in PBS and further processed.

Protein analysis and rotary shadowing electron microscopy

Embryoid bodies were sequentially extracted with TBS and EDTA as described (Paulsson et al., 1987). The EDTA extract was passed over a Hi-Trap heparin affinity column (Pharmacia). Bound laminin was eluted with a linear NaCl gradient as previously described (Paulsson et al., 1987). Rotary shadowing electron microscopy was performed as described (Paulsson et al., 1987).

For radioimmuno inhibition assays (RIA) embryoid bodies from three independent homozygous mutant and control cells and embryos were extracted in RIPA buffer (Sasaki et al., 1996) and analyzed as described (Sasaki et al., 1998). For immunoblotting, 5 μg total protein was separated under non-reducing conditions on 5-15% SDS-PAGE gels, transferred onto PVDF membranes (Millipore) and incubated with the primary antibodies. Using goat-anti-rabbit antibodies conjugated with horseradish peroxidase (BioRad), specific bands were detected after visualizing enzyme activity with ECL (Amersham).

MATERIALS AND METHODS

Construction of the targeting vector

A 164 bp cDNA fragment coding for γ1IIl4 was used for probing a lambd FIX II genomic library (Stratagene) of the mouse 129SvJ strain. Exons coding for domain III of the laminin γ1 chain were identified by hybridization with radiolabeled oligonucleotides according to the published human genomic organization of the Lama1 gene (Kallunki et al., 1991). The nidogen-binding module γ1IIl4 was contained within a 4.7 kb HindIII/BamHI fragment. Deletion of γ1IIl4 was achieved within a unique 400 bp Apal/NheI fragment using 5'-GATCGGGCGTCGAGCTGCAAAAGTGTAGTCAGGGCA GACTCGCTACG and 5'-ACAGAGAGCTGTCATGCAATAAAA- CACATTGCC, and verified by sequencing. After subcloning the mutated Apal/NheI fragment into the HindIII/BamHI genomic clone, the HSV-TKNeo cassette flanked by two loxP sites was inserted into the NheI site in the following intron. To identify Cre recombination, an additional EcoRV restriction site was introduced after the second loxP site. The mutated clone was then extended by a 3’-following 5.2 kb BamHI genomic fragment.

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Northern RNA blotting and RT-PCR

Total RNA was prepared using Trizol, according to the suppliers protocol (GibcoBRL). Samples (10 μg) of each genotype were electrophoresed in a 1.2% denaturing formaldehyde gel and transferred to HybondN membrane (Amersham) and hybridized against 32P-oligolabeled cDNA probes for laminin γ1, laminin γ1III4, perlecain, nidogen 1 and nidogen 2. For standardization, the same membranes were reprobed with a GAPDH probe.

For RT-PCR, 5 μg of total RNA was reverse transcribed with MuLV reverse transcriptase (Appligene). For second strand synthesis 2 μl of the reaction mixture were used in PCR reactions. After 35 cycles, PCR products were analyzed on a 2% agarose gel. Control reactions were carried out without reverse transcription. Forward and reverse primers were chosen from different exons and were as follows: for demonstrating deletion of the nidogen-binding module γ1III4, 5’-AGATGGAAACTCTTGAGTGAC and 5’-TTGTAGTAGACGCCAGGTCACAAGTA; for LE modules 6-7 of the laminin γ1III domain, located downstream of the deletion site, 5’-GTGTAAGCCATGTGAC-3’ and 5’-CTTCAACAGTCTGTAACAGCGG; for GAPDH, 5’-CTGCCAAGTGATGACATCA-3’ and 5’-TACCTCTTGAGGCCCAGTAGG; yielding in reaction products of 369 and 201 bp for wild-type and mutated alleles for the deletion site, and 317 and 252 bp products for LE6-7 and GAPDH, respectively. The PCR products were verified by sequencing.

Histochemistry and immunohistochemistry

Embryos and tissues were dissected, fixed for 2-4 hours in 4% paraformaldehyde in PBS and either embedded in OCT or in paraffin wax. Paraffin wax-embedded sections (5-10 μm) were processed for Hematoxylin and Eosin staining.

For immunostaining 7-10 μm cryosections were stained with the respective antibodies. Paraffin wax-embedded sections were dewaxed and treated for 10 minutes with 100 μg/ml proteinase XXIV (Sigma) at 37°C. After a blocking step with PBS, 5% normal goat serum (NGS), first antibodies were applied in PBS, 2% NGS for 1 hour at 37°C in a humidified chamber. After washing with PBS, the sections were incubated with secondary Cy3- or Cy2-conjugated goat-anti-rabbit or goat-anti-rat antibodies. After final washing with PBS, sections were mounted and analyzed on an Axiophot fluorescence (Zeiss) and an Mountar PM 2000. Hematoxylin and Eosin staining.

Electron microscopy

Tissue specimens (1 mm²) were fixed in 3% paraformaldehyde and 3% glutaraldehyde in PBS for 2 hours at 4°C as described (Mayer et al., 1997). They were then postfixed for 1 hour in 1% osmium tetroxide and embedded in Epon. Ultrathin sections were collected on formvar coated copper grids, stained for 10 minutes with uranyl acetate, 5 minutes with lead citrate and examined with a Zeiss EM 109 electron microscope. For quantification of lung morphology, eight tissue samples from wild-type and heterozygous mice were obtained and did not show any obvious differences compared with those devoid of the nidogen-binding site with respect to size and spontaneous contraction.

RESULTS

Targeted inactivation of the nidogen-binding site within the Lamac1 gene abrogates nidogen binding

To explore the biological role of the laminin-nidogen interaction, a targeting vector was prepared that lacked the nidogen-binding LE module, γ1III4, within domain III of the laminin γ1 chain. The nidogen-binding module is encoded together with part of the preceding LE module in one exon. Deletion of γ1III4 was achieved using a PCR-based strategy, leaving 7 bp of the original sequence at the 3′-end of the exon, thus allowing fusion of γ1III3 directly to γ1III5 after splicing. The inserted selection cassette was flanked by two loxP sites and excised after transient transfection with Cre recombinase (Fig. 1A). Homologous recombination in embryonic stem cells (ES) was followed by the appearance of a 4.5 kb band for the targeted allele, which was further reduced in size to 2.7 kb after Cre-mediated recombination (Fig. 1B).

To analyze the role of the laminin-nidogen interaction in vitro, we generated ES cells homozygous mutant for the deletion (Fig. 1B). RT-PCR analysis with primers flanking γ1III4 showed the correct size of the amplified products for the wild-type and mutated alleles, indicating that the deletion close to the exon border did not interfere with transcription and correct splicing (Fig. 1C). To prove that the deletion of γ1III4 completely abolishes nidogen-binding activity, laminin was purified from wild-type and γ1III4-deficient embryoid bodies by heparin affinity chromatography. In agreement with earlier studies (Paulsson et al., 1987), nidogen 1 was retained together with laminin in the matrix in wild-type extracts, while it was exclusively detected in the flow through fraction in mutant extracts, demonstrating that mutant laminin had lost nidogen-binding activity (data not shown). This was further confirmed by rotary shadowing electron microscopy showing nidogen 1 closely associated with the short arm of laminin in controls, whereas a related structure was absent in the mutant laminin (Fig. 2). Furthermore, the typical cruciform shape and the correct number of globular domains in the short arms of the mutant laminin (Fig. 2) excluded the possibility that the deletion had interfered with folding and assembly of laminin into a trimer.

Embryoid bodies grown from wild-type and heterozygous cells did not show any overt differences compared with those devoid of the nidogen-binding site with respect to size and spontaneous contraction.

Kidney organogenesis is dependent on the laminin-nidogen interaction

Heterozygous mice derived from two independent ES cell lines were obtained and did not show any obvious phenotype up to one year of age. Immunostaining with antibodies specific for the interfaces of the tandem arrays of LE modules in the wild-type or mutant (Mayer et al., 1998) demonstrated that both the mutant and the wild-type laminin γ1 chain were present in basement membranes (data not shown). Genotyping of more than 800 offspring of heterozygous crossings at weaning failed to reveal any surviving homozygous mutant animals (+/+ or +/− or −/−; 280:524:0). Dead pups were routinely observed shortly after birth of which most were homozygous for the mutation. This suggested that mice deficient for the nidogen-binding module γ1III4 died soon after birth.
only ~60% of the mutant embryos developed to term, whereas ~40% died before 11.5 dpc for so far unknown reasons. Embryos were removed at 18.5 dpc by Caesarian section. The majority of the mutant embryos were about 20% smaller than their littermate controls. Strikingly, visual inspection demonstrated the lack of kidneys in most homozygous embryos (Fig. 3C,E) whereas they were present in all wild-type (Fig. 3A,D) and heterozygous embryos analyzed. Both kidneys were identifiable in only a small percentage of mutant embryos (Fig. 3B), while ~90% showed either bilateral (80%; 46 out of 59 animals) or unilateral (10%; 5 out of 59 animals) renal agenesis, demonstrating that the laminin-nidogen interaction plays a pivotal role during kidney organogenesis. The low penetrance of metanephric development was independent of the ES clone or genetic background being observed on a heterogeneous (C57/129Sv) as well as on a 129Sv background. Despite the lack of kidneys, adrenal glands, testis and ovary were normally present with gross and histological analysis failing to reveal any abnormalities in these organs, while the uterus in female and vas deferens and the seminal vesicles in male mutants were absent (Fig. 3C,E).

**Growth of Wolffian duct is impaired in g1III4-deficient mice**

An essential step during kidney development involves growth of the Wolffian duct and its subsequent fusion with the cloaca, a prerequisite for the outgrowth of the ureteric bud (Saxen et al., 1987). The ureteric bud subsequently induces the metanephric mesenchyme to condense and to undergo a mesenchymal-epithelial conversion. As kidney organ culture studies have demonstrated that less new epithelium formed around the tips of the ureteric bud in the presence of antibodies inhibiting the laminin-nidogen interaction (Ekblom et al., 1994), defective epithelialization could have led to subsequent degeneration of the kidney anlage, resulting in renal agenesis. We therefore analyzed kidney development at 13.5 dpc with standard histological techniques. In all wild-type and heterozygous embryos, multiple branches of the ureteric bud surrounded by condensed metanephric mesenchyme were observed (Fig. 4A) that were indistinguishable in number and...
morphology from those mutant embryos in which either one or two metanephric kidneys had been induced (Fig. 4C). However, in the majority of γ1III4-deficient embryos, only remnants of uncondensed metanephric blastema were observed (Fig. 4B).

This raised the possibilities that either the ureteric bud failed to branch from the Wolffian duct at earlier stages or that its growth towards the metanephric mesenchyme was defective. At 11 dpc, the first dichotomous branching of the ureteric bud was visible in all wild-type, heterozygous and a few mutant embryos (Fig. 4D,F). In most of the mutant embryos, however, we were not able to identify any signs of the ureteric bud (Fig. 4E) and thus defective or delayed growth followed by cell death could be excluded. Surprisingly, a morphologically identifiable Wolffian duct was also absent in the caudal region of the embryos. During embryonic development, two pairs of renal organs, the pronephros and the mesonephros, are formed in a spatial and temporal sequence before the metanephros becomes the adult functional kidney. As all three organs are formed by mesenchymal-epithelial transition, kidney agenesis could have been explained by a general failure of this conversion. Yet, the mesonephros was properly formed and in size and number of tubuli similar to wild-type embryos at 11.5 dpc (Fig. 5A,B). At the same stage the caudal progression of the Wolffian duct towards the cloaca could be followed in serial sagittal and transverse sections in wild-type embryos (Fig. 5C). In eighteen γ1III4-deficient embryos devoid of metanephric kidney induction, however, the Wolffian duct was blind-ending within the upper level of the urogenital ridge (Fig. 5D). In support of these data, at 13.5 dpc we also observed a blind-ending Müllerian duct, which is known to develop in parallel to and to be dependent on an inductive influence of the Wolffian duct (Jacob et al., 1999). Despite the absence of a ureteric bud, we could identify distinct uninduced metanephric mesenchyme in all cases.

Growth of the Wolffian duct is still poorly understood. We speculated that a defective basement membrane along the duct might cause retardation of its growth. Yet, immunostaining for laminin revealed normal, linear basement membrane structures (Fig. 5F). A closer inspection by confocal microscopy, however, demonstrated that discontinuities close to the growing tip of the Wolffian duct existed in deeper layers of the sections, and that the normally tightly connected epithelial cells were partially separated (Fig. 5G). These results
were further supported by electron microscopy showing that basement membranes were interrupted by gaps along the Wolffian duct (see Fig. 9D,E). Epithelial cell shape and polarity, however, appeared normal. To determine whether growth retardation is correlated with a decrease in cell proliferation, we performed BrdU-labeling in utero and found that the number of BrdU-positive cells was not significantly changed compared with controls (Fig. 5G). Similarly, staining for Pax2, a transcriptional regulator of the paired-box family known to result in Wolffian duct dysgenesis (Torres et al., 1995) was normal (Fig. 5H,I). From these data, we conclude that growth inhibition of the Wolffian duct in γ1III4-deficient embryos is dependent on a locally restricted defective basement membrane, but independent of a proliferation defect or differentiation processes regulated by Pax2.

Kidney dysgenesis caused by the absence of the laminin-nidogen 1 interaction

Kidney agenesis occurred in 90% of the animals. In the 10% of mutant embryos in which kidney development did occur, we observed normal kidney architecture with multiple branches of the ureteric bud between 11.5 and 13.5 dpc (Fig. 4C,F). At 18.5 dpc, however, mutant kidneys seemed to be non-functional, as the urinary bladders were morphologically empty. Dysgenesis of variable severity was apparent in the mutant kidneys. While the outer developing cortical region was comparable with the control tissues, the quantity of tubular structures in the mature region was greatly diminished (Fig. 6A,B). Furthermore, histological sections revealed abnormalities in the glomeruli of the inner cortical region. The total number of mature glomeruli counted in kidney sections obtained from three individual animals was 30% less than in wild-type kidneys. Of these, 60% were normally developed (Fig. 6C), whereas the rest had either a reduced (25%) or missing (15%) capillary tuft, and consequently the Bowman’s capsule was partially or completely filled with red blood cells (Fig. 6D,E). The distal and proximal tubuli, as well as all Comma- and S-shaped bodies were surrounded by an ultrastructurally distinct basement membrane. The same was found for the majority of glomerular basement membranes (GBM), although when the capillary tuft was missing the GBM was not linear but interrupted (data not shown).

In addition, some of the mutant kidneys analyzed showed cystic enlargements, which were only observed in male homozygotes. To define the defect more closely, we followed the ureter in transverse sections from three male mutant embryos at 18.5 dpc caudally towards the bladder. A hydromephric kidney with a thinned parenchyma was apparent (Fig. 6F) and the ureter connected aberrantly either to the vas deferens (Fig. 6G) or, in one case, ended directly in the epididymis, both derivatives of the Wolffian duct. During dissection of the urogenital tracts, we observed a vesicular structure in close vicinity to the testis in mutant embryos (Fig. 3B), which was identified as epididymis. These data therefore strongly suggest that in these males, the kidney directly drained into the genital tracts, thus leading to loss of the normal architecture of the epididymis.

Nidogen 1 is not retained in basement membrane structures

To determine whether the deletion of γ1III4 within the laminin γ1 chain affects the localization of nidogen, we performed immunostaining of various tissues. Nidogen 1 was present in all basement membranes of wild-type and heterozygous animals, but, with a few exceptions, barley detectable in basement membrane structures of the mutant animals. Only basement membranes around large blood vessels and at the dermal-epidermal junction of the skin showed staining intensities comparable with the control tissues (data not shown). This raised the question as to whether the recently newly identified, highly homologous isoform, nidogen 2 (Kohfeldt et al., 1998), is upregulated. Double-immunostaining for nidogen 1 and nidogen 2 in kidney sections, however, indicated that nidogen 2 was neither differently deposited nor
was its staining intensity increased relative to the controls (Fig. 7A-D). Staining intensities similar to that of wild-type controls were also observed for the laminin γ1 chain (Fig. 7E,F), collagen IV and perlecan (data not shown).

Former studies have demonstrated that nidogen 1 is highly susceptible to proteases and that complex formation with laminin protected the C-terminal globular domain G3 from degradation (Mayer et al., 1993a). Surprisingly, immunoblotting of various tissue extracts showed normal levels of non-degraded nidogen 1 (Fig. 7G), excluding the possibility that nidogen 1 is lost due to excessive proteolysis. Comparable RNA and protein levels for nidogen 1, nidogen 2, laminin and perlecan were also found by radioimmuno inhibition assays and Northern blotting (data not shown). We therefore conclude that through the deletion of the nidogen-binding site, no compensatory mechanisms are induced, but that nidogen 1 is lost during embedding procedures, thus strongly arguing for a weakened integration into basement membranes caused by the deletion of its binding site.

**Lung defects might cause perinatal lethality in γ1III4-deficient mice**

Homozygous mice die very soon after birth and, despite the fact that the kidney phenotype should lead to a perinatal lethality within the first 2 days, we have been unable to identify any living offspring within the first postnatal day. This indicated that another defect causes death of the homozygous animals. Indeed, some of the mutant dead pups had a cyanotic appearance indicative of respiratory problems.

Mutant lungs dissected from 18.5 dpc embryos, were normally developed with respect to number of lobes. Cross-sectioning through the lung revealed that the major bronchial trees had formed in the homozygotes, but they were more compact and smaller compared with wild type (Fig. 8A,B). At higher magnification, it became apparent that the prealveolar sacculi were immature and only poorly inflated, and mesenchymal thickening between the terminal airspaces was observed (Fig. 8C,D). Interaction between mesenchyme and epithelium is a crucial factor throughout lung development. Yet, between 11.5 and 13.5 dpc, no differences in number of buds or sizes of the lungs were observed compared with control embryos (Fig. 8E,F). The presence of surfactant protein C (SP-C) positive cells in 18.5 dpc mutant lungs further indicated that differentiation of precursor epithelial airway cells had proceeded (Fig. 8G,H).

Formation of the thin air-blood barrier is a crucial event for gas exchange and hence crucial for survival. Ultrastructural analysis of four individual γ1III4-deficient embryos at 18.5 dpc demonstrated that only ~10% of the prealveolar sacculi had formed, when compared with controls. In addition, when the air-blood barrier of the homozygotes had formed, the basement membranes showed severe abnormalities, while in the wild type, endothelial and epithelial cells were separated by a fused basement membrane (Fig. 9A-C). The alveolar basement membranes were apparently normal in 30% of the mutants (Fig. 9B), whereas only amorphously deposited material was detected in a further 30% (Fig. 9C). Remarkably, in the remaining alveoli, basement membranes were completely missing and epithelial cell protrusions were seen entering into the empty space (Fig. 9B). Together, these data suggest that the laminin-nidogen interaction is crucially important for the formation of the air-blood interface. Insufficient gas exchange, therefore, may have resulted in perinatal death of the mutant animals.

**DISCUSSION**

The laminin-nidogen interaction is not a prerequisite for basement membrane formation

During embryonic development, laminin 1 and nidogen 1 are first expressed at the four- and 32-cell stages, respectively...
including a high-affinity interaction to the perlecan core protein (Hopf et al., 1999). The finding that normal immunoreactivity was only apparent in a few basement membrane structures, strongly argues that nidogen 1 is preferably integrated into basement membranes via laminins containing the γ1 chain.

It has previously been shown that inactivation of the Nid1 gene in mice and C. elegans did not interfere with basement membrane formation and the animals are viable and were fertile (Murshed et al., 2000; Kang and Kramer, 2000). Yet, compensation by nidogen 2, which has a highly homologous domain structure and binding repertoire (Kohfeldt et al., 1998), might be the reason for the lack of phenotype in Nid1 mutant mice (Murshed et al., 2000), and, vice versa, the same may be true for mice with a mutation in Nid2 (Mitchell et al., 2001). Although human nidogen 2 binds with only low affinity to the nidogen 1-binding site within the laminin γ1 chain (Kohfeldt et al., 1998), the murine homolog has now been shown to be similar in binding affinity as nidogen 1 (T. Sasaki and R. Timpl, personal communication). It will therefore be of interest to see whether the double mutation of both genes reflects the phenotype described here for the deletion of the nidogen-binding site. Together, the genetic analyses supports the conclusion that neither nidogen 1 (Murshed et al., 2000; Kang and Kramer, 2000) nor, as shown here, its binding site on the laminin γ1 chain are crucial for basement membrane assembly and function in most tissues. Only a few tissues, including the cortex and neural tube, showed subtle discontinuities in basement membranes and developmental abnormalities (Halfter et al., 2002), arguing for a specific physiological function of the laminin-nidogen interaction. However, we cannot exclude the possibility that other, presently unknown functions of the nidogen-binding module of the laminin γ1 chain might contribute to the described phenotype.

The laminin-nidogen interaction is crucially important for Wolffian duct growth

Genetic data obtained from human diseases and transgenic mice have implicated transcription and growth factors, cell-surface receptors and extracellular matrix components in metanephric development (Lechner and Dressler, 1997; Müller and Brändli, 1999). Renal agenesis to a variable degree was manifested by mutations in several of these genes. Perturbed communication between cellular receptors and their respective ligands on either the metanephric blastema or the ureteric bud have been suggested in null-mutations for the glial cell line-derived neurotrophic factor, GDNF, and the tyrosine kinase receptor Ret (Lechner and Dressler, 1997), as well as in integrin α8-deficient mice (Müller et al., 1997), leading to defective metanephric kidney induction. However, renal agenesis in the absence of the laminin-nidogen 1 interaction is due to defects in Wolffian duct elongation and therefore manifests earlier during development.

The Wolffian duct develops from the intermediate mesoderm and gives rise to parts of the male genital system. In agreement with a primary defect in Wolffian duct growth, vas deferens and the seminal vesicles in male mutant embryos were missing, while the testes were normally present. The female genital tracts originate from the Müllerian duct, which forms later in development in parallel to the Wolffian duct, and its growth was shown to be dependent on an inductive influence of the
former (Jacob et al., 1999). In support of these data, we found both ducts to be blind-ending in the urogenital ridge in 13.5 dpc embryos and the female mutants to lack the uterus and occasionally the oviduct.

The Wolffian duct develops in the absence of the laminin-nidogen interaction, excluding its contribution to the first mesenchymal-to-epithelial transition that occurs in kidney development (Saxen, 1987), but its growth caudally towards the cloaca is inhibited. A similar phenotype was described for the targeted ablation of Pax2, a member of the paired-box family of regulatory transcription factors. As γIII4-deficient mice, Pax2 mutants fail to form parts of the genital tract and the ureteric bud (Torres et al., 1995), suggesting that growth of the Wolffian duct is affected in a similar manner. One possibility therefore is that deletion of the nidogen-binding site interferes with Pax2 expression in the epithelial tube. Yet, normal immunostaining of Pax2 in the mutant Wolffian duct indicates that it is not affected through the deletion.

Although we do not know the detailed mechanism responsible for the phenotype of our γIII4 mutants, these mice now provide a model system to further our understanding of Wolffian duct growth and elongation, which has long been a subject of interest because of its central position in urogenital development. Once formed, the duct progressively elongates; so far two mechanisms have been proposed for this process: one suggests autonomous growth through cell proliferation at the tip of the duct, whereas a second model proposes that mesenchymal cells are continuously added by an epithelial transition (Saxen, 1987). The data presented here provide evidence that Wolffian duct elongation through autonomous cell proliferation at its growing tip can be excluded. First of all, the number of BrdU-positive cells were similar in mice with the nidogen-binding site deletion as in controls, and did not accumulate at the tip of the growing duct, but uniformly spread over the length of the duct as it has been described in amphibians (Overton, 1959), indicating that the underlying mechanisms are not due to reduced cell proliferation. Strikingly, however, we identified subtle local ruptures in the basement membrane close to the tip of the duct that may well interfere with duct elongation. Nidogen-binding to laminin could modulate the conformation of laminin or affect the spatial relationships between basement membrane components, which facilitate their interaction with cellular receptors. We therefore propose that the locally restricted discontinuities seen in basement membranes around the Wolffian duct are due to a perturbed cell-matrix interaction, which in turn interferes with signaling cascades necessary to induce the genetic program to recruit mesenchymal cells.

The function of the laminin-nidogen interaction for epithelial branching morphogenesis in kidney and lung

The laminin-nidogen interaction has been suggested to be crucial for epithelial branching morphogenesis in general (Ekblom et al., 1994; Kadoya et al., 1997). Our data indicate that early branching morphogenesis of the lung is not disturbed in the absence of the laminin-nidogen interaction. However, shortly before birth, the lungs were only poorly inflated, with a mesenchymal thickening of the distal airspaces. Consequently, a respiratory problem caused with high probability the death of all the mutant pups. Interestingly, a similar phenotype has been reported in the absence of TGFβ3 (Kaartinen et al., 1995). The phenotype in TGFβ3-deficient mice has been correlated with reduced number of airway precursor cells (Shi et al., 1999). Although we identified these cells in γIII4-deficient lungs by staining for SP-C, a detailed statistical analysis at different developmental stages is required to determine whether they are formed in similar numbers as in wild type. Ultrastructural analysis demonstrated that the majority of the distal airspaces in the mutants were uninflated.
GDNF (Schuchardt et al., 1996; Moore et al., 1996; Müller et al., 2000) have independent lethal phenotypes at different developmental stages. For example, deficiencies for the integrin α4 and αv subunits or perlecan, have independent lethal phenotypes at different developmental stages, depending on the individual animal (Yang et al., 1995; Bader et al., 1998; Costell et al., 1999). Furthermore, variable penetrance of impaired kidney development has been observed upon mutations of the integrin α8 subunit, Ret or GDNF (Schuchardt et al., 1996; Moore et al., 1996; Müller et al., 1997). Similarly, a minority of the γ1III4-deficient animals displayed metanephric development, which may be due to either intact or regenerated basement membranes along the Wolffian duct at crucial stages of development. However, the number of tubuli and glomeruli in the mature part of the cortex in 18.5 dpc mutant kidneys was reduced even in those mutants in which induction of the metanephric blastema and branching of the ureteric bud proceeded without any obvious defects. Inhibition of the laminin-nidogen interaction by antibodies led to reduced tubulogenesis in an in vitro organ culture model, although in contrast to the present observations in vivo, basement membranes were largely distorted in the in vitro analysis (Ekblom et al., 1994). Thus, although the basement membrane architecture in the γ1III4-deficient kidneys appeared normal, it is tempting to speculate that locally restricted and transient basement membrane ruptures also occur during kidney morphogenesis, resulting in the differentiation defect.

Unexpectedly, the bladder of the mutant animals was empty even when kidneys had developed. The histological analysis in male mice demonstrated that the ureter failed to open into the bladder but instead stayed connected aberrantly either to the vas deferens or the epididymis, both derivatives of the Wolffian duct. The cysts seen in these kidneys are therefore likely to be secondary to a defect in urine outflow. These results also indicate that the reduction of the glomerular tufts is attributed to increased pressure through a tailback of the urine, rather than by a primary defect in mesangial cells as described in Pdgfb mutant mice (Leveen et al., 1994), and their GBM may rupture because of mechanical stress. Interestingly, this phenotype resembles congenital anomalies of the kidney and urinary tract (CAKUT) found in humans and mice (Pope et al., 1999). An abnormal ureter connection in conjunction with a hydronephric kidney has been reported for heterozygous Bmp4 mutant mice (Miyazaki et al., 2000). In these mice it is suggested that a deficit in BMP4 levels inhibits branching of the ureterovesicle junction into the cloaca as a result of impaired elongation of the ureter (Miyazaki et al., 2000). This model is reminiscent to the growth defect of the Wolffian duct in the urogenital ridge in the γ1III4 mutant animals. We therefore propose that a similar mechanism to that described above leads to the aberrant ureter fusion. However, we cannot exclude at the moment that the absence of the laminin-nidogen interaction causes subtle changes in the supramolecular organization of basement membranes, which then in turn could interfere with the sequestration of BMP4 or other growth and morphogenetic factors, or that specific unknown physiological functions exist for the nidogen-binding module, γ1III4 of the laminin γ1 chain.

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