Genetic and Cellular Studies of The Podocyte in Focal Segmental Glomerulosclerosis

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Genetic and Cellular Studies of The Podocyte in Focal Segmental Glomerulosclerosis

by
Haiyang Yu

A dissertation presented to the Graduate School of Arts & Sciences of Washington University in partial fulfillment of the requirements for the degree of Doctor of Philosophy

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*Washington University in St. Louis*

*May 2015*
Dedicated to my parents and my wife.
ABSTRACT OF THE DISSERTATION

Genetic and Cellular Studies of the Podocyte in Focal Segmental Glomerulosclerosis

by

Haiyang Yu

Doctor of Philosophy in Biology and Biomedical Sciences

Molecular and Cellular Biology

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Professor Andrey Shaw, Chair

The podocyte forms the outer layer of the filtration barrier in the glomerulus to prevent albumin leakage. Podocyte damage leads to focal segmental glomerulosclerosis (FSGS), a leading cause of chronic kidney disease. The cause of the majority of FSGS cases is unknown and referred to as sporadic FSGS. Genetic studies have identified genes as monogenic causes of FSGS in patients with a strong family history, but these cases account for only a small proportion of the FSGS population. Whether genetic susceptibility contributes to sporadic FSGS and which cellular process in the podocyte initiates the pathogenesis of FSGS are important questions that remain to be elucidated. To answer these questions, my research followed two different lines of inquiry. I performed a genetic analysis of both familial and sporadic FSGS patients, and I investigated the role of the actin cytoskeleton in podocytes. Based on expression analysis, we identified a new FSGS susceptibility gene, ARHGAP24, and showed that it was mutated in a family with FSGS. Since ARHGAP24 functions to maintain high Rho and low Rac levels, my work suggested that this balance might be important in FSGS. Using an inducible transgenic
mouse model and multi-photon intravital microscopy, we validated that high activity of Rac1, one of the Rho family GTPases, is responsible for podocyte foot process effacement, increased membrane dynamics, and podocyte shedding into the urine, three important processes that lead to proteinuria and FSGS. By sequencing a large cohort of sporadic FSGS patients, I identified 16 potential FSGS susceptibility genes that were novel. Using a novel podocyte-specific indicible RNAi mouse model that I developed, four of these genes were validated. Some of these genes function as regulators of the actin cytoskeleton. Our genetic study further reinforces the role of actin cytoskeletal regulation in the pathogenesis of FSGS.
Chapter 1. Introduction

1.1 Podocytes form the outer layer of the kidney filtration barrier

The kidney is the organ where the urine forms. In adults, each kidney has 0.33~1.4 million glomeruli [1]. The primary function of the glomerulus is to filter small solutes in the blood and to generate the primary urine. During the filtration process, the blood flows into a tuft of capillaries from the afferent arteriole. Molecules that are smaller than 40 kilodaltons (kD), including glucose, urea, inorganic ions and water, pass through the filtration barrier, and enter the Bowman’s space (Figure 1.1A). The filtration barrier is a three-layer structure: the inner layer is the fenestrated endothelium; the middle layer is glomerular basement membrane (GBM), which is formed by negative-charged and glycosylated extracellular matrix proteins; the outer layer is formed by a kind of visceral epithelial cells, the podocyte (Figure 1.1B) [2].

Podocytes are unique-shaped epithelial cells that wrap around the glomerular capillaries. They extend major (primary and secondary) processes from the cell body. The tertiary processes (foot processes) interdigitate with those from neighboring podocytes to form a mesh-like network (Figure 1.1B and 1.2A). The foot processes are connected by an electron-dense, ~40nm-wide structure, the slit diaphragm, which composed of many adhesion proteins such as nephrin, NEPH1, P-cadherin, and FAT (Figure 1.2B). As terminal-differentiated cells, podocytes rarely proliferate post developmental period.

Pathological changes usually cause foot processes and the slit diaphragm to disappear.
The interface between two neighboring podocytes is flattened (Figure 1.3). This morphological abnormality is usually referred to as “foot process effacement”. The loss of the organized foot process–slit diaphragm meshwork is accompanied by the loss of the barrier function, and thus large size proteins in plasma could leak into the Bowman’s space, which further results in proteinuria. Albumin (MW ~70kD) leakage is measured as an indicator for the barrier function.

1.2 Focal Segmental Glomerulosclerosis

Focal Segmental Glomerulosclerosis (FSGS) is a glomerular disease that causes end stage renal disease. As one of the leading causes, FSGS accounts for about 4% of the patients who developed ESRD, and 40% of adult patients with nephrotic syndrome [3]. About ninety years ago, Theodor Fahr, a German pathologist, published the first drawing of FSGS glomerulus with great details [4]. Later, in 1957, A. Rich described that the cardinal characteristic of FSGS is progressive scaring in some glomeruli [5]. In the early stage of this disease, sclerotic tissue (scar) is found in a portion of the glomerulus (segmental), which only occurs in a subset of glomeruli (focal) (Figure 1.4). The scarred glomerulus often exhibits other pathological features such as mesangial hypercellularity and accumulation of foam cells in the capillaries [6]. Global sclerosis can be diagnosed as FSGS progresses [7]. Recent studies suggest that the incidence of FSGS has been increasing since the first clinical-pathological study in 1970s [3]. The major cell type that is affected in this disease is the podocyte. At electron microscopic level, the cardinal feature of FSGS is the effacement of podocyte foot processes. FSGS can be classified into familial FSGS and sporadic FSGS base on the patient family history of this disease. Sporadic FSGS is
more common in the patient population.

Although the details of the pathogenesis of FSGS are not fully uncovered, genetic studies of familial FSGS and animal models show that both genetic and environmental factors are important in this disease. Genetic approaches have identified 8 genes, which are all specifically expressed in the podocytes, as FSGS susceptibility genes: ACTN4 [8], ANLN [9], CD2AP [10–12], INF2 [13], MYH9 [14, 15], NPHS2 [16], and TRPC6 [17, 18]. In African American population, APOL1-G1 and -G2 alleles cause susceptibility to HIV-associated FSGS and hypertension-associated kidney disease [19, 20]. The combined heterozygosity of Cd2ap and Synpo induces FSGS in mice with incomplete penetrance: in 25% of the mice after 6 months of age [21]. This result suggests that genetic background could contribute to the susceptibility to FSGS. A nephrotoxic drug-induced FSGS model also supports this hypothesis. Adriamycin induces severe glomerular injury and FSGS in BALB/cJ, 129X1/SvJ and 129SvImJ mouse strains, but C57BL/6J strain is relatively resistant to this drug [22]. Recently Papeta et al discovered the genetic cause of this phenomenon, C6418T SNP in Prkdc gene, by meiotic mapping and genome sequencing [23]. This variant protects podocyte from mitochondria DNA depletion during Adriamycin treatment. Thus, there could be other triggers, like drug treatment or virus infection, which initiate the pathogenesis of FSGS.

Aging is also an important factor that contributes to the susceptibility of FSGS. Since podocytes are terminal differentiated and they do not undergo mitosis to generate new podocytes, the capacity of healthy podocytes to cover extra space is limited [24]. Each day, hundreds of podocytes are shed into the urine, and capillary surface left will be covered by the
remaining healthy podocytes through a process called podocyte hypertrophy [25]. Diseases and environmental factors that dramatically reduce podocyte number will be an important aspect of FSGS pathogenesis, because when the capillary surface exceeds the capacity of the remaining podocytes, the barrier system is permanently damaged and the damaged glomerulus will undergo sclerosis. Studies that use diphtheria toxin induced podocyte depletion in transgenic rats show that >40% of podocyte depletion induces segmental to global glomerulosclerosis [26].

1.3 The actin cytoskeleton supports the morphology and function of podocytes

Actin cytoskeleton is a dynamic cytoskeleton network in all eukaryotic cells. The polymerization and depolymerization of actin filament networks allow the cell to form dynamic structures that facilitate migration, cytokinesis, endocytosis, and other biological processes that are essential to maintain the normal functions of a cell. Actin filaments can also form stable bundles, which are named stress fibers, to facilitate the adhesion between cells and the extracellular matrix. Actin network is responsible for generating and transmitting force. Transverse arches and dorsal stress fibers are the structures that are involved in generating force on the cell body and transmit them to stress fibers and branched actin networks [27].

The fine structure of podocyte foot processes is supported by actin cytoskeleton network [28]. Transmission Electron Microscopy (TEM) shows that there electron dense actin bundles are enriched in the center of the foot processes, which are surrounded by loose cortical actin networks (Figure 1.5A,B). These actin bundles initiate from the major processes. Curved actin
bundles connect the straight bundles of two neighbor foot processes that initiate from the same major process [29] (Figure 1.5C). Details about how these structures form during podocyte differentiation are still not known. The actin cytoskeleton network in the foot processes connects the adhesive complex in the slit diaphragm, the focal contacts near the basement membrane, and the apical cell membrane of the foot processes.

Slit diaphragm is an adhesive complex that maintains interaction between adjacent podocytes. It connects to the actin cytoskeleton network inside the foot processes [30] (Figure 1.2B). Nephrin and Neph1 are transmembrane proteins that form the “zipper” like structure in the slit diaphragm. They have large extracellular domains that bind to another Nephrin or Neph1 molecules from the adjacent podocyte. Nephrin has a small intracellular domain that binds to several other cytoplasmic proteins like CD2AP and Nck [31–33]. Cd2ap directly bind to actin filaments, cortactin, and capping proteins [34–37]. Nck recruit WASp, another nucleation factor of actin filaments [38]. These proteins regulate branched actin network, and are believed to form a signal hub that maintains the normal morphology of foot processes.

Other actin-associated proteins also play important roles in podocytes. Alpha-actinin4 (encoded by ACTN4) connects between actin fibers to form stable bundles [39]. INF2 (Inverted formin 2) antagonizes the Rho-activated formin mediated actin polymerization and also the localization of formin in cultured podocytes [40]. Non-muscle myosin IIA is responsible for tension generation in response to the upstream signaling on the actin bundles, and regulates podocyte adhesion and migration [41]. Synaptopodin also functions to facilitate the polymerization of G-actin [42]. It rescues the tropomyosin defects in drosophila and human cells [43].
The focal complexes that mediate the podocyte-GBM interaction also connect to actin network in the foot processes. Focal complexes transmit the force that is generated by the cytoskeleton networks inside the cell to the extracellular matrix. They also can sense the force change outside the cell and convert it into intracellular signals. Integrin complexes are the transmembrane proteins mediating the inside-out and outside-in signals [44]. In podocytes, the most common integrin isoforms are α3β1 and αvβ3 [45, 46]. The extracellular domain of α3β1 and αvβ3 recognize laminin and collagen. The intracellular domains of multiple integrin heterodimers assemble the platform of the focal complexes. Focal adhesion molecules, including talin, zyxin, paxilin and vinculin form layers and connect to actin stress fibers. Kinases such as FAK, SRC and ILK are regulated by the focal complex proteins and regulate the actin cytoskeleton structure \textit{in vivo} [47].

1.4 Rho Family GTPases are the main regulators of Actin cytoskeleton

Rho family guanosine triphosphatases (hereafter called Rho GTPases) are a family of molecular switches that mainly regulate actin cytoskeleton structures [48]. The most studied members are Rac1, RhoA and Cdc42. Rho GTPases cycles between GTP-bound, active conformation and GDP-bound, inactive conformation. GTP-bound Rho GTPases activate the downstream pathway by interacting with their effector molecules (Figure 1.6A). The C-terminus of most Rho GTPases is modified by farnesyl or geranylgeranyl isoprenoid lipid, which is required for membrane targeting. Active Rac1 recruits WAVE complex to the plasma membrane and induces actin polymerization into branched network. Cells with high Rac activity forms large
lamellipodia (Figure 1.6B,C). GTP-bound RhoA, on the other hand, activates formin mediated actin polymerization, resulting in long linear actin fibers and bundles (Figure 1.6B,C). Active RhoA induces the assembly of stress fibers. Active Cdc42 induces finger-like extensions through recruiting Wiskott–Aldrich Syndrome protein (WASp) to the plasma membrane and release its auto-inhibitory conformation (Figure 1.6B,C). WASp further recruits Arp2/3 complex and assembles thin, linear actin filaments, which protrude from the plasma membrane, and form finger-like structures.

Direct modulation of the activity of Rho GTPases affects podocyte morphology and function [49–51]. Podocyte-specific expression of CA-RhoA induced late onset of proteinuria and FSGS [49]. Podocyte specific deletion of Cdc42 in mice causes congenital nephrotic syndrome and foot process effacement [50]. However, deletion of either Rac1 or RhoA in podocytes does not induce any defects in mice. Podocyte specific deletion of Rac1 in mice causes resistance to protamine sulfate perfusion-mediated transient podocyte damage and foot process effacement, but susceptibility to a chronic model of podocyte injury in UNX/DOCA-salt-hypertensive mouse model [52].

1.5 Regulators of Rho GTPases

Three families of regulators mainly control the activity of Rho GTPases. Two classes of Rho GTPase regulators control the processes of cycling between GTP- and GDP-bound states: Guanine nucleotide exchange factors (GEFs) and GTPase-activating proteins (GAPs) (Figure 1.6A) [53]. The Rho GTPases are slow GTPases by themselves. The exchange between
GDP-bound and GTP-bound forms is also ineffective. GEFs facilitate GTP loading, and increase the level of active Rho GTPases. GAPs enhance the catalytic activity of Rho GTPases, converting the active Rho family GTPases to the inactive state. A third family of regulatory proteins are the guanine nucleotide dissociation inhibitors (RhoGDIs) (Figure 1.6A), which sequester Rho GTPases at GDP-bound state, and pull it out of the cycling pool. RhoGDIs are shown as “chaperones” for small GTPases because they stabilize GDP-bound state, and also mediate the translocation of Rho GTPases between different membrane compartments [54, 55].

Although the exact physiological function of the Rho GTPase regulators has not been extensively studied in podocytes, several studies indicate their crucial roles in regulating podocyte morphology and function. RhoGDI-alpha knockout mice (Arhgdia−/−) exhibit congenital nephrotic syndrome, and the knockout mice have high Rac activity in the kidney [56]. Tyrosine phosphorylation on nephrin endoplasmic tail could also regulate p190RhoGAP through Rac [57]. Human protein atlas project has also identified GAPs and GEFs that are highly expressed in podocytes, including ARHGAP12, ARHGAP28, ARHGAP35, SRGAP2, ARHGEF11, and ARHGEF12 [58].

Some podocyte specific and/or FSGS susceptibility genes also regulate the activity of Rho GTPases. Nephrin controls the Rac activity through PI3K-p85 signaling pathway [57]. Synaptopodin, another podocyte-specific protein, protects Rho from degradation by Smurf-mediated ubiquitination, and inhibit the Cdc42-IRSp53 interaction [59, 60]. Mutations in TRPC6, encoding a calcium channel, are associated with familial FSGS [17, 18]. Recently studies showed that TRPC6 forms complex with RhoA, and TRPC5 forms complex with Rac1.
Calcium influx through TRPC6 increases active RhoA, while through TRPC5 increases active Rac1. PLCE1, a risk gene for steroid resistant nephrotic syndrome and FSGS, is responsible for efficient activation of Rac1 [62].

1.6 Large-scale genetic studies discover FSGS susceptibility genes

Next-generation sequencing (NGS) technologies, revolutionary sequencing methods developed in early 2000s, allow researchers to access to personal genome in a faster and more economic than traditional sanger sequencing method [63]. These methods read millions of sequencing reactions simultaneously by high throughput monitoring approaches [64]. Using NGS methods, researchers can identify specific genetic variants in a large cohort of patients. These methods have been applied to investigate cancer genomes [65], to identify susceptibility variants that associate with diseases such as autism and diabetes [66, 67], and to map the epigenetic modification loci in the genome. Since the cost of whole genome sequencing for each patient is still high, Hildebrandt’s group combines linkage analysis of familial nephrotic syndrome with NGS recently identified several new genes that could cause FSGS, including ADCK4 [68], ARHGDIA [69], and EMP2 [70].

Genome-wide association studies (GWAS) have been widely used to identify the linkage between common variants that associate with diseases before NGS technologies became popular. This method scan hundreds of thousands common DNA variant in human. Several GWAS have identified 24 susceptible loci for chronic kidney disease, including SHROOM3, UMOD, and DACH1 [71, 72]. GWAS method was also applied to study HIV-associated FSGS in
African American population, and identified genetic locus between \textit{APOL1} and \textit{MYH9} that are strongly associated with HIV-associated FSGS [14]. G1 and G2 variants of \textit{APOL1} are the coding variants for FSGS susceptibility [19, 20].

\textbf{1.7 Current mouse genetic models of podocyte diseases}

Using \textit{in vitro} models such as podocyte cell culture to study the function of podocytes is limited since cultured podocytes do not generate foot processes. Thus \textit{in vivo} models are necessary to investigate and truly understand podocyte functions. Currently, transgenic and knockout mouse models are the most widely used \textit{in vivo} models for podocyte diseases. Rat models are also commonly used and they are more susceptible to kidney damage than mouse models. However, rats require more space and longer breeding period, and also the transgenic/knockout tools in rat models are not as efficient as in mouse models. Knockout mouse models have shown that many podocyte-specific genes and FSGS disease genes are essential for maintaining the podocyte foot process structure \textit{in vivo}, including \textit{Nphs1} [73], \textit{Nphs2} [74], \textit{Neph1} [75], \textit{Actn4} [76], and \textit{Cd2ap} [77]. Loss of function of some other genes in mice does not cause any podocyte dysfunction, but increases the susceptibility to artificial kidney injury models, such as \textit{Synpo} [21], \textit{Trpc6} [78], and \textit{Myh9} [79]. The combinations of heterozygosity in \textit{Cd2ap/Synpo} and \textit{Cd2ap/Fyn} also induce proteinuria and podocyte foot process effacement, the early signs of FSGS in aged mice, indicating there could be genetic complexity in FSGS patients [21].
1.8 Intravital imaging by multi-photon microscopy

Since podocyte cell culture cannot recapitulate the foot process architecture in petri dishes, intravital imaging is the only option to investigate the podocyte cell dynamics at the physiological conditions. Currently multi-photon microscopy (MPM) is the best method to perform non-invasive live imaging *in vivo*. This method based on a theory that multiple low-energy, long wavelength photons can excite a fluorophore simultaneously (Figure 1.7A). The interaction between multiple photons and the fluorophore will induce sequential electronic transition that equals to a single high-energy photon [80]. The first MPM microscope was invented by Denk *et al* in 1990 [81]. Since low-energy, long-wave length photon is the source of excitation, MPM achieves three following advantages: first, it penetrates tissue up to several hundred micrometers, which is much deeper than another microscopy system; second, it reduces phototoxicity and bleaching of the fluorophores; third, it reduces out-of-focus background signal because fluorophores are excited only at the laser beam focus (Figure 1.7B) [82, 83]. Since the excitation is not linear, second harmonic signal can be generated to visualize membrane structures like kidney capsule and fascia by visualize their native fluorescence [84].

All the features make MPM an excellent approach to track podocytes *in vivo*. Using MPM, Hackl *et al* showed podocyte migration in response to kidney injuries in podocin-GFP and podocin-CRE confetti mice [85]. Endlich *et al* observed podocyte migration during the development of zebrafish glomeruli. They also observed that zebra fish podocytes and their
branch patterns are stationary at physiological conditions [86]. Peti-Peterdi’s group also reported increased calcium waves in podocyte during unilateral ureteral obstruction induced kidney injury by using podocyte specific GCaMP mice [87].

1.9 Conclusions

The cell body of podocytes and the slit diaphragm between the interdigitating foot processes from adjacent podocytes form the outer layer of the filtration barrier to prevent albumin leakage into the primary urine. The organized structures of foot process and the slit diaphragm are the indicators of the intact barrier function. In podocyte diseases, the effacement of foot processes and the loss of slit diaphragm are often observed. These pathological changes are also correlated with albuminuria. Actin network is the major cytoskeleton structure inside the foot processes. Cortical Actin network connects the cell-cell adhesion of adjacent podocytes (slit diaphragm) to the actin bundles in the center of the foot processes, which is thought to be the cytoskeletal support of the foot process. Actin bundles are also connected to the focal complexes that mediate the interaction between the podocytes and GBM. As the major regulators of actin cytoskeleton, Rho family small GTPases are very important in maintaining foot processes and slit diaphragm. Previous studies have indicated that dysregulation of the activities of Rho GTPases could cause foot process effacement and glomerulosclerosis. However, it is not fully understood about which small GTPase play the most important role in the podocyte damage.

FSGS, a leading cause of end stage renal failure, is one of the diseases that caused by podocyte damage. It remains unknown about how much genetic risk causes susceptibility to
FSGS. FSGS can be classified as familial FSGS and sporadic FSGS based on whether another family member is diagnosed with the same disease or not. Genetic studies of familial FSGS cases have identified many FSGS causing genes, most of which are components of the slit diaphragm, and/or regulate the actin cytoskeleton structure. Gain or loss of function mutations of FSGS genes could cause actin cytoskeleton remodeling in the podocytes, which could be the reason of foot process effacement. However, FSGS genes do not explain the sporadic form of this disease, because they are rarely mutated in sporadic cases. Whether there is a genetic factor and how much of the genetic risk is in the sporadic FSGS remain to be investigated.
1.10 Figures

![Diagram of the glomerulus structure](image)

**Figure 1.1 The structure of the glomerulus**

(A) The image showed the composition of the glomerulus. Afferent arteriole branched into capillaries ball inside the Bowman’s capsule. During the filtration process, water and small solutes can penetrate through the filtration barrier and become the glomerular ultrafiltrate. The rest of the blood including blood cells and proteins remain in the capillary loops, which combine to form the efferent arteriole. (B) The structure of the filtration barrier. The capillaries inside the Bowman’s capsule are fenestrated. They are covered by podocytes. Between foot processes of the podocyte, big transmembrane proteins form an adhesive structure called slit diaphragm (filtration slits). (Adjusted from *Human Physiology*, Chapter 12, [https://humanphysiology2011.wikispaces.com/12.+Urology](https://humanphysiology2011.wikispaces.com/12.+Urology), by Rausch A and Kortleever C)
Figure 1.2 The podocyte and the slit diaphragm
(A) Scanning microscopy of the podocyte. FP: foot processes; P: primary processes. (B) A cartoon of the cross-section view of foot processes and slit diaphragm. Important proteins and their locations are also shown [88, 89].

Figure 1.3 Foot process effacement
(A) A cartoon showed the effaced podocytes lose their foot processes. The effaced area looks like lamellipodia. (B) Transmission electron microscopy shows normal and effaced foot processes.
The upper glomerulus shows features of glomerulosclerosis on the left lobe. The lower glomerulus is normal. The red-staining area on the left lobe of the upper glomerulus shows sign of scarring (sclerosis). (A mouse kidney was stained by a method called periodic acid-Schiff stain)

Figure 1.5 The actin cytoskeleton in the podocyte foot processes
(A) An cross-section image of a foot process imaged by transmission electron microscopy. The arrow showed the actin bundle in the center of the foot processes, the star showed the loose actin network at the bottom of the foot processes and surrounding the actin bundle. The arrowheads show the slit diaphragm. (B) A cartoon of a foot process. AB: Actin bundles, CAN: cortical actin network, SD: slit diaphragm, GBM: glomerular basement membrane, and En: endothelial cells. (C) A bird view image of foot processes. The open arrow shows curved actin bundles that connects adjacent foot processes extended from the same podocyte; the closed arrows show the straight actin bundles inside the foot processes. This figure is adapted from Sakai et al [29].
Figure 1.6 Rho Family GTPases: regulation and function

(A) The regulated catalytic cycle of Rho family GTPases. Lipid modification allows Rho GTPases attach to the membrane structures in the cell. GEFs facilitate the exchange of GDP to GTP and convert GDP-bound Rho GTPases to GTP-bound form. GTP-bound Rho GTPases activate the downstream effectors through allosteric mechanisms. GAPs stimulate the GTPase activity of Rho family GTPases and convert the GDP-bound form to GDP bound form. RhoGDI proteins bind the GDP-bound form, and pull it out from the membrane structure. (B, C) Active Rho stimulates stress fiber formation through mDia, activate Rac stimulates lamellipodia formation through WAVE complexes, and active Cdc42 stimulates filopodia formation through WASP. This figure is adapted from Tubulewicz et al. [90] and http://www.mechanobio.info/modules/go-0051893
**Figure 1.7 The mechanism and advantage of multi-photon microscopy (MPM)**

MPM reduces the fluorescent background compared to single photon microscopy. This figure is adapted from [http://cleoqels2010.blogspot.com/2010/05/expo-idea-generation-and-multiphoton.html](http://cleoqels2010.blogspot.com/2010/05/expo-idea-generation-and-multiphoton.html)
Chapter 2. Arhgap24 Inactivates Rac1 In mouse Podocytes, and A mutant Form is Associated with Familial Focal Segmental Glomerulosclerosis

2.1 Abstract

Podocyte has a complex actin-based cytoskeleton that maintains efficient barrier function of glomeruli. Disruption of components of the actin cytoskeleton results in podocyte damage and cell loss, which may lead to a prototypic injury response called focal segmental glomerulosclerosis (FSGS). From genes that are highly expressed in mouse podocytes, we identified a RhoA-activated Rac1 GTPase-activating protein, Arhgap24, which was upregulated in podocytes as they differentiated, both in vitro and in vivo. Decreased Arhgap24 expression results in high level of active Rac1 and Cdc42, which influenced the cell shape and membrane dynamics. Consistent with a role for Arhgap24 in maintaining normal podocyte functions in vivo, we identified a mutation in Arhgap24 that impaired its Rac1-GAP activity by sequencing FSGS patients and that was associated with disease in a family. Thus, Arhgap24 contributes to the careful balancing of RhoA and Rac1 signaling in podocytes, the disruption of which may lead to kidney disease.

2.2 Introduction

The kidney filters plasma and reabsorb salts and nutrients to maintain the appropriate extracellular environment. The proximal component of the nephron, the glomerulus, is the primary filtration barrier that prevents the loss of serum proteins into the primary filtrate. The
glomerular filtration barrier consists of fenestrated endothelial cells, a thick glomerular basement membrane (GBM), and specialized epithelial cells (podocytes) arranged in series. Diseases affecting the filtration barrier, especially of the GBM or the podocyte, result in the leakage of serum proteins into the urine, progressive damage to the glomerulus, and loss in renal function [91].

The podocyte has a complex cellular architecture composed of an octopus-like cell body that attaches to the GBM through actin-based foot processes [88]. In kidney diseases that are associated with proteinuria, such as focal segmental glomerulosclerosis (FSGS) and minimal change disease, podocytes rearrange their actin cytoskeleton network, which results in retraction or effacement of foot processes [92]. While it is unclear how this change leads to the leakage of serum proteins, foot process effacement appears to be a key step in the breakdown of the filtration barrier. Studies to understand the molecular basis of foot process effacement in vitro have shown that, in response to stress, podocytes switch from a RhoA-dependent stationary state to a Cdc42- and Rac1-dependent migratory state [28, 93]. These studies suggest the intriguing possibility that altered membrane dynamics and increased cell motility are the mechanisms underlying foot process effacement in vivo.

Since podocyte membrane reorganization is a common feature of proteinuric kidney diseases, we sought to understand the regulation of membrane dynamics of these cells. Using an in vitro model of podocyte differentiation, we found that podocytes reduced their membrane ruffling activity as they were differentiated. We found that decreased membrane ruffling in differentiated podocytes was dependent on the presence of a GTPase-activating protein (GAP),
Arhgap24. Previous works from Stossel and colleagues have shown that Arhgap24 (also known as Filamin A-binding RhoGAP [FilGAP]) is a GAP for Rac1, and it binds to branched actin network and suppresses lamellipodia formation and cell spreading downstream of RhoA signaling [94]. Their work also shows that the highest level of Arhgap24 transcript is in the kidney. Here we show that Arhgap24 was specifically expressed in podocytes in the kidney. Its expression increased as these cells differentiate in vivo. The ARHGAP24 gene is highly conserved, implying an important role for the gene product. We sequenced the DNA from patients with FSGS, and identified a loss-of-function mutation in the ARHGAP24 gene in kindred with familial kidney disease. Taken together, these results suggest that Arhgap24 controls the RhoA-Rac1 signaling balance in podocytes that could be dysregulated in proteinuric kidney diseases, such as FSGS.

2.3 Methods

2.3.1 Podocyte cell culture

Generation and propagation of conditionally immortalized murine podocyte cell lines has been described previously [95, 96]. Briefly, podocytes were propagated on collagen I–coated dishes at 33°C (permissive temperature) in RPMI supplemented with 10% fetal bovine serum (FBS) and 10 U/ml of recombinant mouse interferon-γ (IFN-γ, a gift from Robert Schreiber, Washington University School of Medicine). To induce differentiation, the medium was changed to RPMI with 5% FBS without IFN-γ, and the cells were shifted to 37°C (nonpermissive temperature) for 7 to 14 days. Under these conditions, cells underwent growth arrest, increased
in size, and developed elongated cell processes.

For live cell imaging experiments, podocytes were stably transduced with YFP-actin by lentivirus transduction, and purified by automated cell sorting. Control and knockdown cell lines were generated using a bicistronic lentiviral vector incorporating the targeting shRNA and YFP-actin expressed downstream of an internal ribosomal entry site. The control knockdown sequence targeted the firefly luciferase gene (Fluc). The 2 Arghap24 knockdown constructs targeted the sequences 5′-TTAAGGAGCTAATGAAACA-3′ (line 451) and 5′-TAACGATGGTCATAAGAAA-3′ (line 756), respectively. Stably transduced cell lines were generated by automated cell sorting for YFP expression.

2.3.2 RNA isolation

Isolation of primary mouse podocytes using Dynabead perfusion and flow cytometric cell sorting has been described previously [97, 98]. RNA was extracted from 6,000 primary podocytes and from cultured podocytes grown at the permissive or nonpermissive temperatures using an RNeasy Kit (Qiagen) following the manufacturer’s protocol. RNA quality was verified by gel electrophoresis and to ensure that the 260/280 nm absorbance ratio was greater than 1.8.

2.3.3 Arhgap24 antiserum production

Amino acids 390–604 of isoform 1 (NCBI accession no. NP_083546) of mouse Arhgap24 were cloned downstream of glutathione-S-transferase (GST) in the pGEX4T-1 expression vector. This portion of Arhgap24 is well conserved across species and lies downstream of the GAP domain in both isoforms of Arhgap24. GST-tagged Arhgap24 was expressed in BL21 (DE3) pLysS E. coli. After induction with IPTG and sonication of bacteria, soluble GST-Arhgap24 was
batch purified using glutathione-agarose beads. GST-Arhgap24 was eluted with reduced glutathione and dialyzed against PBS to remove excess glutathione. This antigen was emulsified in complete Freund’s adjuvant (Sigma-Aldrich) and was used to immunize rabbits and Armenian hamsters. Hamster handling and immunization were performed by the Washington University School of Medicine Hybridoma Center. Specificity for Arhgap24 was confirmed by immunoblotting FLAG-tagged Arhgap24 transfected into HEK293 cells. Specific signal in immunoblotting and staining experiments was confirmed by quenching of signal with antiserum preincubated with antigen.

2.3.4 Confocal imaging

Podocytes grown on sterilized collagen I–coated coverslips at 33°C or 37°C were fixed with 4% PFA in PBS for 10 minutes. Cells were then blocked and permeabilized for 1 hour with PBS with 2.5% FBS and 0.1% saponin. Primary rabbit anti-Arhgap24 antiserum (1:300) in blocking buffer was applied to the cells for 1 hour at room temperature. After four 5-minute washes with blocking buffer, coverslips were mounted using ProLong Antifade mounting medium (Invitrogen) according to the manufacturer’s protocol. Images were captured with confocal settings using an Olympus FluoView FV1000 microscope. A similar protocol was used to stain Arhgap24 in formalin-fixed paraffin-embedded mouse kidney after antigen retrieval. Podocytes were stained using a mouse anti-synaptopodin monoclonal antibody (a gift from Peter Mundel, Massachusetts General Hospital, Boston, Massachusetts, USA) [99]. For wound healing experiments, differentiated knockdown podocytes were plated to confluence on collagen I–coated coverslips. A scratch was created using a sterile 200 µl pipette tip. Loosely adherent cells were washed
away by 3 vigorous washes with PBS. The cells were transferred to culture medium, and, at various time points, coverslips were harvested and fixed in 4% PFA in PBS. Cells were identified by nuclear staining with DAPI (4’,6-diamidino-2-phenylindole).

2.3.5 Tissue isolation and immunoblotting

Whole mouse tissues (~200 mg) were snap frozen on dry ice and homogenized in cold lysis buffer containing 1% NP-40 and protease inhibitors using a dounce homogenizer. Post nuclear supernatants were loaded to equalize actin levels by immunoblotting. Glomerular isolation/enrichment experiments using magnetic particles were performed as described previously [97].

2.3.6 Live cell imaging and kymograph analysis

Podocytes that were lentivirally transduced with YFP-actin were cultured in glass bottom dishes at 33°C or differentiated at 37°C (nonpermissive condition) for 7 to 14 days and then imaged. Similarly, YFP-actin transduced Arhgap24 knockdown (lines 451 and 756) and control knockdown (Fluc) cell lines were differentiated on glass bottom dishes for 7 to 14 days before imaging experiments. Sequential images were captured every 15 seconds for a 20-minute duration using an Olympus FluoView FV1000 microscope. Movies were assembled using Olympus Fluoview software. The Multiple Kymograph plug-in of ImageJ (http://rsbweb.nih.gov/ij/) was used to generate kymographs at 3 different locations of maximal membrane ruffling for each cell imaged [100]. For each kymograph, up to 13 actin spikes were measured, and the average length was computed as described previously [101]. A value of 1 was assigned to kymographs without a measurable actin spike. Ruffling movies were generated by H. Suleiman, and the
kymographic analysis was performed in a blinded fashion by S. Akilesh. Images were assembled in Adobe Photoshop.

2.3.7 Sequencing

DNA was obtained from affected (n = 310) and control (n = 180) individuals. DNA from 96 of the control individuals was purchased from the Coriell Institute. PCR primer pairs were designed to completely sequence exons and exon-intron junctions of ARHGAP24. Bidirectional sequencing using ABI Big Dye 3.1 sequencing chemistry on an ABI PRISM 3730 sequencing platform (Applied Biosystems) was performed. Sequences were aligned using Sequencher software (Gene Codes) and manually verified. The Q158R variation was determined by A.S. Shaw’s laboratory and verified by M.P. Winn’s laboratory. In addition, the exon incorporating this variation was sequenced in an additional 554 control chromosomes to rule out the possibility that it represented a low-frequency noncausal variation.

2.3.8 Haplotype analysis

MSM primer sequences for markers D4S1538, D4S1534, D4S2409, and D4S2460 were obtained from UniSTS (http://www.ncbi.nlm.nih.gov/unists) or designed with Primer3 software (http://frodo.wi.mit.edu/primer3/). Carboxyfluorescein succinimidyl ester–labeled (FAM-labeled) oligonucleotides were synthesized by Invitrogen Life Technologies and were run on the 3730 DNA Analyzer (Applied Biosystems) and analyzed using GeneMapper Software v4.0 (Applied Biosystems). The analysis was carried out by visual inspection, assigning the most likely linkage phase by minimizing the number of recombinants in the pedigree.

2.3.9 Arhgap24 dimerization assay
FLAG- or GFP-tagged wild type or Q156R Arhgap24 were cotransfected into HEK293 cells. Cell lysates were immunoprecipitated with M2 mouse anti-FLAG antibody (Sigma-Aldrich) with protein A conjugated to sepharose beads. The immunoprecipitates were resolved by SDS-PAGE and then immunoblotted with JL-8 mouse anti-GFP antibody (Invitrogen).

2.3.10 Rac1, Cdc42, and RhoA pull-down assays

The GST-tagged Pak1-PBD used for the pull-down assays was expressed in BL21(DE3) E. coli. The recombinant protein was bound to glutathione-agarose beads stored at –80°C until use. For the active Rac1 pull-down assay, HEK293T cells were seeded in 6-cm dishes and transfected at 80%–90% confluence using Lipofectamine 2000 reagent according to the manufacturer’s protocol (GIBCO BRL). Cells were transfected with FLAG-tagged wild-type or mutant Q156R Arhgap24 or a titration of the 2 constructs, keeping the total DNA transfected constant. Twenty-four hours after transfection, the cells were lysed, and a sample of the lysate was retained for measurement of total Rac1 and FLAG-Arhgap24 levels. Equal volumes of the remaining lysates were incubated with GST-PBD bound to glutathione-agarose beads. Rac1-GTP bound to the beads (active Rac1) was eluted with Laemmli sample buffer and separated on a 12% polyacrylamide gel. After transfer to a nitrocellulose membrane, active and total Rac1 were detected with the 23A8 mouse anti-Rac1 monoclonal antibody (Upstate Biotechnology). FLAG-tagged Arhgap24 was detected with the M2 mouse anti-FLAG antibody. For Cdc42 activity assays, active Cdc42 was immunoprecipitated with GST-PBD bound to glutathione-agarose beads and detected with B-8 mouse anti-Cdc42 (Santa Cruz Biotechnology Inc.). For RhoA activity assays, active RhoA was immunoprecipitated with GST-Rhotekin bound
to glutathione-agarose beads and detected with 26C4 mouse anti-RhoA (Santa Cruz Biotechnology Inc.). The immunoblot signal was detected using a LI-COR Odyssey Imaging System using their infrared dye-labeled secondary reagents. Data are representative of at least 3 independent experiments.

2.3.11 Statistics

Data are represented as mean ± SD. In all cases, group differences were assessed by ANOVA with post-test correction (Bonferroni-Holm). A P value of less than 0.05 was considered significant.

2.3.12 Study approval

Patient and control DNA samples were obtained after written informed consent and with local Institutional Review Board approval (Washington University School of Medicine, Duke University Medical Center, Université René Descartes, Brigham and Women’s Hospital, and the NIDDK). All animal experiments were conducted with approval of the Washington University Animal Care and Use Committee.

2.4 Results

2.4.1 Differentiated podocytes reduce ruffling of their cell membranes

Podocytes can be propagated in vitro by conditional expression of a temperature-sensitive SV40 large T antigen at the permissive temperature of 33°C. Shifting the cells to the nonpermissive temperature (37°C) induces destabilization of the large T antigen, growth arrest, and morphologic changes that mimic podocyte differentiation in vivo [95]. During development,
podocytes reorganize their cell membranes from broad sheets into sieve-like foot processes [88]. Conversely, in proteinuric diseases, podocytes lose this membrane complexity and exhibit effacement of their foot processes [92]. Therefore, we hypothesized that the membrane dynamics of podocytes would be regulated in this in vitro model of cell differentiation. When undifferentiated podocytes were cultured at the permissive temperature, they exhibited highly ruffled plasma membranes (Figure 2.1A). In contrast, the plasma membranes of the differentiated podocytes had a very smooth, flat appearance. In order to quantitate these membrane dynamics, we performed time-lapse imaging of live podocytes transduced with yellow fluorescent protein–actin (YFP-actin). While undifferentiated podocytes rapidly ruffled their membranes, differentiated podocytes had reduced membrane motility, correlating with the reduced frequency of ruffled cell membranes (Supplemental Videos 1 and 2; supplemental material available online with this article; doi: 10.1172/JCI46458DS1). This difference is also apparent by kymographic analysis (Figure 2.1A and 2.1B).

2.4.2 Podocytes upregulate Arhgap24 when they differentiate

Next, we evaluated the expression of the small G proteins and their regulators that might control membrane motility in this in vitro model of podocyte differentiation. We isolated RNA from undifferentiated and differentiated mouse podocytes and performed gene expression analysis using microarrays. Our cultured podocytes expressed genes for several known podocyte-specific proteins, such as Wilms tumor protein 1 (Wt1), podocin (Nphs2), CD2-associated protein (Cd2ap), podocalyxin (Podxl), synaptopodin (Synpo), α-actinin-4 (Actn4), and to a lower extent nephrin (Nphs1). We focused our analysis on the expression patterns of known regulators of the
actin cytoskeleton and membrane dynamics, such as GAPs and guanine nucleotide exchange factors (GEFs) for Rac1, RhoA, and Cdc42 (a list of these genes is available as a supplemental data table in the Gene Expression Omnibus (GEO) database, series GSE23856; http://www.ncbi.nlm.nih.gov/gds/). Of these known GAPs and GEFs, the message for Arhgap24 was highly upregulated when podocytes were differentiated in vitro. Arhgap24 was an intriguing gene, because previous work from Stossel and colleagues had shown high transcript levels in the kidney [94]. However, the cell expressing Arhgap24 within the kidney was unknown. We found that RNA from in vivo–isolated podocytes also had a high level of Arhgap24 transcript [96]. Next, we examined Arhgap24 gene expression in publicly available microarray data (the GenitoUrinary Molecular Anatomy Project; www.gudmap.org) generated from microdissected glomeruli isolated at various stages of kidney development. The Arhgap24 message was upregulated as glomeruli progressed from the E12.5 renal vesicle stage to the E15.5 S-shaped body to the almost mature E15.5 renal corpuscle (Figure 2.2A).

We confirmed the microarray results by quantitative RT-PCR. Arhgap24 transcript levels increased ~70 fold when podocytes were differentiated (Figure 2.2B). To confirm upregulation at the protein level, we generated Arhgap24-specific antiserum in Armenian hamsters and rabbits. Preabsorption of these antisera with Arhgap24 antigen abolished specific signal (Supplemental Figure 2.1). Podocytes upregulated a 95-kDa band corresponding to the Arhgap24 protein when they were differentiated in vitro (Figure 2.2C). The lower band seen in the undifferentiated podocytes (at approximately 50 kDa) did not correspond to the predicted molecular weight of known splice isoforms of Arhgap24. This 50kDa bind likely represents a specific degradation
product of Arhgap24, perhaps due to increased turnover of the protein in undifferentiated podocytes. Confocal imaging of podocytes showed that Arhgap24 was enriched in structures at the base of the cell, and the staining intensity at these sites increased with differentiation (Figure 2.2D). Since Arhgap24 has been described to localize to focal adhesions [94], we verified that these structures were in fact focal adhesions by co-labeling with vinculin (Figure 2.2E). These experiments showed that Arhgap24 is highly expressed in the focal adhesions of cultured podocytes and is upregulated as these cells are differentiated.

2.4.3 Arhgap24 is expressed in kidney podocytes \textit{in vivo}

Next, we tested Arhgap24 expression and localization in murine tissues and its specific localization within the kidney. Immunoblotting of lysates from several tissues showed that the highest level of full-length Arhgap24 protein (95-kDa band) was in the kidney. Again, the approximately 50-kDa putative breakdown product was seen in lysates from the brain, kidney, and liver. We then evaluated whether Arhgap24 was expressed in the glomeruli of the kidney. We trapped magnetic particles within the glomeruli by beating heart perfusion and then used magnetic separation to isolate whole glomeruli to more than 95% purity (Figure 2.3B). Arhgap24 was enriched in the glomerular fraction (verified by podocin immunoblotting) compared with the flow through fraction that contained mostly tubule fragments (Figure 2.3C). Next, we stained sections of mouse kidney and detected the greatest signal within glomeruli (Figure 2.3D). This signal was specific since it was completely abolished by preabsorption of the antiserum with Arhgap24 antigen (Data not shown). Within the glomeruli, the Arhgap24 signal colocalized with the podocyte marker synaptopodin, confirming that Arhgap24 is expressed in podocytes \textit{in vivo}.
2.4.4 Arhgap24 knockdown restores membrane ruffling in differentiated podocytes

Having established that Arhgap24 is expressed in podocytes in vitro and in vivo, we next asked whether it was responsible for the dramatic decrease in membrane ruffling that we observed in differentiated podocytes (Figure 2.1). Membrane ruffling is dependent on the activity of the small G protein Rac1 [102]. Rac1 cycles between an active, GTP-bound state and an inactive, GDP-bound state. When activated by GEFs, Rac1 interacts with multiple downstream effectors to stimulate actin nucleation and branching required for lamellipodia formation and membrane ruffling. Subsequently, Rac1 can be inactivated by GAPs that stimulate the intrinsic GTPase activity of Rac1, resulting in the hydrolysis of GTP to GDP. Previous work has shown that Arhgap24 is a GAP protein that inactivates Rac1 [94]. Since Arhgap24 is upregulated in podocytes as they differentiate, it was a logical candidate to inactivate Rac1 and thereby slow membrane ruffling in differentiated podocytes.

We investigated whether Arhgap24 upregulation was responsible for the reduced Rac1 activity and membrane motility of differentiated podocytes. We generated podocyte cell lines (lines 451 and 756) with 2 different Arhgap24 lentiviral knockdown constructs that coexpressed the YFP-actin reporter. Compared with that of the control vector (Fluc), the podocyte cell lines transduced with the 2 knockdown constructs resulted in approximately 85% (line 451) or 60% (line 756) downregulation of the Arhgap24 protein in differentiated podocytes (Figure 2.4A). After differentiation, the control and 2 Arhgap24 knockdown cell lines were assayed for their membrane motility as before. Compared with that of the control knockdown, Arhgap24
knockdown resulted in increased membrane motility in differentiated podocytes (Figure 2.4B and Supplemental Videos 3–5). Kymograph analysis showed that Arhgap24 knockdown significantly increased membrane motility (Figure 2.4C).

Since Arhgap24 inhibits Rac1 and Cdc42 activity, we asked whether Arhgap24 knockdown resulted in increased levels of active Rac1 and Cdc42 in differentiated podocytes. We measured active Rac1, Cdc42, and RhoA by standard pull-down assays and found that the Arhgap24 knockdown cell lines had higher levels of active Rac1 and Cdc42 compared with those of the control knockdown cell line (Figure 2.5A). Interestingly, Arhgap24 knockdown did not affect RhoA activity. Another assay of Rac1 activity is epithelial monolayer wound closure. When a confluent monolayer of differentiated control knockdown podocytes was scratched/wounded, the cells migrated into the gap but did not completely close the wound in 24 hours. In contrast, both Arhgap24 knockdown cell lines showed accelerated wound closure kinetics, consistent with higher Rac1 activity (Figure 2.5B). Thus, Arhgap24 knockdown increases levels of active Rac1 and Cdc42. Two measures of increased Rac1 activity, membrane ruffling and epithelial wound closure, are enhanced when Arhgap24 levels are reduced in podocytes, consistent with its function as a Rac1-GAP.

2.4.5 Sequencing ARHGAP24 in patients with FSGS identifies several nonsynonymous changes in a highly conserved gene

Given the podocyte-specific pattern of expression of Arhgap24 and its effects on membrane dynamics and epithelial wound healing, we next asked whether variations in ARHGAP24 were associated with kidney disease. The ARHGAP24 gene consists of at least 8
exons (depending on the splice isoform) that span approximately 500 kilobases on the long arm of human chromosome 4. Complete exon sequencing of the ARHGAP24 gene in 310 patients (620 chromosomes) with biopsy-proven FSGS and 180 controls (360 chromosomes) identified 13 nonsynonymous sequence variations in the 2 main splice isoforms of Arhgap24 (Table 2.1). Two of these (P417A and F539L) were found in both patients and controls and also in the 1000 genomes database (www.1000genomes.org) (Table 2.2). However, seven of the nonsynonymous changes were only found in patients and not in any of the controls. These results suggest that variation in the ARHGAP24 gene may be linked with susceptibility to FSGS.

2.4.6 Identification of an ARHGAP24 variation in familial FSGS

Most of the nonsynonymous changes that we detected in patients with FSGS were in the uncharacterized portions of the Arhgap24 protein. However, 1 of the sequence variations (Q158R) is located in the GAP domain of Arhgap24 close to the catalytic arginine (residue 175). This variation is predicted to result in a coding change that would substitute a positively charged arginine for a neutral polar glutamine at position 158 (isoform 1; NCBI accession no. NP_001020787) or position 65 (isoform 2; NCBI accession no. NP_112595). Remarkably, this residue is conserved in the ARHGAP24 gene across several model organisms that have been sequenced (Table 2.3). Of the other 2 nonsynonymous variations for which pedigree DNA was available (T97I and P417A), neither variation reliably correlated with disease status (Figure 2.6).

We first identified the Q158R variation in a Hispanic proband and explored the association of this variation with kidney disease in his family (Figure 2.7). The proband (patient number 1) had elevated serum creatinine levels (16 mg/dl), and a biopsy that was performed at age 20 that
showed FSGS. His sister (patient number 101) also had biopsy-proven FSGS that had progressed to end-stage kidney disease by age 12. The mother (patient number 1001) presented at a late stage and died at age 29 of renal failure. Sequencing showed that the patient’s affected sister and his mother both had the Q158R variation in the ARHGAP24 gene. In contrast, the proband’s 2 other siblings did not possess this variation. The brother (patient number 100) remains healthy, but the sister (patient number 102) had pregnancy-related proteinuria and hypertension. A biopsy was not performed on this individual, and she was lost to follow-up. However, an extended haplotype consisting of 4 flanking microsatellite markers (MSMs) showed that the haplotype of this sister was similar to that of the unaffected brother, making a disease-contributing rearrangement in this region of the genome unlikely (Figure 2.7).

In addition, the exon incorporating this variation was sequenced in an additional 554 control chromosomes (for a total of 914 control chromosomes) to decrease the probability that it represented a low-frequency noncausal variation. Altogether, we had sequenced over 900 control chromosomes across this span and did not detect the Q158R change. We also sequenced members of the affected kindred for mutations in genes known to cause hereditary FSGS and nephrotic syndrome, such as NPHS1, NPHS2, ACTN4, transient receptor potential cation channel, subfamily C, member 6 (TRPC6), phospholipase C, e1 (PLCE1), and WT1 (exons 8 and 9) and inverted formin, FH2 and WH2 domain containing (INF2) (data not shown). There were no suggestive mutations in any of these genes, ruling out known monogenic causes of FSGS in this family. Therefore, in kindred with familial FSGS, the Q158R variation of the ARHGAP24 gene tracked with severe, early-onset kidney disease.
2.4.7 Arhgap24 Q158R has reduced Rac1-GAP activity

Another possibility was that a cosegregating mutation closely linked to the ARHGAP24 gene was responsible for early-onset kidney disease. We therefore tested whether the Q158R variation produced a defective form of the Arhgap24 protein. The Q158R variation is located close to the catalytically active site arginine residue of the GAP domain, and so we evaluated whether this change affected the Rac1-GAP activity of Arhgap24. For these experiments, we used the mouse Q156R protein, which corresponds to Q158R in human Arhgap24. As expected, wild type Arhgap24 reduced the level of active Rac1 in cell lysates, showing again that it has GAP activity against Rac1 (Figure 2.8A). In contrast, transfection of the Q156R-mutated Arhgap24 resulted in a marked increase in the level of active Rac1. Titrating increasing levels of Q156R Arhgap24 against the wild-type protein confirmed that the Q156R mutation impairs the GAP activity of Arhgap24. Next, we tested whether the Q156R variant Arhgap24 could homodimerize and/or heterodimerize with the wild-type protein. FLAG- and GFP-tagged wild-type and Q156R versions of murine Arhgap24 were cotransfected into HEK293 cells. In bidirectional pull-down assays, we detected that both wild type and Q156R Arhgap24 homodimerized and heterodimerized (Figure 2.8B). These experiments showed that the Q158R mutation reduces the enzymatic activity Arhgap24. The fact that Arhgap24 dimerizes may explain the observed dominant effect of this mutation in our family with inherited kidney disease.

2.5 Discussions

Arhgap24, also known as FilGAP and p73RhoGAP, is the protein product of a highly
conserved gene encoded on human chromosome 4. The longer isoform of Arhgap24, which we have used exclusively in this report, is highly expressed in the kidney [94]. GAPs such as Arhgap24 and GEFs for small G proteins comprise a large family of proteins that have diverse patterns of spatial and temporal expression [103]. Such a restricted pattern of expression may allow GAPs and GEFs to behave as cell-type specific effectors for ubiquitously expressed actin regulatory proteins. Here, we show that Arhgap24 is one such candidate for cytoskeletal regulation of the kidney podocyte.

The small G proteins, Rac1, Cdc42, and RhoA, control cell shape and motility through their effects on the actin cytoskeleton. In many systems, RhoA and Rac1 are mutually antagonistic [104, 105]. However, the molecular basis of this counterregulation had been unclear until the discovery of Arhgap24. Stossel and colleagues showed that, in response to RhoA activation, Arhgap24 is phosphorylated by the RhoA effector kinase, ROCK, and this modification stimulates its GAP activity for Rac1 and Cdc42 [94]. By catalyzing the conversion of Rac1 and Cdc42 to their inactive GDP-bound state, Arhgap24 inhibits cell spreading and cell motility. Recently, a similar mechanism was demonstrated for the closely related family member, Arhgap22, in melanoma cells [106]. The diversity of GAPs likely ensures that specific ones may play a role in Rac1-RhoA counterregulation in different cell types and tissues.

Recent work from several groups has shown that the balance of RhoA and Rac1 signaling is carefully regulated in podocytes. Normally, the podocyte appears to be a stationary cell with predominantly active RhoA signaling. The podocyte-enriched adaptor protein synaptopodin controls a signaling module that promotes RhoA activity via 2 different mechanisms.
Synaptopodin blocks Smurf1-mediated ubiquitination and subsequent proteasomal degradation of RhoA [59]. Synaptopodin also disrupts IRSp53 binding to the Cdc42-Mena complex, interrupting Cdc42 signaling that opposes RhoA-mediated stress fiber formation [60]. Synaptopodin itself is phosphorylated and protected from degradation by binding to 14-3-3 [107]. Conversely, when it is dephosphorylated by calcineurin, synaptopodin is degraded by cathepsin L [107]. The net result is that reduced synaptopodin levels lead to decreased RhoA levels and activity [59].

The RhoA-Rac1 signaling balance in podocytes is also responsive to hormonal stimulation. Binding of the vasoactive hormone angiotensin II to its receptor, angiotensin receptor type I (AT1R), leads to a calcium flux through the channels Trpc5 and Trpc6. Recent work by Tian et al. has shown that Trpc5 associates with Rac1 in membrane microdomains, while Trpc6 has a separate association with RhoA [61]. In response to AT1R stimulation, Trpc5 activation leads to Rac1 activation and RhoA inactivation. Triggering Trpc6 has the opposite response, with increased RhoA activity and reduced Rac1 activity. Of note, mutations in the TRPC6 gene have already been associated with human FSGS [17]. Therefore, this recent study by Tian et al. provides new insights into how signaling events downstream of the angiotensin receptor may lead to dynamic cytoskeletal reorganization in podocytes via selective modulation of RhoA and Rac1 activity through calcium signaling.

Decreased RhoA activity and increased Rac1 activity is associated with proteinuric kidney disease, consistent with the idea that an imbalance in the RhoA-Rac1 signaling balance is harmful to the podocyte. HIV infects podocytes and modulates the cytoskeleton of the cell such
that patients with HIV are susceptible to collapsing FSGS [108]. In podocytes, the HIV adaptor protein nef interacts with p190RhoGAP that then inactivates RhoA [109]. In addition, nef binds to diaphanous interacting protein and the Rac1-GEF vav2, resulting in Rac1 activation [110]. Nef also downregulates synaptopodin expression [111] and may further destabilize RhoA signaling via the synaptopodin pathway outlined above. The net effect of HIV infection is a downregulation of RhoA signaling activity and a shift toward Rac1 signaling with consequent podocyte injury.

There is further evidence from mouse models that disrupting the RhoA-Rac1 signaling balance in podocytes can cause kidney disease. Studies on mice deficient for the Rho-guanine nucleotide dissociation inhibitor, RhoGDIα (Arhgdia), demonstrate that Rac1 activation in podocytes is harmful. RhoGDIα binds to and stabilizes Rac1 and RhoA, and, in its absence, levels of these small G proteins and their activity levels are dysregulated [112]. In particular, Rac1 is relatively overactivated, leading to podocyte foot process effacement and proteinuria [56, 113]. Interestingly, treatment with a Rac1 inhibitor reduced proteinuria and the severity of the kidney damage in this model.

Taken together, these studies highlight that the careful control of relative RhoA and Rac1 activity is critical for the proper functioning of podocytes. However, we believe that until this report a direct regulator of Rac1 activity in response to RhoA signaling had not been described in podocytes. Here, we show that Arhgap24 is highly expressed in podocytes and inhibits Rac1-dependent membrane ruffling and epithelial wound healing. Our work predicts that Arhgap24 would enforce the normal RhoA-dependent nonmotile podocyte phenotype by inactivating Rac1. Consistent with a role for Arhgap24 in vivo, we identified a mutation in the
GAP domain of Arhgap24 in kindred with familial FSGS. This mutation reduces the Rac1-GAP activity of Arhgap24. We also show that Arhgap24 dimerizes, and this may explain the dominant pattern of inheritance of the Q158R mutation [114, 115]. Our studies show that Arhgap24 is a potential candidate gene to explain a subset of inherited FSGS. Perhaps, more informatively, Arhgap24 adds to the emerging model that aberrant Rac1 activation is a key step in podocyte dysfunction, leading to proteinuric kidney disease. Since Arhgap24 is selectively expressed in podocytes, modulating its function can alter relative RhoA/Rac1 activity, with potentially minimal systemic side effects, and provides an intriguing pathway for the therapy of proteinuric kidney disease.
Acknowledgements

We are grateful for the support of Washington University George M. O’Brien Center for Kidney Disease Research (P30 DK079333) for some of the patient specimens. We would like to thank Jeffrey Miner for helpful discussions and Jiancheng Hu for technical assistance and reagents for the GTPase activity pull-down assays. This work was supported in part by the NIDDK Intramural (to J.B. Kopp) and Extramural Research Programs (RO1DK058366, to A.S. Shaw), Howard Hughes Medical Institute, and the Medical Scientist Training Program Grant to Washington University School of Medicine.

* This chapter is adapted from Akilesh S., et al. Arhgap24 (FilGAP) inhibits Rac1 activity in mouse podocytes and is associated with familial focal segmental glomerulosclerosis. J Clin Invest. 2011; 121(10): 4127-37.
2.6 Tables and Figures

Table 2.1 Incidence of ARHGAP24 nonsynonymous sequence variations in patients with biopsy-proven FSGS (n=310) and controls (n=180)

<table>
<thead>
<tr>
<th>Variation</th>
<th>No. affected</th>
<th>No. controls</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Isoform 1</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>T97I</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>R142C</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Q158R</td>
<td>2</td>
<td>0</td>
</tr>
<tr>
<td>L215V</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Q359R</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>S396L</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>P417A</td>
<td>9</td>
<td>2</td>
</tr>
<tr>
<td>T451I</td>
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<td>0</td>
</tr>
<tr>
<td>T481M</td>
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</tr>
<tr>
<td>F539L</td>
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<td>5</td>
</tr>
<tr>
<td>N587I</td>
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<tr>
<td><strong>Isoform 2</strong></td>
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</tr>
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<td>1</td>
</tr>
<tr>
<td>R5L</td>
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<td>0</td>
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The isoform 1 protein reference sequence is NCBI accession no. NP_001020787, and the isoform 2 reference sequence is NCBI accession no. NP_112595.
Table 2.2 Non-synonymous SNPs in ARHGAP24

Non-synonymous SNPs identified in the transcript of isoform 1 of ARHGAP24 from over 1000 human genomes sequenced. (www.1000genomes.org) SNPs in bold are known polymorphisms in the gene.

<table>
<thead>
<tr>
<th>Residue</th>
<th>Variation</th>
<th>SNP ID</th>
<th>Alleles</th>
</tr>
</thead>
<tbody>
<tr>
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<td>rs36067390</td>
<td>G/A</td>
</tr>
<tr>
<td>Q359R</td>
<td>YRI_P3:4:87134934:G</td>
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<td>D368G</td>
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<td>A/G</td>
<td></td>
</tr>
<tr>
<td>P417A</td>
<td>rs35521695</td>
<td>C/G</td>
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</tr>
<tr>
<td>V420T</td>
<td>LWK_P3:4:87135089:A</td>
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<td></td>
</tr>
<tr>
<td>K421Q</td>
<td>YRI_P3:4:87135092:C</td>
<td>A/C</td>
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<tr>
<td>G473V</td>
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</tr>
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</tr>
<tr>
<td>E721*</td>
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<td>G/T</td>
<td></td>
</tr>
</tbody>
</table>

Table 2.3 Q158 is a conserved residue

Sequence alignment of the Arhgap24 protein across various species in the region of the patient variation (Q158) shows complete conservation of the glutamine residue.

<table>
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<tr>
<th>Species</th>
<th>aa</th>
<th>Sequence</th>
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<th>Accession no.</th>
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<td>Pan troglodytes</td>
<td>330</td>
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</tr>
<tr>
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<td>168</td>
<td>NP_083546</td>
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<tr>
<td>Rattus norvegicus</td>
<td>140</td>
<td>vryekgrnr lapmlveQcv dfirqrqlke</td>
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<tr>
<td>Callithrix jaccus</td>
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<tr>
<td>Equus caballus</td>
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<td>XP_001495166</td>
</tr>
<tr>
<td>Bos taurus</td>
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<td>Canis familiaris</td>
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<td>Gallus gallus</td>
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<td>vryekgrnr lapmlveQcv dfirqrqlke</td>
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<tr>
<td>Monodelphis domestica</td>
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<td>Danio rerio</td>
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<td>vryergrnk mapmlveQcv dfirwgglre</td>
<td>173</td>
<td>XP_0013399010</td>
</tr>
</tbody>
</table>

The location of the region of interest in each protein is designated by the amino acid start and finish positions. The glutamine residue is shown in bold. NCBI accession numbers are shown.
Figure 2.1 Differentiated podocytes show reduced membrane ruffling

(A) YFP-actin–transduced podocytes were cultured at the permissive temperature (33°C) or differentiated for 7 to 14 days at the nonpermissive temperature (37°C). Kymographs (insets) obtained at the sites of active ruffling (hatched boxes) show prominent ruffling at 33°C that is reduced at 37°C (also see Supplemental Videos 1 and 2). DIC, differential interference contrast. Original magnification, ×400; approximately ×1,000 (insets and time-lapse panels). (B) Quantification of actin spike lengths in kymographs shows that differentiated podocytes (n = 11) have a significant decrease in ruffling activity compared with that of undifferentiated podocytes (n = 15). Individual symbols represent data from individual podocytes. P = 3.29 × 10^{-6} by ANOVA with post-test correction.
Figure 2.2 Arhgap24 transcript and Arhgap24 protein are specifically expressed in podocytes

(A) MBEI values for microarray data for in vitro–cultured podocytes (33°C, 37°C), ex vivo–isolated podocytes (E13.5, E15.5, and adult), and laser-capture-microdissected glomeruli (E12.5 renal vesicle, E15.5 S-shaped body, and E15.5 renal corpuscle) show an increase in the Arhgap24 transcript with differentiation in vitro and in vivo. (B) By quantitative RT-PCR, differentiated podocytes have higher Arhgap24 mRNA levels compared with those of undifferentiated podocytes, after normalization to 18S rRNA. (C) Immunoblotting shows that differentiated podocytes have higher levels of Arhgap24 protein than undifferentiated podocytes. (D) Confocal imaging of cultured podocytes also shows an increase in levels of Arhgap24 in differentiated podocytes, concentrated in punctate structures at the base of the cell. (E) Arhgap24 colocalizes with the focal adhesion marker vinculin at the tips of actin stress fibers. Original magnification, ×600 (D and E).
Figure 2.3 Arhgap24 is expressed in kidney podocytes *in vivo*

(A) Immunoblotting of tissue lysates shows that Arhgap24 is expressed in the kidney (expected size ~95 kD). Smaller bands likely represent specific degradation products. The positive control (+) is a lysate from HEK293 cells transfected with FLAG-tagged Arhgap24. The negative control (−) is HEK293 cell lysate. Brn, brain; Kid, kidney; Liv, liver; Lng, lung; Spl, spleen. (B) Magnetic separation of glomeruli after beating heart perfusion of mice with magnetic beads results in more than 95% pure isolated glomeruli (top). The flow through consists of tubule fragments (bottom). Original magnification, ×100. (C) Arhgap24 is enriched in the glomerular (Glom) fraction that also contains the podocyte protein, podocin. Tub, tubule. (D) A low-magnification view (original magnification, ×200) of a mouse kidney stained for Arhgap24 shows that the highest signal is detected in the glomeruli. (E) Within mouse glomeruli, the Arhgap24 signal colocalizes with that of the podocyte marker synaptopodin. Original magnification, ×600.
Figure 2.4 Arhgap24 knockdown in differentiated podocytes increases membrane ruffling

(A) Two lentivirally transduced cell lines targeting different portions of the Arhgap24 transcript show marked reduction of Arhgap24 protein compared with that of an irrelevant knockdown (Fluc, 100%; line 451, 15%; line 756, 40%). Results are representative of at least 3 independent experiments. (B) Representative images of the 3 knockdown cell lines after differentiation at 37°C for 7 to 10 days. Kymographs (insets) generated from the hatched box area and time-lapse sequences show that the 2 knockdown cell lines have increased membrane ruffling compared with that of the control (also see Supplemental Videos 3–5). Original magnification, ×400; approximately ×1,000 (insets and time-lapse panels). (C) Quantitation of actin spikes in kymographs shows that both the 451 (n = 19) and 756 (n = 11) Arhgap24 cell lines have significantly greater membrane ruffling activity compared with that of the control knockdown Fluc cell line (n = 18) (*P < 0.00001). The ruffling activity is not significantly different between the 2 knockdown cell lines (P = 0.57). Group differences were analyzed by ANOVA with post-test correction.
Figure 2.5 Arhgap24 knockdown in differentiated podocytes increases active Rac1 and Cdc42 levels and accelerates epithelial monolayer wound closure

(A) Pull down of active (GTP-bound) Rac1, Cdc42, and RhoA shows that the Arhgap24 knockdown cell lines (lines 451 and 756) have increased levels of active Rac1 and Cdc42 compared with those of the control (Fluc). However, active RhoA levels are similar across all 3 cell lines. Results are representative of 3 independent experiments. (B) Arhgap24 knockdown cells migrate and close a scratch made in a confluent monolayer faster than the control knockdown cell line. Cell nuclei were stained with Hoechst dye. Original magnification ×100.
Figure 2.6 Pedigree information for T97I and P417A variations

DNAs from one family for the T97I variation and three families for the P417A variation were available for sequence analysis. The presence of kidney disease (clinical and/or biopsy proven) is denoted by the filled-in symbols. If DNA was analyzed for sequence variation, the result of a non-synonymous variation is denoted within the individual symbol (Wt = wildtype; Ht = heterozygous). The reference protein sequence (NP_001020787) was considered wildtype. In no instance did the variation consistently correlate with kidney disease.
Individuals with end-stage kidney disease are denoted by black symbols. Deceased family members are represented by diagonal lines. Flanking MSM analysis shows that the unaffected siblings have a distinct haplotype that is different from that of the affected individuals. The columns of numbers and letters under each symbol refer to the alleles that individual carries at the given genetic markers. Where these numbers are within brackets, the haplotype is inferred. The solid black rectangle beneath individuals 1, 101, and 1001 represents the inherited disease haplotype.
Figure 2.8 Arhgap24 Q158R has defective Rac1-GAP activity and dimerizes with the wild-type protein

(A) Wild-type FLAG-tagged Arhgap24 (first lane) transfected into HEK293 cells reduces active Rac1 levels compared with Q156R Arhgap24 (last lane). Titration of increasing proportions of Q156R Arhgap24 produces increased levels of active Rac1 (middle lanes). Total Rac1 and FLAG-Arhgap24 protein levels are similar across all lanes. Results are representative of 3 different experiments. (B) FLAG- or GFP-tagged wild-type (W) or Q158R Arhgap24 (Q) constructs were cotransfected into HEK293 cells. Cell lysates were immunoprecipitated with anti-FLAG antibody and then immunoblotted for GFP-tagged Arhgap24 to assess for protein dimerization. Whole cell lysates (WCLs) were immunoblotted for GFP and FLAG to ensure protein expression.
Chapter 3. Rac1 Activation in Podocytes Induces Rapid Foot Process Effacement and Proteinuria

3.1 Abstract

The kidney’s vital filtration function depends on the structural integrity of the glomerulus, the proximal portion of the nephron. Within the glomerulus, the architecturally complex podocyte forms the final cellular barrier to filtration. Injury to the podocyte results in a morphologic change called foot process effacement and this is a ubiquitous feature of proteinuria. The exact nature of foot process effacement is not known but recently it has been proposed that this might reflect activation of the Rac1 GTPase. To test this hypothesis, we generated a podocyte specific, inducible transgenic mouse line that expressed constitutively active Rac1. We observed a rapid onset of proteinuria that began to remit spontaneously in one week. Using super resolution imaging, we verified that the induced transgene was expressed in damaged podocytes. The kinetics of this podocyte injury model differs from previously reported studies and highlights the complex balance of Rho-GTPase signaling that is required for proper regulation of the podocyte cytoskeleton.

3.2 Introduction

The structural integrity of the proximal portion of the nephron, the glomerulus, is vital to the kidney’s filtration function. Within the glomerular capillary tuft, the kidney’s filtration barrier is a biomechanical composite of fenestrated endothelial cells, a thick glomerular basement
membrane and complex visceral epithelial cells called podocytes. Podocytes lie on the outer aspect of glomerular capillaries and extend cytoplasmic processes (foot processes) that interdigitate with those from neighboring podocytes to form a mesh-like network that forms the final barrier to filtration. Podocyte foot processes are built around highly organized actin bundles that are reorganized during injury with flattening and simplification (“effacement”) of foot processes leading to spillage of serum proteins into the urine (proteinuria). Defects in actin regulatory proteins lead to irreversible podocyte injury and focal and segmental glomerulosclerosis (FSGS) in humans and in animal models [2].

Recent studies show that rather than being a static filter, the cytoskeleton and therefore the shape of the podocyte are both tightly regulated and dynamic [52]. Small GTPases of the Rho family (exemplified by RhoA, Cdc42, Rac1) are the central organizers of the actin cytoskeleton [48]. After receiving diverse signaling inputs, the Rho-family of small GTPases act through their effectors to polymerize and organize actin filaments into various configurations that deform the cell membrane and change cell shape. Of the three major Rho-family GTPases, Cdc42 has been shown to be critical for podocyte development, while both RhoA and Rac1 seem dispensable in early stages [28].

After this initial phase, RhoA and Rac1 seem to play more important roles in podocyte cell biology. In many biological systems, including podocytes, RhoA and Rac1 antagonize each other’s activation and function [50, 116]. Recent studies have shown that constitutive activation of RhoA causes podocyte foot process effacement and proteinuria after several weeks [49, 117], suggesting that inappropriate RhoA activation is pathogenic to podocytes. This is surprising
given that 1) activation of Rho family GTPases causes rapid cytoskeletal rearrangement \textit{in vitro} [118, 119]; and 2) introduction of dominant-negative RhoA produces a similar phenotype [49].

On the other hand, it has been proposed that excessive Rac1 activation or inhibition of Rho activity might be the key step in podocyte injury. Synaptopodin, a podocyte actin-binding protein, reinforces RhoA signaling and suppresses Cdc42 signaling to promote proper cytoskeletal architecture [59, 120]. Genetic ablation of synaptopodin in mice results in increased susceptibility to proteinuria [60, 121]. Deletion of RhoGDI (a negative regulator of Rho-family GTPases) in mice results in foot process effacement and proteinuria that correlates with increased Rac1 activity [56]. Mutations in the GTPase activating protein Arhgap24 result in increased Rac1 activation \textit{in vitro} and are correlated with podocyte injury and FSGS in patients [122].

The effects of Rac1 activation in podocytes have only been studied indirectly through the manipulation of upstream regulatory proteins [21, 56, 122, 123]. To test directly whether Rac activation induces podocyte foot process effacement, we generated a double transgenic system in mice in which GFP-tagged constitutively active Rac1 (Rac1Q61L) is expressed in podocytes after doxycycline (DOX) induction. We observed rapid onset of proteinuria within 2 days of DOX induction. The degree of proteinuria correlated with the levels of active Rac1 expression. However, proteinuria in this system was not durable and gradually decreased over the course of a month despite continuous exposure to DOX. Thus, activation of Rac1 in podocytes rapidly causes foot process effacement and proteinuria \textit{in vivo}. These results are distinct from the effects of RhoA activation in podocytes and emphasize the complex interplay of small GTPase signaling in the regulation of podocyte shape and function.
3.3 Methods

3.3.1 Generation of EGFP_CA-Rac1 knock-in transgenic mice

We chose the X-linked \( Hprt1 \) locus for targeting because it is a nonessential housekeeping gene that encodes a selectable marker [124, 125]. The pHPT targeting vector was generated on a pBluescript SKII(-) backbone by PCR amplifying the left arm (5.1 kb fragment upstream of Exon 1) and right arm (2.1 kb fragment downstream of Exon 1) of the \( Hprt1 \) gene from a bacterial artificial chromosome (RP24-335G16). The tetracycline responsive promoter element (TRE), EGFP, Rac1Q61L, and bovine growth hormone polyadenylation signal (bGH-polyA) sequences were amplified by PCR, and inserted sequentially into the pHPT targeting vector. The KH2 ES cell line harboring the Rosa26-M2rtTA insertion was used for transfection. Cells with homologous recombination of the transgene into the \( Hprt1 \) locus were selected based on their growth in the presence of 6-thioguanine, which is toxic to cells expressing functional \( Hprt1 \). Appropriate single copy insertion of the EGFP_CA-Rac1 transgene into the \( Hprt1 \) locus was confirmed by PCR. Targeted ES cells were injected into blastocysts to generate chimeric mice. EGFP genotyping primers (a protocol from The Jackson Laboratory) were used for genotyping (Forward_oIMR0872 5’-AAGTTCCATCTGCACCACCG-3’, Reverse_oIMR1416 5’-TCCTTGAAGAAGATGGTGC-3’; internal positive control Forward_oIMR7338 5’-CTAGGCCACAGAATTGAAAGATCT-3’, Reverse_oIMR7339 5’-GTAGGTGGAATTCTAGCATCATCC-3’). All animal experiments were conducted with approval of the Washington University Animal Studies Committee.
3.3.2 Mouse strains and transgene induction

The *Rosa26*-rtTA mouse strain was purchased from The Jackson Laboratory (#006965). The *NPHS2*-rtTA strain was obtained from Dr. Jeffrey Kopp at the NIH. The *Nphs1*-rtTA-3G strain will be described in detail elsewhere [126]. All mice used in this study were male and therefore carried a single copy of the EGFP_CA-Rac1 transgene. To induce transgene expression, regular chow was substituted with DOX-supplemented chow (2000ppm, TestDiet) for the indicated time periods.

3.3.3 Cell culture and cell based assays

Immortalized murine podocytes were maintained and differentiated as described previously [122]. For live cell imaging assays, podocytes were infected with lentiviral vectors encoding N-terminal EGFP tagged CA-Rac1 and CA-RhoA. An EGFP empty vector was used as control.

3.3.4 Antibodies

Antibodies for immunostaining included rabbit anti-podocin (Sigma Aldrich P0372, 1:400 dilution), goat anti-nephrin (R&D Systems AF3159, 1:100 dilution), rabbit anti-laminin β2 (20) (1:1500 dilution) and chicken anti-GFP (Invitrogen A10262, 1:500 dilution). Antibodies used for immunoblotting were mouse anti-GFP (Clontech 632381, 1:10000 dilution), rabbit anti-ERK2 (Santa Cruz Biotechnology sc-154, 1:5000 dilution) and rabbit anti-podocin (Sigma-Aldrich P0372, 1:500 dilution). Fluorescently-conjugated secondary antibodies were purchased from Jackson Immunoresearch, and the STORM antibodies were conjugated as described [127].

3.3.5 Immunofluorescence assays
Fresh kidney tissue was embedded in OCT compound and snap frozen on dry ice. 8 µm cryosections were applied to charged slides. Cultured podocytes were seeded onto collagen I coated coverslips. For immunofluorescence assays, the tissue sections or coverslips containing podocytes were fixed with 1% PFA in PBS for 5 minutes followed by blocking and permeabilization with 2% FBS in PBS with 0.1% saponin. Primary antibodies at the indicated dilutions were applied for 1 hour at room temperature. After extensive washes, fluorescently conjugated secondary antibodies were applied at 1:500 dilution for another hour at room temperature. After washing, the prepared slides were imaged on a Olympus FV-1000 spinning disc confocal microscope.

3.3.6 Albumin/creatinine assay

Mouse urine samples were collected at the indicated time points and urinary albumin (Bethyl, E90-134) and creatinine (BioAssay Systems, DICT-500) were quantified by ELISA according to the manufacturers’ protocols.

3.3.7 Transmission Electron Microscopy

Portions of kidney cortex were fixed with 2% paraformaldehyde and 2% glutaraldehyde and processed for electron microscopy. Ultrathin sections were prepared and imaged by the Electron Microscopy Core Facility at Washington University.

3.3.8 STORM imaging and STORM-SEM correlation

Kidney tissues were fixed in 4% paraformaldehyde and cryoprotected in 30% sucrose. Tissues were then embedded in OCT compound and semi-thin sectioned at 100-200 nm thickness using an ultra cryomicrotome. These tissue sections were collected on a
carbon-coated #1 coverslip and fixed and stained as described previously [127]. Using a custom STORM microscope, 20,000 imaging cycles were collected and the resulting images were compiled to generate a composite multi-channel image as described earlier [128]. After STORM imaging, the coverslip was floated off and the sample was fixed with 2% glutaraldehyde. This sample was then processed for scanning electron microscopy (SEM). The images obtained from STORM and SEM were superimposed using Photoshop CS5.1.

3.3.9 Live cell imaging and kymograph analysis

Undifferentiated murine podocytes were transiently transfected using Amaxa Nucleofection (Lonza, Allendale, NJ) with plasmids encoding for constitutively active Rac1 or RhoA. Transfected podocytes were cultured on collagen I-coated glass-bottomed dishes overnight, and serum starved for 6 hours to arrest baseline membrane ruffling. Rac1 transfected podocytes were imaged in the serum starved state, while Rho transfected podocytes were imaged following the addition of 10% FBS 10 minutes prior to imaging to induce membrane ruffling. Sequential images were obtained by an Olympus FluoView FV1000 microscope every 10 seconds for a 20-minute duration, and movies assembled using Olympus Fluoview software. The ImageJ plug-in, Multiple Kymograph (http://rsbweb.nih.gov/ij), was used to generate kymographs at 5 different locations of maximum membrane ruffling for each imaged cell [100]. 10 actin spikes were measured for each kymograph, and average length (ruffling index) was determined as previously described [100].

3.3.10 Rac1 pull-down assay

The GST-tagged Pak1-PBD was expressed in BL21(DE3) E. coli and purified using
glutathione-agarose beads. For the active Rac1 pull-down assay, whole kidney lysates were generated by homogenizing the kidneys in cell lysis buffer (50mM Tris-HCl, pH7.5, 150mM NaCl, 5mM MgCl₂, 10% Glycerol, 1% NP-40, 1mM DTT, 1mM PMSF, 10µg aprotinin, 10µg leupeptin; aprotinin, leupeptin, DTT and PMSF are fresh added), and isolating the supernatant. Equal volumes of the lysates were incubated with GST-PBD beads. Rac1-GTP bound to the beads (active Rac1) was eluted with Laemmli sample buffer and examined by western blot.

3.4 Results

3.4.1 Rho family GTPases induce distinct effects on the actin cytoskeleton in cultured murine podocytes

A network of highly organized actin cytoskeleton structures forms the structure of podocyte foot processes. During podocyte foot process effacement, these actin bundles are reorganized into broad membrane sheets that resemble lamellipodia seen in cultured cells. Numerous cell culture systems point to a critical role for Rho family GTPases in actin cytoskeleton remodeling with RhoA activation inducing actin bundling and Rac1 activation inducing lamellipodia [48]. Therefore, we hypothesized that the remodeling of the podocyte actin cytoskeleton seen during the process of effacement may represent an alteration in the balance between Rac1 and RhoA activities [122].

To test the effects of Rho GTPase activation on the podocyte actin cytoskeleton in vitro, we first transfected a murine immortalized podocyte cell line with enhanced green fluorescent protein (EGFP) fused to constitutively active versions of Rac1 (EGFP_CA-Rac1) and RhoA
(EGFP_CA-RhoA). Actin fibers and focal adhesions were visualized with phalloidin and vinculin respectively. Constitutively active Rho (CA-Rho) increased the number of stress fibers and focal adhesions in podocytes (Figure 3.1A). In contrast, CA-Rac1 expression in podocytes induced membrane spreading and lamellipodia formation resulting in large, round and flattened cells (Figure 3.1A).

Since foot process effacement might be an expression of podocyte motility [93], we next asked if these changes in cell morphology correlated with changes in podocyte membrane dynamics. Using live cell imaging, we quantitated membrane motility using kymograph analysis as described previously [122]. Podocytes expressing CA-Rac1 show increased membrane ruffling compared with wild type cells after serum starvation (Figure 3.1B, and Videos S1 and S2). Addition of serum induced membrane ruffling in wild type cells, and this was largely suppressed in podocytes expressing CA-RhoA (Figure 3.1C, and Videos S3 and S4). These experiments confirmed that activation of Rac1 and RhoA produce marked changes in the podocyte actin cytoskeleton and on membrane dynamics. Active Rac1 induced lamellipodia formation and increased membrane motility in podocytes, while active RhoA stabilized the cytoskeleton and suppressed membrane motility.

3.4.2 Generation of inducible EGFP_CA-Rac1 transgenic mice

Our in vitro results suggested that Rac1 activation might have significant effects on podocyte morphology. This is in contrast to recent studies that showed RhoA activation could play a pathogenic role in podocytes in vivo [49, 117]. To test the effects of Rac1 activation in podocytes in vivo, we generated a transgenic mouse model that would allow for inducible
expression of EGFP_CA-Rac1. Using homologous recombination, we targeted the EGFP_CA-Rac1 transgene into the Hprt1 locus in ES cells [124, 125] (Figure 3.2A) containing the tetracycline inducible transactivator (rtTA) inserted into the Rosa26 locus (Rosa-rtTA) [129]. Cells with successful homologous recombination of the transgene for the Hprt1 locus were selected based on their growth in the presence of 6-thioguanine, which is toxic to cells harboring a functional Hprt1 allele. Targeted gene insertion was confirmed by PCR (Figs. 2 B and C) and recombinant ES cells were microinjected into blastocysts to generate chimeric mice.

To test inducible expression of the EGFP_CA-Rac1 transgene, mice were fed doxycycline (DOX) for one week, and multiple tissues were harvested and examined for EGFP expression by fluorescence microscopy (Figure 3-S1). DOX induced strong transgene expression in multiple tissues. However, we could not detect EGFP expression in podocytes by fluorescence microscopy. These experiments revealed that the Rosa26-rtTA allele was ineffective at driving efficient transgene expression in podocytes.

3.4.3 EGFP_CA-Rac1 expression in podocytes causes rapid onset but transient proteinuria

To induce podocyte-specific expression of the CA-Rac1 transgene, we crossed our mice with transgenic mice with the rtTA driven by the human podocin promoter (NPHS2-rtTA) [130]. In double transgenic mice (PODxRac1) fed DOX, EGFP expression was detectable as early as 4 days after induction. To validate the functionality of the CA-Rac1 transgene product, we used the p21 binding domain of PAK1 (PBD) to precipitate CA-Rac1 from whole kidney lysates of DOX-treated double transgenic mice (Figure 3.2D). EGFP_CA-Rac1 protein was easily detectable in the DOX-treated mice but was undetectable in the absence of DOX. Expression of
the transgene was barely detectable in the whole kidney lysate, which was not surprising given that podocytes constitute only a small fraction of cells in the kidney. Expression of the CA-Rac1 transgene in podocytes was confirmed by colocalization with the podocyte-specific marker (podocin) by immunofluorescence microscopy (Figure 3.3A). Surprisingly, the CA-Rac1 transgene was expressed in only a small fraction of podocytes, and this expression was variable between glomeruli and between mice.

After 2 days of induction with DOX, however, PODxRac1 mice developed significant proteinuria compared to single transgenic control mice. Proteinuria reached its peak on day 4 and then began to abate around 1 week post-induction (Figure 3.3B, and 3-S2). DOX treatment for up to one month did not result in progressive renal dysfunction or significant histologic alterations. Given the variable expression of the transgene, we assessed whether the level of proteinuria correlated with the level of expression (Figure 3.3C). The magnitude of proteinuria positively correlated with the frequency of EGFP-positive glomeruli. The kinetics of proteinuria, however, were similar among all PODxRac1 mice beginning around day 2 and abating after day 7 (Figure 3-S2). These experiments showed that podocyte-specific expression of CA-Rac1 induces rapid and transient proteinuria that correlated with the level of transgene expression.

3.4.4 Nphs1-rtTA-driven EGFP_CA-Rac1 expression in podocytes results in higher transgene expression and more severe proteinuria

We considered that the patchy and uneven expression of CA-Rac1 might be due to issues with silencing of the NPHS2-rtTA transgene. To circumvent this problem, a new transgenic mouse line (NEF) expressing a modified rtTA (rtTA-3G) under the control of the mouse nephrin
(Nphs1) promoter was generated [126] and bred to our CA-Rac1 mice. Examination of kidneys from double transgenic (NEFxRac1) mice after four days of DOX treatment showed a greater proportion of glomeruli and higher numbers of podocytes expressing the transgene compared to the PODxRac1 mice. Yet, the expression in podocytes was still not 100% (Figure 3.4A). This increased expression and distribution of the CA-Rac1 transgene correlated with a faster onset and higher levels of proteinuria (Figure 3.4B). In contrast to PODxRac1 mice, the proteinuria in NEFxRac1 mice persisted after one week of DOX treatment (Figure 3.4C, 3-S2). However, similar to PODxRac1 mice, proteinuria peaked at day 4 and gradually decreased over time (Figure 3.4C). After exposure to DOX for 28 days, we could not detect any EGFP expressing podocytes in kidney sections suggesting that the transgene-positive podocytes had been lost (Figure 3.5B).

3.4.5 EGFP_CA-Rac1 induces foot process effacement, but without other histological changes in the glomerulus

The morphology of podocytes expressing CA-Rac1 was assessed by both light and electron microscopy. Glomeruli from DOX-induced Rac1 single transgenic mice and NEFxRac1 mice were unremarkable by light microscopy (Figure 3.5A). No obvious abnormalities were detected even after 1 month of continuous DOX induction. Transmission electron microscopy showed segmental effacement of podocyte foot processes in PODxRac1 glomeruli (Figure 3.6A) consistent with variable transgene expression. To test this, we used a super resolution fluorescence imaging method, Stochastic Optical Reconstruction Microscopy (STORM). STORM allows nanometer resolution of fluorescently tagged molecules such as labeled antibodies by
capturing thousands of sequential images of a specimen illuminated with a very low energy excitation source [127]. Sections of kidney tissue from NEFxRac1 mice on day 4 after DOX induction was stained with fluorescently tagged antibodies for EGFP (to localize CA-Rac1) and laminin β2 (to detect the glomerular basement membrane) and examined by STORM. After STORM imaging, the same tissue sample was processed for freeze etch electron microscopy and the images were correlated (Figure 3.6 B and C). This showed that only the EGFP_CA-Rac1 expressing podocytes had effaced foot processes, while adjacent non-expressing podocytes had intact foot processes (Figure 3.6C). These studies provide an explanation for the segmental foot process effacement seen by TEM.

3.4.6 CA-Rac1 decreases podocin and nephrin levels via proteasomal degradation

During our examination of the kidney with immunofluorescence microscopy, we noted that in both PODxRac1 and NEFxRac1 mice CA-Rac1 expression was inversely correlated with podocin and nephrin level (Figure 3.7 A and B). Quantitative correlation of EGFP pixel intensity with that of podocin and nephrin showed a significant negative correlation (Figure 3.7 C, D, E and F). To test whether Rac1 activation cause a direct downregulation of podocin expression, cultured podocytes were transiently transfected with empty vector, EGFP_CA-Rac1 and EGFP_CA-RhoA, and podocin levels were analyzed by immunoblotting. Compared with the vector control, CA-Rac1 diminished podocin levels by about 50%, while CA-RhoA had no effect (Figure 3.7G). We measured the levels of podocin mRNA in these podocytes by quantitative PCR and found no significant difference, which suggested that the podocin decrease was not mediated by transcriptional downregulation (Figure 3-S3). The decrease in podocin levels could
be blocked by the proteasome inhibitor MG132, suggesting that Rac1 activation induces podocin degradation via a proteasomal pathway (Figure 3.7H).

3.4.7 CA-Rac1 expression stimulate membrane dynamics in podocytes in vivo

To further investigate that CA-Rac1 expressing podocytes in vivo, we used multiphoton intravital imaging (MPM) technique to directly assess podocyte membrane dynamics in live mice. Since CA-Rac1 is tagged with EGFP at the N-terminus and only some of the podocytes expressed this transgene after DOX induction, we could directly observe EGFP+ podocyte membrane dynamics. Dylight594-labeled lectin molecules were injected intravenously before imaging. Endothelial cells and Tubular epithelial cells can absorb lectin and be labeled in the red channel. During the imaging process, each frame was taken at 30-second time interval by using resonance scanner. We observed dramatic membrane dynamics in EGFP_CA-Rac1+ podocytes (Figure 3.8). Our result suggests that CA-Rac1 induce membrane ruffling in podocytes in vivo. The membrane ruffling is likely caused by actin cytoskeleton rearrangement induced by high Rac activity.

3.4.8 The shattered podocytes could form new interaction with the interstitial endothelium in the renal tubules

In our intravital imaging experiments, we observed EGFP+ cells that formed stable interaction with the renal tubules (Figure 3.9). Because the expression of rtTA transgene is restricted to podocytes [126], these cells could be podocytes that shattered from the glomerulus. Surprisingly, 3D images of these cells showed that they establish stable interactions with the epithelial cells in the tubules (Figure 3.9). Some of these cells extended protrusions that crossed
the basement membrane of the tubular epithelium and formed new interactions with the endothelial cells of the interstitial capillaries. This result suggests that high Rac activity could cause podocyte shattering from the glomerulus. However, shattered podocytes could form new connections with other endothelial cells outside the glomerulus.

3.5 Discussions

The structural integrity of the glomerular filtration barrier is essential for selective excretion of waste products and the retention of cells and large serum proteins within the circulation. The glomerular podocyte, with its arbor of interdigitating foot processes, is the critical final component of the kidney’s filtration barrier. The elaborate actin-based cytoskeleton of the podocyte’s foot process is simplified in many diseases in which serum proteins are lost into the urine. Since actin reorganization mediated by Rho family GTPases is a well-established mechanism for cell shape change, we and others have attempted to directly ask how specific Rho GTPase activation regulates podocyte morphology in vivo.

Previous studies used an approach similar to ours to study the role of RhoA in podocytes. Both constitutively active and dominant negative (DN) inducible RhoA transgenic mouse lines were generated [49, 118]. While both approaches resulted in podocyte dysfunction, the changes detected occurred relatively slowly, over weeks to months. Given how rapidly Rho family GTPases induce actin cytoskeletal changes in vitro, it raises the possibility that proteinuria in those systems is due to indirect effects of transgene expression in podocytes and not due to a direct signaling effect of RhoA. In addition, the mechanism of foot process effacement directed
by RhoA activation was not directly addressed in these studies.

In our mouse, expression of constitutively active Rac1 (CA-Rac1) produced rapid onset (~48 hours) proteinuria that correlated with the degree of transgene expression. While the \(Nphs1\)-rtTA transgene was expressed in a larger percentage of podocytes than the \(NPHS2\)-rtTA transgene, neither driver promoted expression in all of the cells. This was supported by our finding of segmental foot process effacement detected by electron microscopy. Using STORM imaging and freeze-etch electron microscopy correlation techniques, we confirmed that effaced foot processes correlated with GFP positive cells, while the foot processes of podocytes not expressing the transgene remained intact [128]. The rapid onset and dose-responsive nature of proteinuria induced by CA-Rac1 expression provides strong evidence for a direct signaling role for Rac1 activation in generating foot process effacement. This in vivo biologic correlate further supports the use of the membrane-ruffling assay in cultured podocytes as a useful reporter of podocyte injury in vitro. Our results demonstrate for the first time that Rac1 activation in podocytes can directly cause foot process effacement and proteinuria.

While onset of proteinuria in the CA-Rac1 expressing mice was rapid, the proteinuria induced was only transient. As GFP could not be detected in kidneys of mice treated long-term with DOX, we suspect that chronic Rac1 activation is toxic to podocytes. The efficiency of CA-Rac1 transgene expression differed between the two different rtTA driver lines and therefore incomplete CA-Rac1 expression is related in part to the efficiency of the promoter driving rtTA expression. It is well-known that epigenetic silencing can suppress transgene expression over time [131]. It is also possible that the incomplete expression of our CA-Rac1 transgene may be
related to other epigenetic factors that can affect expression of both the driver and CA-Rac1 transgenes.

We did not observe any progressive renal dysfunction or focal and segmental glomerulosclerosis (FSGS) in the CA-Rac1 expressing mice after prolonged DOX treatment, in contrast to the CA-RhoA transgenic mice [118]. This may be attributed to the low frequency of EGFP_CA-Rac1-expressing podocytes in our systems, since in a rat model of tunable podocyte loss, approximately 20 to 40% podocyte depletion was required before FSGS was consistently observed [26].

The presence of CA-Rac1 reduced podocin protein expression both in vivo and in cultured podocytes. This was mediated, at least in part, by proteasomal degradation as it could be blocked in vitro with proteasome inhibitors. Because we could not detect CA-Rac1-positive cells after extended DOX treatment, we suspect that the expressing podocytes were eliminated and replaced either by remaining podocytes or by a progenitor population [126, 132]. These compensatory mechanisms might explain the resolution of proteinuria over time in the CA-Rac1 expressing mice. Therefore, future studies should focus on the development of tools to detect activation of endogenous Rac1 and RhoA in podocytes. The balance and localization of the activity of these GTPases is likely to be tightly and dynamically regulated for proper podocyte function.
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3.6 Figures

Figure 3.1 Constitutively active Rho family GTPases exert opposing effects on the actin cytoskeleton of podocytes

(A) Immunofluorescence imaging was performed on differentiated mouse podocytes that were stably transduced with EGFP_CA-Rac1 or EGFP_CA-RhoA (green). The filamentous actin network was delineated with phalloidin (red) and focal adhesions and contacts were identified with vinculin (blue). Compared to control, untransduced cells, EGFP_CA-Rac1 induced flattening of the cell and lamellipodia formation while EGFP_CA-RhoA induced numerous stress fibers and cell contraction. (B) EGFP_CA-RhoA suppresses serum-induced membrane ruffling in podocytes. Left panel, using kymographic analysis, under conditions of serum starvation, podocytes exhibit minimal membrane ruffling activity. Introduction of EGFP_CA-Rac1 significantly increases membrane dynamics and ruffling (*** P < 0.0001 by unpaired T-test). Right panel, exposing starved cells to serum also induces membrane ruffling which is suppressed by introduction of EGFP_CA-RhoA (*** P < 0.0001 by unpaired T-test).
**Figure 3.2 Generation of inducible EGFP_CA-Rac1 transgenic mice**

(A) Strategy for targeted insertion of an inducible EGFP_CA-Rac1 into Exon 1 of the murine \( \text{Hprt1} \) locus on chromosome X. The tetracycline response element (TRE) allows for DOX inducible EGFP_CA-Rac1 expression when crossed to lineage-specific rtTA-transgenic mouse lines. (B) Confirmation of targeted insertion and verification of EGFP_CA-Rac1 activity in transgenic mice. Long genomic PCR for the 5' and 3' insertion sites confirmed homologous recombination in ES cell clone 36 (parental KH2 ES cells are used as the negative control). Lanes 1 and 3: Primer1 (in 5' EGFP transgene sequence) and Primer3 (400bp down stream of the 3' arm). Lanes 2 and 4: genomic PCR by Primer2 (in Rac1 transgene sequence) and Primer3. (C) XbaI digestion produced specific digestion bands (400bp and 600bp) that confirmed the specificity of the 5' and 3' targeted locus PCR products from Figure 3.2B. Two replicates are shown. (D) Transgenically expressed EGFP_CA-Rac1 is functionally active. EGFP_CA-Rac1 transgenic mice were crossed to \( \text{NPHS2}-\text{rtTA} \) inducer mice to generate PODxRac1 mice. EGFP_CA-Rac1 transgene was induced by feeding DOX to the mice. Transgenically expressed EGFP_CA-Rac1 binds to GST-PBD, which specifically recognizes the active conformation of Rac1. Transgenic EGFP_CA-Rac1 is not induced and is not present to bind to GST-PBD without DOX induction. Representative data from two induced and non-induced PODxRac1 mice are shown.
Figure 3.3 Podocyte specific expression of CA-Rac1 causes proteinuria

(A) Upper panels, without DOX induction, EGFP_CA-Rac1 is not expressed in PODxRac1 mice. Lower panels, after a 4-day DOX induction, EGFP_CA-Rac1 transgene (green) is specifically expressed in glomerular podocytes, confirmed by immunofluorescence colocalization with the podocyte marker, podocin (red). (B) DOX treatment induced fast onset of proteinuria in PODxRac1 mice. Urine samples were collected at the indicated timepoints from the single transgenic EGFP_CA-Rac1 and DOX induced double transgenic PODxRac1 mice at the indicated timepoints. Proteinuria was quantitated by measuring the albumin/Creatinine ratio (Al/Cr) for each sample. Each point represents the Al/Cr ratio from a single mouse measured at the indicated timepoints. (C) The frequency of EGFP+ glomeruli correlated positively with the level of proteinuria (Al/Cr ratio) (Pearson’s r=0.8831, P< 0.0001).
Figure 3.4 EGFP_ CA-Rac1 expression driven by Nphs1-rtTA induces more robust transgene expression and transient proteinuria

(A) Nphs1-rtTA mice were generated and crossed to EGFP_ CA-Rac1 transgenic mice to generate NEFxRac1 mice. After 4 days of DOX treatment, there was robust expression of the EGFP_CA-Rac1 transgene (green) in podocytes, labeled with the podocyte marker, podocin (red). (B) As before, NEFxRac1 mice were treated with DOX for various time periods and proteinuria was measured in the collected urine samples. Compared to the single transgenic (EGFP_CA-Rac1 only) control, NEFxRac1 mice exhibit significant proteinuria as early as 2 days after induction. Each point represents the Al/Cr ratio from a single mouse measured at the indicated timepoints. (C) Similar to PODxRac1 mice, the proteinuria in NEFxRac1 mice peaks around day 4 and then returns to baseline in 28 days. Each point represents the Al/Cr ratio from a single mouse measured at the indicated time points.
Figure 3.5 No obvious pathological changes were detected in histological analysis, and the transgene positive podocytes were lost after prolonged DOX treatment

(A) hematoxylin and eosin (H&E) and periodic acid-Schiff (PAS) stained kidney sections from NEFxRac1 mice induced with DOX for 4 days. The control kidney samples are shown on the left, and the NEFxRac1 tissues are shown on the right. (B) The kidney samples were harvested from 3 NEFxRac1 treated by DOX for 4 days or 28 days. For each kidney sections, the percentage of EGFP+ positive glomeruli was counted.
Figure 3.6 EGFP_CA-Rac1 expression in podocytes is associated with foot process effacement in vivo

(A) Left panel, examination of kidney tissues from DOX treated PODxRac1 mice by transmission electron microscopy demonstrates segmental effacement (E) of podocyte foot processes while the foot processes of neighboring podocytes are intact and appear normal (N). The panels on the right demonstrate the same areas at higher magnification. (B) Scanning electron microscopy (SEM, left panel) and STORM (right panel) imaging techniques were performed on the same glomerulus from a NEFxRac1 mouse treated with DOX for 4 days. For STORM imaging, the sample was stained for laminin β2 (red) and EGFP (green) to label EGFP_CA-Rac1 expressing podocytes. The boxed area is examined in detail in Figure 3.4C. (C) Correlation of STORM and SEM images identifies that EGFP-CA-Rac1 expressing podocyte have effaced foot processes while neighboring, non-transgene expressing podocytes retain intact foot processes. The left panel shows capillary loops with effaced foot processes (arrowheads) and intact foot processes (arrows). The middle panel presents the STORM imaging of the same area with laminin β2 (red) and EGFP (green). The right panel shows a schematized representation of the STORM data overlaid on the SEM image. The glomerular basement membrane marked by laminin 2 is outlined in red. The EGFP_CA-Rac1 podocyte with its effaced foot processes is outlined in green. The intact foot processes of non-transgene expressing podocytes are outlined in blue. (M-mesangial cell, En-endothelial cell)
Figure 3.7 EGFP_CA-Rac1 expression correlates with reduced expression of podocin and nephrin

A NEFxRac1 mouse was induced with DOX for 4 days and glomeruli were stained for nephrin (A) and podocin (C), both shown in red. There is decreased expression of nephrin and podocin in podocytes expressing the EGFP_CA-Rac1 transgene. Quantitation of pixel intensity across the glomerulus demonstrates separation of nephrin (B, Pearson’s r = -0.3840, P < 0.0001) and podocin (D, Pearson’s r = -0.1432 P < 0.0001) signals compared to EGFP_CA-Rac1 signals. Additional measurement of three other glomeruli are shown in (E, nephrin) and (F, podocin). (G) EGFP_CA-Rac1, but not EGFP_CA-RhoA expression in cultured podocytes reduces podocin protein levels. Left panel, cultured podocytes were electroporated with EGFP_CA-Rac1 or EGFP_CA-RhoA expression plasmids. Protein lysates were immunoblotted for podocin, Erk2 and EGFP. Compared to the empty vector control and EGFP_CA-RhoA, EGFP_CA-Rac1 reduced podocin protein levels. Right panel, densitometric quantitation of three independent experiments normalized to Erk2 levels. * P < 0.05 by unpaired t-test. (H) The reduction of podocin levels can be rescued by treatment with the proteasomal inhibitor MG132, but not with vehicle (DMSO) alone.
Figure 3.8 Time-lapse intravital MPM imaging of DOX induced NEFxRac1 mice
Here shows nine images of the same glomerulus in 32 minutes of intravital MPM imaging experiment. The green channel is EGFP_CA-Rac1 positive podocytes. The red channel shows the capillaries in the glomerulus. The blue channel shows second harmonic signal from collagen matrix. The white arrow shows the membrane dynamics of the podocytes over time.
Figure 3.9 Z-stack images of EGFP_CA-Rac1+ podocyte that attach to the epithelium of the renal tubules

The green channel is EGFP_CA-Rac1 positive podocytes. The red channel shows the tubular epithelium and interstitial capillaries that absorbed Dylight594-labelled lectin. The images were generated from a 3D image file.
Figure 3.10 EGFP_CA-Rac1+ podocytes extended protrusions that penetrates the tubular epithelium and touched the basal side of the interstitial endothelial cells

The green channel is EGFP_CA-Rac1 positive podocytes. The red channel shows the tubular epithelium and interstitial capillaries that absorbed Dylight594-labelled lectin. The arrows show an interstitial capillary. L: the lumen of a renal tubule. P: podocytes that express EGFP_CA-Rac1.
Chapter 4. A Role for Genetic Susceptibility in Sporadic Focal Segmental Glomerulosclerosis

4.1 Abstract

Focal segmental glomerulosclerosis (FSGS), a disease of kidney podocytes is a significant cause of chronic kidney disease. FSGS can be caused by multiple factors including genetics, medication toxicity, obesity, inflammation and viral infection. In the fraction of FSGS subjects with a family history, highly penetrant disease genes have been identified. The identification of APOL1 as a susceptibility factor for FSGS in African Americans with HIV suggests that genetic factors may play a broader role in enhancing susceptibility to FSGS. Here we used sequencing to investigate whether genetics plays a role in the majority of FSGS cases called primary or sporadic FSGS where medications and viruses have been ruled out as causes. We identified 16 potential new FSGS genes and found that over 46% of FSGS subjects may carry susceptibility genetic variants. Using a novel mouse method based on the manipulation of a murine embryonic stem cell line with a genetic background susceptible to FSGS, we validated three of the top four FSGS candidate genes. Our work supports the feasibility of studying the role of genetic background in the susceptibility to disease in humans.
4.2 Introduction

The glomerulus of the kidney is a specialized capillary bed that generates an ultrafiltrate that after modification by the kidney tubule system becomes urine. Diseases of the glomerulus often lead to chronic kidney disease, a major health care problem affecting between 5-10% of the adult population in developed countries [3]. Treatment options are limited, in part owing to the poor understanding of the pathogenesis of disease. Better insights into the root cause of disease offer hope for eventual improvement of this situation.

One of the most common glomerular syndromes is focal segmental glomerulosclerosis (FSGS). The pathologic change of FSGS is a scar that develops focally (in some but not all glomeruli) and segmentally (in only part of a glomerulus). While originally considered a disease, FSGS is now thought to consist of a variety of different syndromes. These include primary (idiopathic) FSGS that is thought to be caused by a circulating factor and secondary FSGS, which may be caused by viruses, medications and genetic mutations. The most common form of secondary FSGS follows glomerular hyperfiltration arising from mismatch between metabolic load and glomerular capacity, associated with obesity, low birth weight, reduced renal mass as well other causes. Genetic mutations can be sufficient by themselves to cause disease (Mendelian), or can increase susceptibility to FSGS by potentiating the effects of environmental factors.

The glomerulus is composed of three different cell types: endothelial cells, mesangial cells and epithelial cells known as podocytes. The podocyte is an unusual cell that covers the outside of the capillary wall and interdigitates with other podocytes to create small slits that allow the
passage of fluid and small solutes into urinary space. It is now clear that podocyte dysfunction is responsible for FSGS as well as other glomerular diseases such as minimal change disease, membranous glomerulopathy and congenital nephrotic syndrome. Current models suggest that increased podocyte loss is the primary lesion in FSGS [24, 26, 133, 134].

Over the last 10 years, highly penetrant disease genes has been identified in the fraction of FSGS subjects with a family history [135–137]. However, they do not explain the majority of the disease population, which are non-familial. In terms of number of people affected, the most significant genetic contributor to FSGS susceptibility identified to date is APOL1. FSGS associated alleles of APOL1, called G1 and G2, are common in West African populations, possibly as a consequence of providing resistance to trypanosomiasis [19, 138, 139]. The presence of two variant alleles significantly increases the risk of arterionephrosclerosis (hypertensive nephropathy) (Odds Ratio (OR) = 7), FSGS (OR = 17) or HIV associated nephropathy (OR = 29) in African [19, 20]. Approximately 13% of African Americans carry two variant alleles and are at increased risk for chronic kidney disease. As these variants are absent from individuals lacking any African ancestry, they are not documented to play a role in FSGS susceptibility in other ancestries, and by themselves largely explain the increased frequency of FSGS among African-Americans. Despite this, mechanisms by which APOL1 variants cause or predispose to glomerular damage remain unknown.

Here, we used high-throughput sequencing of FSGS subjects to investigate whether genetics plays a broader role in the majority of FSGS cases that cannot be explained by the familial FSGS genes. Since FSGS is considered to be a disease of podocytes, we focused our
sequencing analysis on 2500 genes that are highly and/or specifically expressed in podocytes. DNA from 214 FSGS subjects including 192 sporadic cases and 22 familial cases were sequenced (Table 4.1). DNA samples of FSGS subjects were mostly from a multi-center NIH study of biopsy-confirmed FSGS [20], but also included some subjects diagnosed with FSGS by kidney biopsy at Washington University. All subjects provided informed consent for genetic studies. Control subjects sequenced for an autism study were used as controls [66]. Because the controls were of European ancestry, we focused on FSGS subjects of similar genetic ancestry.

A major challenge of large scale sequencing studies will be to develop strategies that will allow for candidate susceptibility genes identified through association studies to be experimentally validated as functionally relevant to disease. Since there is no in vitro assay for podocyte injury leading to FSGS, validating candidate genes here requires the use of an animal model. We developed a system based on embryonic stem (ES) cells from a susceptible background for FSGS that allows for efficient method for targeted delivery of shRNAs, and that uses a method to generate mice that are close to 100% derived from the ES cells eliminating the need for breeding. Scaling up our system will allow for large numbers of candidate genes constituting the network of FSGS genes to be validated which will provide critical insight into the pathogenesis of this disease syndrome. In addition, our experimental approach should be broadly applicable to studying other oligogenic diseases.

4.3 Methods

4.3.1 Exon capture and sequencing
Sample preparation and sequencing were carried out using standard protocols for targeted capture and Illumina sequencing. In brief, genomic DNA was fragmented to 150-200bp using a Covaris E220 focused ultra-sonicator. The ends of the fragmented DNA were repaired using a mixture of T4 DNA polymerase, Klenow polymerase and T4 polynucleotide kinase. Subsequently, adapters for Illumina sequencing were ligated onto the fragments. These libraries were then hybridized to biotinylated DNA probes from regions of interest (manufactured by MyGenostics, Baltimore, MD). After washing away DNA libraries that bound non-specifically to the probes, DNA of interest was recovered using Dynabeads® MyOne™ Streptavidin T1 (Life Technologies). Resulting DNA libraries were amplified, if needed, to provide enough products for sequencing on an Illumina HiSeq 2500.

4.3.2 Variant calling and data quality control

We performed alignment of the raw sequencing data and variant calling according to GATK best practices with BWA/Picard/GATK software pipeline of the Broad Institute. To insure that we are working with equally well-covered loci in both cases and controls we have performed a QC separately on cases and controls genotypes applying following filters: (1) Keep only SNPs; (2) Keep variants that PASS all GATK quality filters; (3) Keep genotypes with DP>10,GQ>30,AB for hets 0.3<AB<0.7, for homozygous alternative AB<0.3; (4) Keep variants with less than 5% of missing genotypes. After applying these filters we have combined variants from cases and controls keeping only those variants with less than 5% of missing genotypes in both cases and controls. Our final dataset contained 16108 SNPs in 1874 genes.

4.3.3 Principle component analysis and case-control matching
PCA was performed with Eigenstrat software on the common (MAF>5%) variants found in autosomes only. We computed a Euclidian distance from each point on the PCA plot to the origin and plotted distributions of this parameter for both cases and controls. Using 3-sigma rule to remove the samples that appeared to be outlying from the distribution. This resulted in 30 samples of mixed Hispanic ancestry to be removed from initial data.

Sample statistics and case-control matching metrics were computed using Plink-seq. We have used number of variants called per sample, number of heterozygous genotypes per sample and number of genotypes with minor allele per sample as a metrics representing genetic background of the cohort. We established similarity between the genetic background of cases and controls by matching mean and variance of case and control distributions for every metric. We tested the validity of this approach by running Fisher’s exact test on the common variation and QQ-plot of the p-values showed no inflation confirming absence of the population stratification in the case-control dataset (Supplementary Figure 4.1).

4.3.4 Mouse strains and antibodies

Cd2ap<sup>+/−</sup> mice were generated in our previous study [77]. Synpo<sup>+/−</sup> mice were obtained from Peter Mundel’s laboratory [140]. The Nphs1-rtTA3G (NEFTA) strain was a gift from Dr. Jeffrey Miner’s laboratory [126]. Dlg5+/- strain was a gift from Dr. Valori Vasiokin’s lab [141]. All mouse strains were genotyped by published protocol. All animal experiments were conducted with the approval of the Washington University Animal Studies Committee. The antibodies used for immunoblotting were mouse anti-XFP (632381; dilution, 1:10,000; Clontech), rabbit anti-ERK2
(sc-154; dilution, 1:5,000; Santa Cruz Biotechnology), mouse anti-β-actin (A2228; dilution, 1:10000, Sigma), and rabbit anti-CD2AP (generated in our previous study, dilution, 1:10000)

4.3.5 Generation of a male Cd2ap+/-, Synpo+/-, NEFTA+ ES cell line

To generate a male ES cell line that was sensitized to FSGS, we bred Cd2ap+/-, Synpo+/- males with NEFTA+ females. The females were superovulated using standard methods. After mating, the embryos were isolated at the eight-cell stage (morulae), and cultured overnight in KSOM (Millipore MR-121-D) micro-drops overlaid with mineral oil at 5% CO2 and 37C. Blastocysts were transferred one per well, into 48 well plates with gamma-irradiated MEF feeders and standard ESC Medium that contains 15% ES qualified fetal bovine serum (SH30070.03E, Hyclone). The inner cell mass (ICM) was allowed to grow out. The ICM outgrowth was trypsinized after 5-7 days depends on the size and shape of the outgrowth, and cultured until ES colonies were identified. The colonies were expanded, frozen back and genotyped by using standard methods.

4.3.6 Generation of miR30-shRNA knock-in transgenic mice

In our previous studies, we developed a method to integrate a single copy of transgene into Hprt1 locus, and use 6-thioguanine to select for ES clones with homologous recombination [142]. Based on this method, we further developed a double selection method, which significantly improve the chance of obtaining a positive ES clone.

A PGK-Puro cassette was inserted between the left and right arm of the pHPRT targeting vector. The miR30-based shRNA-expressing transgene that was driven by the tetracycline-responsive promoter (TRE) was inserted between the left arm and the PGK-Puro
cassette. Linearized targeting vector was transfected into ES cells that were growing at extension phase. At 24 hours post transfection, the ES cells were treated by 1µg/mL puromycin for 48 hours. Then the survived ES cells were passaged once. After 48 hours, the ES cells were treated by 6-thioguanine for 48 hours. Survived ES cell colonies were picked, expanded, and examined by genomic PCR across the right arm (Forward primer: 5'-CAAGCCCGGTGCCTGATCTAGATCATCAATC-3'; Reverse primer: 5'-CTGTAAGGTCTCTGAACGATTTGCAC-3'). Positive ES cells were then stocked for injection.

4.3.7 Laser assisted microinjection

The ES cells were maintained at extension phase before the injection. Eight ES cells were injected into a recipient embryo at eight-cell stage by following a standard protocol published previously [143]. Since the ES cell line produces mice with agouti coat color, albino B6 (C57BL/6J-Tyr<sup>C2J</sup>) mice were used as host embryos for direct evaluation of chimerism by coat color.

4.3.8 Cell culture and lentivirus infection

Immortalized murine podocytes were maintained and differentiated as described previously [122]. To examine the knockdown efficiency of Cd2ap-sh877, podocytes were infected with lentiviral vectors encoding miR30-sh877. A control lentiviral vector encoding miR30-FF3 that targets fire fly luciferase cDNA was used as a control. The CD2AP expression was examined by immunoblotting of the whole cell lysates.

4.3.9 Design and validation of the miR30-shRNA constructs for genes of interest
The shRNA oligo sequences were picked using an online algorism (http://katahdin.cshl.org/siRNA/RNAi.cgi?type=shRNA) as described [144]. The miR30-shRNA backbone was sub-cloned by PCR from pPRIME-CMV-GFP-FF3 (https://www.addgene.org/11663/) and inserted into pcDNA3.1-Zeo(+) vector to generate pcMIR vector. To examine the knockdown efficiency, the miR30-shRNA construct and its artificial target (Figure 4-S3B) were co-transfected into HEK293T cells at molar ratio 5:1. The expression of EGFP in whole cell lysates was examined by immunoblotting.

4.3.10 Albumin-creatinine assay

Mouse urine samples were collected at the time points indicated in the figures, and urinary albumin (E90-134; Bethyl) and creatinine (DICT-500; BioAssay Systems) were quantified by enzyme-linked immunosorbent assays (ELISA) according to the manufacturers' protocols.

4.3.11 Transmission electron microscopy

Portions of kidney cortex were fixed with 2% paraformaldehyde and 2% glutaraldehyde. Specimen processing, ultrathin sectioning, and imaging were performed by the Electron Microscopy (EM) Core Facility at Washington University.

4.4 Results

4.4.1 Identification of podocyte genes for sequencing

To identify susceptibility genes for FSGS, we designed a custom exome capture reagent that focused on genes that were highly expressed in podocytes and/or previously implicated in FSGS. We began with five genes reported in OMIM in which heterozygous mutation confers risk
to idiopathic FSGS: TRPC6 [17, 145], INF2 [13], APOL1 [19], ACTN4 [8], and CD2AP (Kim et al., 2003). Pathway analysis was performed to identify ~200 genes that are functionally linked to known FSGS genes and other syndromic or recessive genes. 677 genes were added based on their high expression in microdissected human glomeruli [147] and 1600 were the human orthologs of the most highly expressed genes identified by DNA microarrays in mouse podocytes [122, 148, 149]. Our total capture, which we term the “podocyte exome”, included about 2500 genes comprising a total of 7Mb. (Figure 4.1A).

4.4.2 Variant association analysis of FSGS patients of European ancestry using Next Generation sequencing

DNA from 225 biopsy confirmed FSGS patients that were of European ancestry was sequenced. Patient DNA samples were mostly from a multi-center NIH study of FSGS [150], but also included some FSGS patients diagnosed at Washington University; all subjects provided informed consent for genetic testing. The patient population included mostly sporadic but also some cases of familial FSGS patients. Patients sequenced for an autism study, lacking evaluation for kidney phenotypes were used as controls [66]. Because the controls were of European ancestry, our plan was to focus on FSGS patients of similar genetic ancestry. The podocyte exome was captured from the FSGS patients and sequenced using standard technologies with an average coverage of 200X.

Since the FSGS patient sequences were generated using a different platform and sequenced at a different institution than were the controls, we needed to validate that the two data sets were comparable as this is critical for unbiased analysis. After eliminating two case
samples with <20x average coverage, data from all case and control samples were processed in a single batch with raw data was aligned to the human genome using the BWA/Picard pipeline at the Broad Institute and subsequent variant calling was performed using GATK Unified Genotyper (Figure. 4.1B) [64, 151, 152].

The depth of coverage of protein coding exons targeted by both capture reagents was compared between cases and controls; only those exons covered adequately (>20X) in both cases and controls were advanced into the analysis stage. In summary, 16784 exons and 2769942 base pairs were confidently covered in both case and control cohorts, resulting in 16,008 SNPs and 1724 genes analyzed in the final dataset. The SNP calls were equally represented in cases and controls (GQ>30 and rate of missing genotypes less than 5% in each cohort) (Figure 4.1C).

Next, principal component analysis (PCA) was performed to define an appropriately ancestry matched case-control sample set (Figure 4.2A,B). Thirty patient samples were removed because a mixture of Hispanic ancestry was detected. Three more samples were removed because the call-rate of SNPs was less than 95%. The number of SNPs, heterozygous genotypes, and genotypes containing an alternative allele per sample were similar between cases and controls. This reassured us that we could move forward with association analysis (Figure 4.2C,D,E). Our final dataset contained 179 cases and 378 controls of European-American ancestry as determined by PCA (Table 4.1). Accuracy of this analysis strategy was further confirmed by resequencing key SNPs using Sanger sequencing.

4.4.3 Identification of new candidate genes by testing single variant associations
An association test examining single variants (minor allele frequency (MAF) >1%) was performed using Fisher’s exact test. Ten common variants from nine different genes were significantly enriched in cases versus controls with p-value < 0.001 and odds ratios > 4 (Table 4.2). These variants were found only in the sporadic cases and the controls. No common variant was identified in the familial cases. Four cases carried a single copy of the APOL1 G1 variant (G1); this allele is present in 29% of African Americans but only 0.03% of European Americans and confers FSGS susceptibility when two APOL1 risk alleles are present [150]. A search of the NHLBI-Exome Sequencing Project database showed that four other single variants enriched in our cohort (WNK4, KANK1, IL36G, ARHGEF17) had MAFs that were more common in African Americans. These variants were distributed across many patients, and no variant was linked to other common African SNPs. The specific enrichment for African SNPs in some of our European ancestry patients largely explains the high odds ratios that were calculated for these variants and suggest that some of the risk variants could be ancestral variants.

4.4.4 Identification of new candidate genes by rare variant analysis

Rare variants (MAF < 1%) were analyzed by using tests that compare the total numbers of rare variants between cases and controls. We used three such tests: the burden test [153], the variable threshold test [154] and the C-alpha test [155]. Because the effect sizes of genetic variants differ, the accuracy of each method can vary depending on the specific situation. Using a p-value < 0.001 as a cutoff, we identified 11 genes as potential FSGS susceptibility genes (Table 4.3): WNK4, APOL1, DLG5, GCC1, XYL1, KAT2B, BPTF, COL4A4, NID1, EPHX1 and EPHB6. Two of these genes, WNK4 and APOL1 were also identified by common variant analysis.
In support of our approach, COL4A4, identified by rare variant analysis, was recently identified as an FSGS susceptibility gene [156]. With the exception of APOL1 and COL4A4, none of the identified genes has been previously identified as FSGS susceptibility genes. Thus, we identified 16 potential new FSGS genes.

4.4.5 Analysis of known familial FSGS genes

Family studies have identified about 20 genes as the cause of familial FSGS [157, 158]. To determine, whether these genes are also involved in sporadic FSGS, we assessed the frequencies of rare deleterious (missense/nonsense) coding variants in 20 of these genes in cases and controls (Table 4.4). Among cases, 46/179 subjects (32%) had at least one rare variant in these genes compared to 13/378 controls (3.4%, p value = 4.7e-14) (Table 4.1 and 4.5). The distribution of variants between familial and sporadic cases was similar and consistent with previous studies that about 30% of steroid resistant nephrotic syndrome cases have a variant in a known disease gene [159]. There was also a difference in the total number of unique rare variants identified in cases (59 variants in 179 cases) versus controls (15 variants in 378 controls). The significance of this finding was tested using a permutation analysis of groups of 20 genes randomly chosen from our dataset. This showed, however, that 27% of random sets of 20 genes had a p-value that was similar to or below 4.7e-14, suggesting the presence of novel FSGS genes with strong genetic effects in our dataset.

4.4.6 Development of a sensitized mouse system to identify potential FSGS disease causing genes

The FSGS syndrome likely involves diverse injury pathways and so no single in vitro
system is available to test whether a particular gene variant might induce FSGS. We therefore developed a genetic system in mouse, to examine the function of candidate genes in vivo in the kidney. Our strategy involved knocking down the expression of candidate genes in a mouse genetic background that is prone to develop FSGS. Previously, we showed that mice that are heterozygous for two podocyte genes, *Cd2ap* and *Synpo* (encoding synaptopodin), developed FSGS with an incomplete penetrance (~25%), with albuminuria, a sign of podocyte dysfunction not apparent until animals are about six months of age [21]. We reasoned that if impairing the function of a candidate gene worsened the phenotype on this transgenic background, by either enhancing the penetrance or leading to an earlier onset of disease, it would validate the specific candidate gene.

We generated ES cells from mice that were *Cd2ap*\(^{+/\!-}\) *Synpo*\(^{+/\!-}\) using standard methods (Figure 4.3A). To induce RNAi expression in a podocyte specific fashion, the ES cells also express a podocyte-specific and doxycycline-inducible trans-activator (*Nphs1*-rtTA3G) [126]. The rtTA3G is a synthetic transcription factor that binds and transactivates promoters that contain the bacterial tet-operator sequence [160]. After confirming the genotype of the ES cells (Figure 4.3B), we confirmed that the ES cells could generate mice with high-level chimerism using the traditional method of blastocyst injection. Using the method of laser-assisted microinjection into 8-cell embryos [143], we also validated that these ES cells could generate mice that were close to 100% derived from the ES cells as assessed by coat color (Figure 4.3C). As expected based on the genotype (Huber et al., 2006), about 50% of the mice generated from these ES cells developed mild proteinuria after three to four months of age (Figure 4.3D).
We then used homologous recombination to integrate a single copy of the RNAi transgene into the mouse *Hprt1* locus to eliminate variability that could result from the random integration of an RNAi transgene [142]. The *Hprt1* locus is on the X chromosome and because the ES cells were male, targeting of the transgene results in complete loss of the *Hprt1* gene resulting in resistance to 6-thioguanine (6-TG) toxicity (Figure 4.4A). This method was efficient as over 90% of the ES cell colonies that survived selection had a correctly targeted recombination event (data not shown).

We validated the system by testing an RNAi for *Cd2ap*. Mice that are *Cd2ap*+/− exhibit normal renal function and have normal lifespans, but mice that completely lack *Cd2ap* expression develop severe proteinuria shortly after birth [77]. To control shRNA expression by doxycycline, shRNA oligos were embedded into the miR30 backbone that allows for RNA polymerase II transcription [161]. We tested multiple *Cd2ap* specific RNAi’s *in vitro* for their ability to inhibit *Cd2ap* expression (Figure 4.4B and 4.5A) and the best one (sh877) was targeted into the *Hprt1* locus of our ES cell line (Figure 4.5B). An RNAi targeting the firefly luciferase gene was used as a control. Laser-assisted microinjection generated 16 animals that were nearly 100% derived from the ES cell based on the completely agouti coat-color.

When the chimeric mice were two weeks of age, half were given doxycycline (DOX) in the drinking water to induce shRNA transgene expression. Four weeks later, oodocyte function was assessed by monitoring albumin leakage into the urine by measuring the ratio of albumin to creatinine ratio. All of the DOX treated mice developed proteinuria with an average albumin/creatinine ratio that was almost 150 fold higher than in the control animals $\left(1.5 \times 10^5\right)$
compared to $1.0 \times 10^3$ (Figure 4.4C). The proteinuria was sustained during eight weeks of DOX treatment and was reversible, as proteinuria abated when DOX was removed (Figure 4.5D). Histological analysis also confirmed the presence of proteinuria, as protein casts were consistently present in tubules (Figure 4.4D). Electron microscopic examination of the kidney showed widespread foot process effacement, another marker of proteinuria (Figure 4.4E). Control RNAi mice targeting firefly luciferase showed no proteinuria after DOX treatment (Figure 4.4F). This verified that our RNAi strategy could be used to test candidate FSGS genes.

4.4.7 Testing candidate FSGS genes using the novel mouse genetic system

To validate our system, we chose four genes, \textit{WNK4}, \textit{DLG5}, \textit{ARHGEF17} and \textit{KANK1}. \textit{WNK4} was chosen as it was identified by single variant analysis and by all three rare variant tests. \textit{ARHGEF17} and \textit{KANK1} were identified by single variant analysis but not by rare variant analysis. \textit{DLG5} was identified by all three rare variant tests but not by single variant analysis. Because the mouse ortholog for human \textit{KANK1} is unclear, we targeted \textit{Kank2} in addition to \textit{Kank1}.

Multiple shRNAs were generated for all five candidate genes and their efficacy was first tested in vitro (Figure 4.6A, and 4.7A,B). Validated shRNAs were then ligated into targeting constructs that were transfected into ES cells. After selection for loss of \textit{Hprt1}, mice were genotyped by PCR to confirm homologous recombination. Two independent clones for each candidate gene were selected for microinjection and 15-30 mice were generated by laser-assisted injection for each clone. Coat color verified that the chimerism for each animal was close to 100%.
At two weeks of age, half of each cohort was given DOX in the drinking water to induce expression of the transgene. Proteinuria was assessed at four weeks and eight weeks after DOX treatment, at which time mice were sacrificed for histological and electron microscopic examination (Figure 4.6B~F). Induction of both the \textit{Wnk4, Arhgef17} and \textit{Kank2} RNAi transgenes induced substantial proteinuria four weeks after induction with doxycycline with a level of proteinuria that was between 150-200 fold higher than the controls. For both lines of mice, proteinuria was attenuated after eight weeks (Figure 4.6B, D, E). The \textit{Kank1} RNAi mice showed only modest proteinuria at the four-week time point but proteinuria increased significantly after that (Figure 4.6C).

In contrast, \textit{Dlg5} RNAi mice did not develop significant proteinuria at any time up to 12 weeks after induction (Figure 4.6G and 4.7C). We validated this result by breeding \textit{Dlg5}^{+/−} mice to our sensitized background (\textit{Cd2ap}^{+/−}, \textit{Synpo}^{+/−}). Up to six months of age, no significant difference was detected between triple heterozygous mice (\textit{Dlg5}^{+/−}\textit{Cd2ap}^{+/−}, \textit{Synpo}^{+/−}) and double heterozygous (\textit{Cd2ap}^{+/−}, \textit{Synpo}^{+/−}) littermates (Figure 4.6F). This suggested that \textit{Dlg5} is not an FSGS susceptibility gene or that it may not be epistatic with \textit{Cd2ap} and \textit{Synpo} in podocytes. As expected, electron microscopic examination of the kidneys showed podocyte foot process effacement from \textit{Arhgef17} \textit{Kank1}, \textit{Kank2} and \textit{Wnk4} RNAi mice, but not in \textit{Dlg5} RNAi mice (Figure 4.7D).

### 4.5 Discussions

We added \textit{KANK1}, \textit{WNK4} and \textit{ARHGEF17} to the list of 20 known FSGS genes and
reanalyzed the differences between cases and controls. Addition of these three genes increased the p-value to 1.6e-23. Testing random sets of 23 genes by permutation analysis in cases and controls showed that only 0.04% of random sets equaled or matched the p-value for these 23 genes. This supports the idea that genetic variants in a specific subset of genes may function more broadly to create a susceptible background for the development of sporadic FSGS.

The role of genetics in the pathogenesis of FSGS was thought to be restricted to those with a family history. Familial studies have identified ~20 FSGS disease genes with specific variants that are highly penetrant in these families. The identification of APOL1 as a sensitizing factor for FSGS in African Americans with HIV demonstrated that genetic factors may also play a role in enhancing susceptibility to FSGS by environmental factors [19, 20]. Here we found that DNA variants in a set of 23 or more genes may play a role in over 46% of non-familial FSGS cases presumably by enhancing podocyte susceptibility to injury. The role of genetics in sporadic cases is likely much weaker than in the familial cases. Common variants were found exclusively in the sporadic cases and in addition, a significant fraction of sporadic cases had more than one variant, a feature that was not seen in any of the controls or the familial cases. Our analysis was likely facilitated by our focus on podocyte-specific genes that we reasoned would have a higher likelihood to be involved in FSGS and by our RNAi mouse model that allowed us to begin validating these genes. Since oligogenic genetic effects and environmental factors are likely to be broadly involved in disease susceptibility, our methods could be widely applicable to the study of other rare and common diseases.
Acknowledgement

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4.6 Tables and Figures

Table 4.1 Distribution of single and rare variants in FSGS subjects and controls
A total of 192 sporadic and 22 familial FSGS cases were sequenced. 157 sporadic and 22 familial FSGS cases remained after quality control and were compared to 378 controls. The number of subjects with predicted deleterious variants in 20 known FSGS genes, single variants in the top 9 genes from the common variant analysis and the top 11 genes from rare variant analyses. The total number of subjects with variants in 20 known FSGS genes + the three genes that were validated are also shown.

<table>
<thead>
<tr>
<th>Total number of samples sequenced</th>
<th>Sporadic FSGS</th>
<th>Familial FSGS</th>
<th>Total # of Patients</th>
<th>Total # of Controls</th>
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</thead>
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<tr>
<td>Number of samples that passed QC</td>
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<td>179</td>
<td>378</td>
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<tr>
<td>Number of patients with deleterious variants in 20 known FSGS genes*</td>
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<td>46</td>
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<td>Number of patients with single variants in 9 genes**</td>
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<td>28</td>
<td>14</td>
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<td>Number of patients with rare deleterious variants in 11 genes***</td>
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<td>4</td>
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<td>Number of patients with variants in 3 validated genes****</td>
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<td>33</td>
<td>8</td>
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<tr>
<td>Percentage of patients with variants in 20+3 genes</td>
<td>46%</td>
<td>32%</td>
<td>44%</td>
<td>6%</td>
</tr>
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</table>

* The list of 20 known genes is in the Supplemental Table 2
** Genes identified by common variant analysis (Figure 1A)
*** Genes identified by 3 rare variant analyses (Figure 1B)
**** Genes validated by our mouse model (ARHGEF17, KANK1 and WWK4)
### Table 4.2 Potential FSGS susceptibility genes identified by common variant analyses

15 potential susceptibility genes identified by common variant analysis. The frequency of common variants (MAF > 1%) was assessed in cases versus controls and high scoring variants with odds ratios greater than 2.5 are shown here ranked by p value. For each common variant, the chromosome position, gene name, reference and alternative base pair, overall mean allele frequency, number of alternative alleles in cases versus controls, total number of genotypes in cases, the allele frequency in European vs African Americans from the ESP database, the p value and the odds ratio and the Polyphen prediction score are shown.

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<th>GEN_NAME</th>
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<th>MINU</th>
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**CHROM_POS** - the chromosome position of a common variant
**REF** - reference allele
**ALT** - alternative allele (variant)
**MAF** - mean allele frequency
**MINA** - number of alternative alleles in cases
**MINU** - number of alternative alleles in controls
**REFA** - number of reference genotypes in cases,
**ESP_EA** - the allele frequency in European Americans in the NHLBI-Exome Sequencing Project
**ESP_AA** - the allele frequency in African Americans in the NHLBI-Exome Sequencing Project
**OR** - odds ratio

Bold - genes that are identified by both Fisher's test (Table 2) and at least one of the three rare variant test
Table 4.3 Potential FSGS susceptibility genes identified by rare variant analyses

11 genes identified by rare variant analyses. Rare missense, and nonsense variants (MAF < 1%) were pooled together for rare variant analysis using the burden, variable threshold and C-alpha tests. The top genes (p-value < 0.001) identified for each test are shown ordered by p value. Genes that were also identified by single gene analysis using Fisher’s exact test are shown in bold.

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Bold - genes that are identified by both Fisher’s test (Table 2) and at least one of the three rare variant test

VT - variable threshold
Genes that are identified by 2 rare variant tests
Genes that are identified by all 3 rare variant tests

Table 4.4 The list of 20 known FSGS genes

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Table 4.5 The list of rare deleterious variants identified in 20 known FSGS genes in the FSGS subjects that we sequenced

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CHROM.POS - the chromosome position of a common variant
GENE_NAME - number of alternative alleles in cases
MINA    - number of alternative alleles in controls
OBSA    -
OBSU    -

NON_SYNONYMOUS_CODING
Figure 4.1 Comparability of variant calls between cases and controls

(A) Principal component analysis (PCA) of variants between cases (+) and controls (x) shows close similarity of ancestry. (B) The PCA data shown in A is depicted as distance from the origin. The 30 case sample with a distance of greater than 0.09 from the origin, suggesting substantial admixture with non-European populations, were removed from further analysis. (C) The number of total variants per sample were similar between cases and controls. (D) The number of heterozygous (HET) genotypes was similar between cases and controls. (E) The number of heterozygous or homozygous genotypes containing an alternative allele was similar between cases and controls.
(A) 2500 genes were clustered and defined as “podocyte exome”. (B) Average sequencing coverage for all patients in FSGS cases. Each bar represents the coverage of a patient DNA sample we sequenced. (C) PCA Plot. (D) Dapple analysis for the interaction network of all gene identified by common and rare analyses.
Figure 4.3 Generating an ES cell line with an FSGS susceptibility genetic background

(A) Male \( \text{Synpo}^{+/-}, \text{Cd2ap}^{+/-} \) and female \( \text{Nphs1-rtTA3G} \) mice were bred together to isolate blastocyst. ES cell lines were derived from these blastocysts. The chance of deriving a male ES cell line that is \( \text{Synpo}^{+/-}, \text{Cd2ap}^{+/-}, \text{Nphs1-rtTA3G} \) is 1/16. (B) Genomic PCR confirms B12 ES cell line is male, \( \text{Cd2ap}^{+/-} \) and \( \text{Nphs1-rtTA3G} \). \( \text{Synpo}^{+/-} \) was also confirmed (data not shown). (C) Mice were generated using laser assisted injection from B12 ES cell line were monitored for the development of proteinuria by measuring the urine albumin/creatinine ratio. Approximately 50% of mice generated from these ES cells treated with or without doxycycline (administered from 2 weeks of age) slowly developed proteinuria over a period of 3-5 months.
Figure 4.4 Generating a system to validate candidate FSGS genes

(A) The targeting strategy used to integrate a miR30-shRNA transgene into Hprt1 locus is shown. A targeting vector with the miR30 transgene and a PGK-puromycin cassette was generated with left and right homology arms containing sequences that flank exon1 of the Hprt1 gene on the X chromosome. Primers (P1, P2, and P3) for PCR validation of homologous recombination are shown. (B) The knockdown efficiency of a miR30 shRNA for Cd2ap (sh877) is shown. A podocyte cell line was transduced with a lentivirus containing the Cd2ap shRNA was blotted for Cd2ap expression. Erk2 was used as a protein loading control. Transduction with an shRNA
targetting firefly luciferase (FF3) serves as a negative control. (C) Mice generated with ES cells with the Cd2ap shRNA recombined in the Hprt1 locus were treated with and without doxycycline for four or eight weeks and urine was analyzed by measuring the urine albumin/creatinine ratio. (D) By light microscopy, Doxycycline treated Cd2ap-RNAi mice manifested proteinaceous tubular casts (asterisks). (E) By electron microscopy, doxycycline dependent foot process effacement was present in Cd2ap-RNAi mice, while podocyte cytoarchitecture is preserved in control mice. (F) Control FF3-RNAi mice were treated with and without doxycycline for four and eight weeks and urine analyzed by measuring the albumin/creatinine ratio.
Figure 4.5 Supplemental figure for Figure 4.4

(A) The efficiency of 3 miR30-shRNAs that targets Cd2ap. A mouse CD2AP-EGFP construct was co-expressed with different miR30-shRNA constructs in HEK293 cells. The expression level of CD2AP-EGFP was detected by immunoblotting. Actin immunoblotting was used as loading control. FF3, A miR30-shRNA that targets fire fly luciferase was used as control. The design of genomic PCR and the example of results that validate homologous recombination in ES clones. Forward primer (P1: 5’-CAAGCCCGGTGCCTGATCTAG ATCATAATC-3’) was designed at the end of puromycin resistant cassette. Two reverse primers were designed out side the Right Arm. (P2: 5’-CTGTAAAGGTCTCTGAACTACCAATTGCAC-3’, and P3: 5’-GAGACTAAGGCAGGAGGATTCCAGGTTTG -3’). (B) PCR validation of homologous recombination. The arrows points the specific PCR products for the PCR reactions by using P1+P2 and P1+P3, and the PCR products were confirmed by restriction digestion with desired sizes of digested DNA fragments. (C) The system of podocyte-specific, DOX-inducible RNAi. (D) Cd2ap-RNAi mice showed DOX-dependent proteinuria. The DOX treatment was stopped after 8 weeks. The urine samples were collected at 4, 8, 12 and 16-week time points. Albumin/creatinine ratio was measured and plotted. (E) Untreated Cd2ap-RNAi mice showed normal foot processes.
Figure 4.6 Validation of five FSGS candidate disease genes
(A) To select the best shRNA construct, multiple shRNAs were tested for Arhgef17, Dlg5, Kank1, Kank2 and Wnk4 by inhibiting expression of a construct containing the GFP-tagged target sequence in 293 cells (Figure 4-S3A). GFP immunoblotting was used to determine the best shRNA for each gene. (B-E) Mice were generated by laser-assisted microinjection of ES cells with the specific shRNAs for the indicated genes. After treatment with or without doxycycline to induce expression of the transgene, urine albumin/creatinine ratio was measured after 4 and 8 weeks. As shown, shRNA knockdown of Arhgef17, Kank1, Kank2, and Wnk4 increased proteinuria at 4 and 8 weeks, (F-G) By contrast, neither Dlg5 shRNA knockdown nor a Dlg5+/− mouse on a susceptible genetic background (Cdap+/− and Synpo+/−) increased albuminuria.
Figure 4.7 Supplemental figure for Figure 4.6

(A) The system was developed to test the RNAi efficiency of a given miR30-shRNA. The ExonX targeted by the shRNA is amplified by genomic PCR and inserted to the 3’ end of EGFP sequence of pEGFP-C1 vector. The pEGFP-C1-ExonX became an artificial target for the shRNA. The pEGFP-C1-ExonX and miR30-shRNA constructs were co-expressed in 293 cells, and the expression level of EGFP was detected by immuneblotting. (B) Immunoblotting results for each miR30-shRNA oligo designed for the candidate genes. The FF3 miR30-shRNA was used as control for each miR30-shRNA oligos. Actin immunoblotting was used as loading control. The expression of EGFP was used as an indicator of the RNAi efficiency, the miR30-shRNA oligo that had the best RNAi efficiency was inserted into pHPR vector to generate RNAi ES cells and mice. (C) Dlg5-RNAi mice showed no proteinuria after 12 weeks of DOX treatment. (D) Electron microscopy of each RNAi mice showed the foot process effacement.
Chapter 5. Conclusions and Future Directions

5.1 Conclusions

5.1.1 The balance between Rho and Rac activity maintains the morphology and function of the podocyte.

To form properly interdigitated foot processes, podocytes established organized actin cytoskeleton. In the foot processes, actin polymerize into at least two different structures that are observed by transmission electron microscopy, the dense actin bundle and the loose cortical actin network. *In vitro*, active Rho promotes the actin bundle formation, while active Rac promotes the branched actin network. Podocytes regulate the spatial balance of Rac and Rho activities. The active Rho could maintain the thick actin bundles along with other actin bundling proteins such as alpha-actinin4 and INF2, while active Rac could keep the actin network next to the silt diaphragm.

Over-active Rac or Rho in podocytes could cause transient or permanent damages. Transgenic mice that express the constitutively active mutant of either Rac (CA-Rac) or Rho (CA-Rho) are lethal in embryonic stage, which indicates the high toxicity of both constitutively active mutants [162]. Inducible expression of CA-Rho in podocytes cause slow onset of proteinuria and permanent FSGS [49]. In contrast, our research showed that inducible expression of CA-Rac1 caused much faster onset of proteinuria, which can be detected as early as 2 days post doxycycline treatment. This observation supports the hypothesis that foot process
effacement is analogous to lamellipodia. Our result also supports that loss of podocyte foot processes is a direct cause of albumin leakage. Long-term expression of CA-Rac1 could lead to chronic damage such as glomerulosclerosis.

Podocytes express high level of ARHGAP24, a Rho-activated Rac/Cdc42 GAP. We identified this gene initially by comparing expression profiles of differentiated versus undifferentiated mouse podocytes, and later we found mutations of this gene associated with familial FSGS. This protein is also called “FilGAP”, because it binds to filamin A, a protein that dimerizes and binds to branched actin filaments [94, 114]. Thus, ARHGAP24 could work as a barrier between the cortical actin network and the actin bundles to keep the active Rac from the center of the foot processes.

5.1.2 The genetic susceptibility could explain over 46% of the non-familial FSGS subjects.

Our genetic study and podocyte-specific RNAi mouse model identified and validated genetic susceptibility to FSGS in non-familial FSGS cases. FSGS was thought mainly caused by environmental factors such as virus infection, and chemical toxins [3]. Studies of familial FSGS cases have identified about 20 FSGS susceptibility genes, but only a few of them were recently found mutated in non-familial cases. Here we identified another 16 genes that could potentially involve in the susceptibility of non-familial FSGS. We validated 4 genes in our podocyte-specific RNAi mouse model, ARHGEF17, KANK1, KANK2 and WNK4. We also observe that variants of different genes presents in the same non-familial FSGS subjects but not in familial FSGS subjects. This observation suggests that a combination of low penetrant genetic risks could cause susceptibility to FSGS, which was first shown by our bigenic mouse models [21]. The
oligogenic genetic factors of non-familial FSGS was not considered as important risk factors before, but our research suggests that non-familial FSGS could be caused by oligogenic genetic factors plus an environmental trigger, such as virus infection, chemical toxicity, and a circulating factor in the serum.

5.1.3 Actin Cytoskeleton pathway is the major pathway of susceptibility to FSGS.

Among the known FSGS genes and the susceptibility genes that we identified from our sequencing approach, we found that a large proportion of these gene are involved in regulation of actin cytoskeleton. In the 20 known FSGS genes (Table 4.4), 9 genes either directly interact with actin filaments (INF2, ACTN4, MYO1E and CD2AP), or regulate the activity of Rho family GTPases (ARHGAP24, ECT2, NPHS1, NPHS2, and TRPC6). 3 genes (COL4A3, COL4A4, and LAMB2) are extracellular matrix genes that connect to intracellular actin network through integrins and focal complexes. Thus over half of the familial FSGS genes influence the actin cytoskeleton. The secretion of extracellular matrix proteins is also regulated by actin cytoskeleton. In the susceptibility genes we validated, 3 out of 4 regulate the actin through Rho family GTPases, KANK1 [163], KANK2 [164] and ARHGEF17 [165]. The enrichment of mutants in actin-associated and actin-regulatory genes in FSGS patients support the hypothesis that the natural susceptibility to FSGS comes from a weakly organized actin cytoskeleton in the podocytes. Previous mouse model and human genetic studies also support this hypothesis. Cd2ap deficient mice develop normal foot processes in the beginning and do not have any protein leakge before 14 days of age. However, after 14 days post natal, Cd2ap deficient mice start to exhibit foot process effacement and proteinuria. In children, NPHS2 mutations account
for about 30% of steroid-resistant nephrotic syndrome [159], which also have proteinuria and foot process effacement. These patients have normal kidney function and foot processes before the syndrome starts. Thus the genetic lesion of NPHS2 might also lead to an actin network that is susceptible to damage. Minimal change disease and steroid-sensitive nephrotic syndrome, however, show transient proteinuria and foot process effacement, because the actin structure in the podocytes could recover from the diseases to regenerate foot processes.

5.1.4 Elevated Rac signal could cause podocyte shattering and shattered podocytes could still be functional to certain extent.

We detected EGFP_CA-Rac1+ podocytes in the urine of DOX induced NEFxRac1 mice. Using intravital multi-photon microscopy (MPM), we observed that EGFP_CA-Rac1+ podocytes enhanced membrane ruffling in the glomerulus and that some EGFP_CA-Rac1+ podocytes attached to the renal tubules. These results indicate that abnormal activation of Rac in podocytes could lead to increased membrane activity, deceased adherence, and podocyte loss. Pathological changes that lead to Rac activation in podocytes could also initiate similar response. At physiological condition, the crosstalk between podocytes and glomerular endothelial cells could suppress the Rac activity, and could also maintain the normal foot process structure and adherence. It is well studied that VEGF secreted by podocytes play an important role in maintaining the endothelium, but which cytokine is maintaining low Rac in podocytes is still not known. Damaging factors that increase Rac activity could compensate this physiological signal, and could cause podocyte shattering into the urine.

The process that shattered podocyte established new interaction with the renal tubules
after was overlooked before, because this event is very rare, and hard to detect by conventional histological techniques. We uncovered this process by intravital MPM experiments. This observation suggests that the defects caused by high Rac activity can be compensated and that shattered podocytes are at least partially functional. How the Rac phenotype is compromised and whether this process has physiological or pathological contribution are interesting questions to answer.

5.2 Future Directions

5.2.1 The temporal change of Rho GTPase activity during podocyte damage.

Given the central role of Rho GTPase signaling in maintaining the normal actin cytoskeleton structure in the podocyte, it is intriguing to know how small GTPases are activated in different podocyte damage models. Do they share similar mechanism? Which GTPase contribute the most to podocyte damage? Intravital imaging is a good approach to investigate these questions. However, new mouse models with novel Rho GTPase sensors are required. The bioluminence resonance energy transfer is a promising candidate sensor technique to investigate dynamic singaling of small GTPases in vivo.

5.2.2 The function of ARHGEF17, KANK1, KANK2 and WNK4 in maintaining podocyte function.

In our sequencing study, we identified multiple candidate genes that could be potential susceptibility gene and validated that decreased expression of ARHGEF17, KANK1, KANK2 and WNK4 in podocytes could cause proteinuria in Cd2ap/Synpo double heterozygous background. It remains unknown how these genes maintain normal podocyte function. Loss of function of
these risk genes might not directly cause podocyte damage, and which environmental factor trigger FSGS under these susceptible background is also an open question.

5.2.4 Can podocytes de-differentiate and become other cell types after they leave the glomerulus?

Using intravital MPM imaging, we observed that podocytes could interact with other cells outside the glomerulus. Is this phenomenon common in other models of podocyte damage? Are these interactions transient? Can podocytes pass the tubular epithelium and enter the interstitial space? Can they de-differentiate into mesenchymal cells? To answer these questions, lineage-tracing experiments are required.
References


Curriculum Vitae

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RESEARCH EXPERIENCE
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• Validate rare genetic variants that associate with focal segmental glomerulosclerosis by an \textit{in vivo} RNAi mouse model (Thesis project).
• Explore how the balance between Rac and Rho activity influences the structure and function of podocytes.
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• Study the mechanism of allosteric activation of Raf kinases.
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• Use RT-PCR to measure the expression level of Eg5 in different pancreatic cancer cell lines

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• Tri-A Outstanding Student Awards, Nankai University, twice in 2005 and in 2008;
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PODIUM PRESENTATIONS


PUBLICATIONS

• Hu J., Yu H., Kornev A.P., Zhao J., Filbert E.L., Taylor S.S. and Shaw A.S., A mutation that
blocks ATP binding creates a pseudokinase that stabilizes the scaffolding function of KSR1, CRAF and BRAF. Proc Natl Acad Sci U S A. 108(15): 6067-72.


**PATENT**