Podocyte-specific knockout of myosin 1e disrupts glomerular filtration

Sharon E. Chase,1 Christina V. Encina,1,2 Lindsay R. Stolzenburg,1 Arthur H. Tatum,3 Lawrence B. Holzman,4 and Mira Krendel1

1Department of Cell and Developmental Biology, SUNY Upstate Medical University, Syracuse, New York; 2Department of Clinical Laboratory Science, SUNY Upstate Medical University, Syracuse, New York; 3Department of Pathology, SUNY Upstate Medical University, Syracuse, New York; and 4Renal-Electrolyte and Hypertension Division, University of Pennsylvania, Philadelphia, Pennsylvania

Submitted 2 May 2012; accepted in final form 16 July 2012

Chase SE, Encina CV, Stolzenburg LR, Tatum AH, Holzman LB, Krendel M. Podocyte-specific knockout of myosin 1e disrupts glomerular filtration. Am J Physiol Renal Physiol 303: F1099–F1106, 2012. First published July 18, 2012; doi:10.1152/ajprenal.00251.2012.—Myosin 1e (myo1e) is an actin-dependent molecular motor that plays an important role in kidney functions. Complete knockout of myo1e in mice and Myo1E mutations in humans are associated with nephrotic syndrome and focal segmental glomerulosclerosis. In this paper, we tested the hypothesis that myo1e is necessary for normal functions of glomerular visceral epithelial cells (podocytes) using podocyte-targeted knockout of myo1e. Myo1e was selectively knocked out in podocytes using Cre-mediated recombination controlled by the podocin promoter. Myo1e loss from podocytes resulted in proteinuria, podocyte foot process effacement, and glomerular basement membrane disorganization. Our findings indicate that myo1e expression in podocytes is necessary for normal glomerular filtration and that podocyte defects are likely to represent the primary pathway leading to glomerular disease associated with Myo1E mutations.

myosin; podocyte; proteinuria

ONE OF THE MAIN FUNCTIONS performed by the kidney is regulation of protein excretion in the urine. In the healthy kidney, glomerular filtration barrier retains the majority of high molecular weight proteins. Glomerular disorders, both inherited and acquired, are associated with proteinuria, which, if not reversed, leads to end-stage renal disease. Genetic studies and experiments using knockout mouse models have implicated many cytoskeletal proteins, particularly those associated with the actin cytoskeleton, in regulation of glomerular permeability (5, 6, 23). One of the cytoskeletal proteins that contribute to the normal functioning of the glomerular filtration barrier is an actin-dependent molecular motor myosin 1e (myo1e). Myo1e-null mice developed by our lab exhibit extensive proteinuria, indicating that myo1e plays a central role in regulating glomerular filtration (12). Two recent genetic studies also identified an association between mutations in the human Myo1E gene and the autosomal recessive nephrotic syndrome (13, 15). These findings underscore the importance of myo1e to normal glomerular functions.

A key question that remains unanswered by the genetic studies is which specific cell type(s) in the glomerulus requires myo1e activity and represents the primary site of injury in the myo1e-null glomeruli. Glomerular filtration apparatus is composed of three main components: endothelial cells of glomerular capillaries, glomerular visceral epithelial cells (podocytes) covering the surface of the capillaries, and glomerular basement membrane (GBM), which is secreted by both endothelial and epithelial cells. In addition to the endothelial and epithelial cells of the glomerulus, another cellular component that contributes to normal glomerular development and functions is glomerular mesangium, composed of contractile cells that shape capillary loops and maintain the overall architecture of the capillary tuft (16). Injury or abnormal development of the cellular components of the glomerulus, particularly podocytes and endothelial cells, results in proteinuria (7, 21).

Since myo1e is highly expressed in podocytes and the phenotype of myo1e-null mice is consistent with a podocyte defect, we set out to test the hypothesis that myo1e expression is necessary for normal podocyte development and functions. As a direct test of this hypothesis, we created a mouse model that allows selective inactivation of myo1e in podocytes through Cre-mediated recombination. Characterization of glomerular functions and organization in this mouse model show that podocyte-targeted loss of myo1e causes proteinuria and ultrastructural defects of podocytes and GBM.

MATERIALS AND METHODS

Animal studies. All animal experiments were conducted in accordance with the protocols approved by the SUNY Upstate Committee for Humane Use of Animals. Complete myo1e-knockout mice and conditional myo1e-knockout mice (previously described in Ref. 12) were backcrossed to C57bl/6 mice for 12 generations. Conditional knockout (“floxed”) mice were then crossed with NPHS2-Cre (Podo-Cre) mice (14) on C57bl/6 background. The following PCR primers were used for genotyping: Myo1e A (TCATGTGTAGCTGTCATCTGACTCCACC), Cre For (GTTCAGGGATCGCCAGGCG), Cre Rev (GGACATGTGTAGCCTGTCATCTGACTCCACC), Myo1e B (TTCCGCTTACGGTGGAAATG), Myo1e C (ACTCATTCTGCATCTGACTCCACC), Cre For (GCATAACAGTAGAAGAACACATCGTCTG), Cre Rev (GGACATGTTCAGGGATCGCCAGGCC). Albumin concentration in mouse urine samples was determined using SDS-PAGE with bovine serum albumin standards followed by gel densitometry using ImageJ software. Creatinine concentration was measured using Creatinine Companion Kit (Exocell).

Antibodies. The following primary antibodies were used for immunostaining: rabbit anti-myoe (20), mouse anti-synaptopodin G1D4 (Meridian Life Science), rat anti-PECAM-1 (BD Pharmingen), mouse anti-desmin (BD Pharmingen), rabbit anti-α-actinin 4 (MABT144, Millipore), rabbit anti-myoe2a (Biomedical Technologies), rabbit anti-podocin (Alpha Diagnostics), rabbit anti-integrin α3 (AB1920, Millipore), rabbit anti-phospho-nephrin (22).

Immunohistochemistry, histology, and electron microscopy. For immunostaining, fresh kidneys were frozen in OCT and sectioned. Cryosections were fixed with 3% paraformaldehyde in PBS for 15 min, permeabilized using 0.25% Triton X-100 in PBS for 3 min, and processed for immunofluorescence or immunohistochemical staining. Immunohistochemical staining was performed using Vector Labs Vectastain Elite ABC anti-rabbit kit following the manufacturer’s
instructions. Histological staining was performed on sections of formalin-fixed, paraflin-embedded kidneys. Images of immunofluorescently labeled kidney sections were collected using a Perkin-Elmer UltraView VoX Spinning Disc Confocal system mounted on a Nikon Eclipse Ti microscope. Images of histology slides were collected using a color Spot camera on a Nikon Eclipse TE-2000 microscope. Kidney fixation and processing for electron microscopy were performed as previously described (12). Morphometry on kidney sections was performed using ImageJ software. Five to fifteen micrographs per animal (obtained from different glomeruli) were included in the analysis, and four animals were analyzed for each genotype. GBM in each electron micrograph was manually outlined in ImageJ to determine its area. The total area of all regions of the basement membrane present in each micrograph was divided by the total length of the basement membrane (measured by drawing a line through the center of each basement membrane segment) to determine its average thickness. Foot process number was manually counted and divided by the length of the basement membrane.

**RESULTS**

Generation of mice with the podocyte-targeted knockout of Myo1e. We previously generated mice containing a conditional (floxed) allele of Myo1e so that Myo1e gene could be removed by Cre-mediated homologous recombination (12). In the previously published study (12), mice carrying floxed Myo1e allele were crossed with mice that expressed Cre in the germline, allowing us to obtain the complete Myo1e knockout (Myo1e-null or −/− mice). In the current study, we used mice that express Cre recombinase under the control of podocin promoter (14) to achieve podocyte-specific removal of Myo1e (Fig. 1). Loss of myo1e from podocytes was verified using immunostaining of mouse kidney sections with the previously characterized myo1e-specific antibodies (Fig. 2) (4). Mice carrying at least one wild-type copy of Myo1e (Myo1e+/−) exhibited normal myo1e expression, with the majority of myo1e colocalizing with the podocyte marker synaptopodin (Fig. 2A). Most mice with the podocyte-targeted Myo1e knockout (Myo1eF/F-PodoCre) showed no expression of myo1e in podocytes (Fig. 2C). However, some of the mice carrying two conditional alleles of Myo1e (Myo1eF/F-PodoCre) exhibited incomplete removal of Myo1e from podocytes, as indicated by a significant amount of glomerular immunostaining for myo1e in such animals (data not shown). To increase the efficiency of myo1e removal from podocytes, we generated mice that carried one copy of the conditional myo1e allele and one copy of the myo1e-null allele (Myo1e−/−PodoCre) so that only one copy of the gene would need to be removed by the Cre recombinase. Mice of this genotype exhibited a highly reproducible loss of Myo1e immunoreactivity from podocytes (Fig. 2). Some residual myo1e staining was observed in the glomeruli of animals with the podocyte-specific Myo1e knockout (Fig. 2). This residual myo1e labeling was observed primarily in cells positive for the smooth muscle marker desmin but also showed some colocalization with cells expressing the endothelial marker PECAM-1 (Fig. 2, E and F). As previously reported (12), myo1e-null mice showed no myo1e staining in the glomeruli (Fig. 2B). Thus, as anticipated, PodoCre-mediated recombination disrupted myo1e expression in podocytes. Unexpectedly, it also revealed a low level of myo1e expression in other glomerular cell populations, particularly in mesangial cells, which are positive for desmin.

Characterization of renal filtration in mice with the podocyte-specific Myo1e knockout. To determine how the loss of myo1e from podocytes affected kidney functions, we measured urinary albumin excretion in control (Myo1e−/−) and podocyte-specific knockout (Myo1e−/−PodoCre) mice. Myo1e−/−PodoCre mice developed moderate proteinuria by 8 wk of age, while control mice had normal glomerular filtration with no evidence of albuminuria up to 6 mo of age (Fig. 3, A and B). We also evaluated proteinuria in mice of the Myo1eF/F/PodoCre genotype (Fig. 3A). Many of these animals developed proteinuria by 8 wk of age. However, those mice that retained myo1e expression in podocytes, as verified by immunostaining (data not shown), did not exhibit proteinuria; some of these animals maintained normal urinary protein excretion up to the end of the screening period (6–8 mo of age).

Comparison of proteinuria levels in mice with the complete and podocyte-targeted myo1e knockout. While urinary albumin excretion in podocyte-targeted myo1e-knockout mice was elevated compared with the controls, the extent of proteinuria in this mouse model was fairly modest, with only an order of magnitude of glomerular immunostaining for myo1e in such animals (data not shown). To increase the efficiency of myo1e removal from podocytes, we generated mice that carried one copy of the conditional myo1e allele and one copy of the myo1e-null allele (Myo1e−/−PodoCre) so that only one copy of the gene would need to be removed by the Cre recombinase. Mice of this genotype exhibited a highly reproducible loss of Myo1e immunoreactivity from podocytes (Fig. 2). Some residual myo1e staining was observed in the glomeruli of animals with the podocyte-specific Myo1e knockout (Fig. 2). This residual myo1e labeling was observed primarily in cells positive for the smooth muscle marker desmin but also showed some colocalization with cells expressing the endothelial marker PECAM-1 (Fig. 2, E and F). As previously reported (12), myo1e-null mice showed no myo1e staining in the glomeruli (Fig. 2B). Thus, as anticipated, PodoCre-mediated recombination disrupted myo1e expression in podocytes. Unexpectedly, it also revealed a low level of myo1e expression in other glomerular cell populations, particularly in mesangial cells, which are positive for desmin.

Characterization of renal filtration in mice with the podocyte-specific Myo1e knockout. To determine how the loss of myo1e from podocytes affected kidney functions, we measured urinary albumin excretion in control (Myo1e−/−) and podocyte-specific knockout (Myo1e−/−PodoCre) mice. Myo1e−/−PodoCre mice developed moderate proteinuria by 8 wk of age, while control mice had normal glomerular filtration with no evidence of albuminuria up to 6 mo of age (Fig. 3, A and B). We also evaluated proteinuria in mice of the Myo1eF/F/PodoCre genotype (Fig. 3A). Many of these animals developed proteinuria by 8 wk of age. However, those mice that retained myo1e expression in podocytes, as verified by immunostaining (data not shown), did not exhibit proteinuria; some of these animals maintained normal urinary protein excretion up to the end of the screening period (6–8 mo of age).

Comparison of proteinuria levels in mice with the complete and podocyte-targeted myo1e knockout. While urinary albumin excretion in podocyte-targeted myo1e-knockout mice was elevated compared with the controls, the extent of proteinuria in this mouse model was fairly modest, with only an order of
magnitude increase over the baseline level. Previously, we observed that albumin excretion in mice with the complete Myo1e knockout (Myo1e<sup>−/−</sup>) on the mixed 129/C57bl/6 genetic background was several orders of magnitude higher than in controls (Myo1e<sup>++/+</sup>) (12). Since all mice in the present study were backcrossed to the C57bl/6 background, which sometimes ameliorates kidney disease (1, 4), we measured proteinuria in mice with the complete myo1e knockout (Myo1e<sup>−/−</sup>) on the C57bl/6 background for comparison with the podocyte-specific knockout mice. Even when the influence of the genetic background was equalized by backcrossing to C57bl/6, myo1e-null mice (Myo1e<sup>−/−</sup>) consistently exhibited much higher levels of proteinuria than mice with the podocyte-targeted myo1e knockout (Fig. 3).
Analysis of glomerular ultrastructure in mice with the podocyte-targeted myo1e knockout. To determine the effects of myo1e removal from podocytes on glomerular ultrastructure, kidney sections from 2- and 4-mo-old mice were analyzed by electron microscopy. These studies were conducted using mice of the Myo1e<sup>−/−</sup>H<sup>11002</sup> and Myo1e<sup>F/F</sup>podoCre genotypes as well as appropriate controls (Myo1e<sup>−/−</sup>H<sup>11001</sup> and Myo1e<sup>F/F</sup>). To limit the ultrastructural analysis to the animals with the successful podocyte-targeted knockout of myo1e, only mice exhibiting detectable levels of proteinuria were used for the analysis of the Myo1e<sup>F/F</sup>podoCre genotype. Pronounced podocyte foot process effacement and basement membrane disorganization and thickening were observed in the 4-mo-old mice with either podocyte-targeted or global myo1e knockout. Since the extent of the changes in the basement membrane and foot process organization was highly variable among individual glomeruli and glomerular segments in each animal, we performed morphometric analysis of the electron micrographs to measure the average basement membrane thickness (area divided by length) and the average number of foot processes per unit length of the basement membrane. Average membrane thickness was increased and foot process number was decreased compared with the control mice (Fig. 4). Some regions of the basement membrane exhibited characteristic electronlucent or “moth-eaten” areas, similar to those observed in the myo1e-null mice and human patients with Myo1E mutations (12, 13), and the overall outline of the basement membrane was jagged and uneven, in contrast to the smooth basement membrane of control animals. In addition, microvillous transformation of podocytes, which represents one of the hallmarks of podocyte damage, was observed in the myo1e-knockout mice.

At 2 mo of age, only a few areas of foot process effacement and GBM thickening were observed in mice with the podocyte-targeted myo1e knockout. This was in contrast to the myo1e-null mice, which exhibited widespread foot process effacement and GBM alterations at 2 mo (data not shown).

Histological characterization of myo1e-knockout kidneys. With routine histology staining, including hematoxylin and eosin, periodic acid Schiff, and trichrome, kidneys of mice with the podocyte-targeted myo1e knockout appeared mostly normal up to 6 mo. At the age of 7–8 mo, some sclerotic glomeruli were observed, along with a few proteinaceous casts indicative of persistent proteinuria (Fig. 5). Mice with the complete myo1e knockout exhibited more pronounced and widespread abnormalities at an earlier age, with multiple sclerotic glomeruli and tubular casts present by 4 mo (Fig. 5). Control mice carrying at least one functional copy of the myo1e allele did not show any signs of glomerular pathology at the ages examined (up to 10 mo).

Analysis of the components of podocyte cytoskeleton and cell adhesion complexes in myo1e-knockout kidneys. Since the abnormalities observed in the myo1e-knockout mice could be caused by defects in the podocyte actin cytoskeletal organization, slit diaphragm structure, or adhesion to the basement membrane, we examined expression and localization of several key markers of the podocyte cytoskeleton and cell adhesion complexes. Expression and localization patterns of the slit diaphragm marker podocin, podocyte-specific cell adhesion receptor α3 integrin, and several key components of the podocyte actin cytoskeleton, including synaptopodin, nonmuscle myosin 2a (encoded in humans by the MYH9 gene), and α-actinin-4, were similar in myo1e-knockout mice and control.
animals (Figs. 1 and 6). We also examined the extent of nephrin phosphorylation as a marker for signaling pathways affecting podocyte cytoskeletal organization (22). No changes in nephrin phosphorylation were observed in the myo1e-knockout mice (data not shown).

DISCUSSION

The key finding of this study is that selective removal of myo1e from podocytes is sufficient to induce proteinuria. Mice with the podocyte-targeted knockout of myo1e also exhibit defects in the glomerular ultrastructure, including foot process effacement and thickening and delamination of GBM. These ultrastructural changes are similar to those observed in the myo1e-null mice and in focal segmental glomerulosclerosis patients with Myo1E mutations (12, 13). Thus, the loss of myo1e from podocytes is sufficient to disrupt glomerular filtration and to induce severe structural defects in the glomerulus.

While mice with the podocyte-targeted knockout of myo1e exhibit defects in protein filtration, the timing of onset and the extent of proteinuria are less dramatic than in myo1e-null mice. Proteinuria in mice with the podocyte-targeted knockout of myo1e develops by 2 mo of age while myo1e-null mice develop proteinuria by 1 mo of age. This delay may be due to the timing of myo1e removal using Cre recombinase expressed under the control of podocin promoter. Since podocin promoter activity is induced fairly late in the glomerular development, during capillary loop stage (14), in the podocyte-targeted knockout mice expression of myo1e may persist up to a fairly late stage in podocyte differentiation. The timing of myo1e loss in the podocyte-targeted mouse model may also explain the difference in the observed proteinuria level. If myo1e plays an important role both in the development of early podocyte precursors and in the functions of mature podocytes, myo1e-null mice would exhibit more severe glomerular defects than mice that lose myo1e expression in podocytes during capillary loop stage.

Moreover, myo1e may be necessary not only for podocyte functions but also for the normal physiological functions of other cells in the glomerulus. Since we observed myo1e expression in mesangial cells, it is possible that myo1e contributes to mesangial cell migration to glomeruli during their development or to normal mesangial cell functions in the mature glomerulus. To address this possibility, we examined the number and morphology of mesangial cells in the kidneys of myo1e-knockout mice. Glomeruli of both myo1e-null and wild-type mice at various ages (newborn, 7-day-old, 21-day-old, 3-mo-old) contained desmin-positive mesangial cells (data not shown) and no mesangial cell abnormalities were apparent from electron micrographs of myo1e-null glomeruli. Therefore, while we cannot exclude the possibility that the loss of myo1e may negatively affect mesangial cell functions, no obvious evidence of mesangial cell loss or defects in the myo1e-null mice was found in this study.

In our study, the severity of proteinuria correlated with the prevalence of histological abnormalities in the kidneys. Mice with the complete knockout of myo1e exhibited widespread glomerulosclerosis by 4 mo of age while mice with the podo-
cyte-specific knockout of myo1e showed first signs of glomerulosclerosis by 8 mo of age. The delayed onset and decreased severity of glomerulosclerosis were in contrast to the ultrastructural glomerular changes, which were highly prevalent by 4 mo even in the podocyte-specific model. Our findings suggest that while the loss of glomerular integrity as a result of myo1e removal from podocytes is rapidly manifested in proteinuria and defects in glomerular ultrastructure, histological signs of glomerulosclerosis develop only with nephrotic range proteinuria and/or with prolonged exposure to moderate proteinuria.

Multiple studies showed that the actin cytoskeleton plays a key role in maintaining podocyte foot process architecture and slit diaphragm organization. Actin cytoskeleton-associated proteins that have been linked to glomerular functions include actin bundling and regulatory proteins (α-actinin-4, synaptopo...

Fig. 5. Histological analysis of kidneys of myo1e-knockout mice. Paraffin-embedded kidney sections obtained from myo1e-null mice (Myo1e−/−), podocyte-targeted knockout mice (Myo1e−/−PodoCre, Myo1e−/−PodoCre), and control mice (Myo1e+/−PodoCre) at the indicated ages were stained using Masson Trichrome (A) or periodic acid Schiff (B). Four-month-old myo1e-null mice had multiple sclerotic glomeruli (arrows) and protein casts (*) while kidneys of 4-mo-old podocyte-targeted knockout mice appeared mostly normal. At 8 mo, kidneys of podocyte-targeted knockout mice contained a small number of sclerotic glomeruli (arrows) while control mouse kidneys appeared normal. Scale bars = 20 μm in A, 100 μm in B.

Fig. 6. Immunohistochemical analysis of podocyte markers in the glomeruli of myo1e-knockout mice. Cryosections of kidneys from control (Myo1e−/−) and podocyte-targeted knockout (Myo1e−/−PodoCre) mice were stained using antibodies against nonmuscle myosin 2a (A), α-actinin-4 (B), podocin (C), and integrin α3 (D). Expression and localization of these markers were similar in control and knockout mice. Bar = 10 μm.
MYO1E KNOCKOUT IN PODOCYTES CAUSES PROTEINURIA


