Polycystic kidney disease (PKD) is a cohort of monogenic disorders that result in development and subsequent growth of renal cysts filled with fluid.\textsuperscript{1–5} Cyst enlargement compromises function of surrounding nephrons and progresses to ESRD.\textsuperscript{1,6} In the more common form of PKD, autosomal dominant PKD (ADPKD), which is caused by mutations of polycystin 1 (PC1) and polycystin 2 (PC2), renal cysts are formed along the full length of the nephron with prevalence to the collecting duct (CD).\textsuperscript{1,7} In the rarer and more severe autosomal recessive PKD (ARPKD), renal cyst formation is virtually restricted to the CD.\textsuperscript{1,2,5,8} Mutations of the PKHD1 gene encoding fibrocytin underlie the genetic basis of the disease.\textsuperscript{6,8,9} Although the exact function of the protein is unknown, fibrocytin was shown to be expressed in primary cilia where it can interact and form complexes with PC2, possibly participating in mechanotransduction.\textsuperscript{10–12}

It is accepted that the CD cells elevate $[\text{Ca}^{2+}]_{i}$ in response to mechanical stress arising from variations in tubular flow or tubular composition.\textsuperscript{13–23} Impaired mechanosensitive $[\text{Ca}^{2+}]_{i}$ responses, reported for both cultured ADPKD\textsuperscript{24} and ARPKD\textsuperscript{25,26}

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cells, point to a possible fundamental role of disrupted \([Ca^{2+}]_i\) signaling in cystogenesis. The central cilia and cilia-associated PC1 and PC2 were proposed to mediate flow-induced cellular responses.\(^{19,27}\) However, homomeric PC2 channels are not mechanosensitive and fail to increase \([Ca^{2+}]_i\) in response to flow and hypotonicity.\(^{28,29}\) Furthermore, intercalated cells, which lack primary cilia, respond to flow changes with comparable increases in \([Ca^{2+}]_i\), as observed in principal cells, which have primary cilia.\(^{16,30}\) Therefore, additional mechanisms conferring mechanosensitivity to the CD cells need to be considered.

Transient receptor potential (TRP) channels are known to participate in cellular responses to a variety of environmental stimuli, including thermosensation, chemosensation, and mechanical forces (reviewed in Song and Yuan\(^{31}\)). Several TRP channels, including TRPC3, TRPC6, and TRPV4, can be detected in the native CD cells and CD-originated cultured lines.\(^{19,32–34}\) Among these channels, TRPV4 has routinely been shown to be activated by mechanical stimuli.\(^{34–38}\) Indeed, we documented that endogenous TRPV4 in M-1 CD cells is stimulated by increases in flow, a response that is abolished by TRPV4 small interfering RNA knockdown.\(^{34,38}\) We further demonstrated a lack of flow-mediated \([Ca^{2+}]_i\) elevations in CD from TRPV4−/− mice.\(^{30}\) Consistently, flow-mediated \([Ca^{2+}]_i\)-dependent K⁺ secretion in the CD is disrupted in TRPV4 knockout animals.\(^{39}\) TRPV4 directly interacts with PC2 to form mechanosensitive heteromeric complexes.\(^{28,29}\) The fact that PC2 interacts with both PC1\(^{10–12}\) and fibrocystin\(^10–12\) suggests that TRPV4 could be an essential part of this mechanotransducing sensory complex.

Current PKD management is directed toward pharmacologic interference with abnormal signaling pathways causing exaggerated cell proliferation, dedifferentiation, apoptosis, and cyst growth.\(^{41}\) Specifically, PKD is associated with elevated circulating vasopressin levels, increased basal cellular cAMP levels, and strong upregulation of cAMP-dependent fluid secretion and proliferation.\(^{42,43}\) V2 antagonism greatly diminishes disease progression in rodent models of both ADPKD and ARPKD.\(^{42,44}\) Elevated cAMP levels might be directly related to the reduced \([Ca^{2+}]_i\), possibly due to impaired ability to sense changes in flow.\(^{42,44,45}\) This raises the possibility that manipulation with the mechanosensitivity in the CD along the TRPV4 axis modulates \([Ca^{2+}]_i\) signalization and, in turn, renal cystogenesis.

In this study, we developed a new approach to isolate native CD-derived cyst monolayers and nondilated CD from a rat model of ARPKD to thoroughly investigate how functional TRPV4 status determines the development and growth of renal cysts. We found that the disease leads to disruption of mechanosensitive \([Ca^{2+}]_i\) signaling and impaired TRPV4 activity specifically in CD cysts but not in nondilated CD cysts. Long-term pharmacologic potentiation of TRPV4 activity gradually restores mechanosensitivity in cyst cells and greatly blunts renal ARPKD progression. From a global prospective, this study establishes a temporal link between disruption of TRPV4-based mechanosensitivity in the CD and cystogenesis. This also suggests pharmacologic potential of targeting TRPV4 activity as a treatment strategy in retarding development of ARPKD.

RESULTS

Accumulating evidence points to impaired flow-dependent \([Ca^{2+}]_i\) signalization in cultured ARPKD cells.\(^{25,26}\) To assess mechanosensitive properties of native cyst cells, we developed and implemented mechanical isolation of CD-derived cyst monolayers from 2-month-old ARPKD PCK453 rats. This novel technique enables real-time monitoring changes in \([Ca^{2+}]_i\) at the single cell level in native tissue in response to mechanical and pharmacologic inputs (Figure 1). CD origin of the isolated cyst fragments was confirmed using staining with AQP2, as discussed in the Concise Methods. We used freshly isolated split-opened CDs from age-matched Sprague-Dawley (S/D) rats as controls. An abrupt 10× elevation in flow (producing shear stress of approximately 3 dyn/cm², as discussed in the Concise Methods) on the apical surface caused a rapid and sustained increase (at least for 10 minutes) (Supplemental Figure 1) in \([Ca^{2+}]_i\) in control CD cells, whereas it had a minimal effect on cyst cells (Figure 2A). Importantly, basal \([Ca^{2+}]_i\) levels are prominently decreased in ARPKD cells.
We demonstrated that genetic deletion of the Ca\(^{2+}\)-permeable TRPV4 channel abolished flow-mediated [Ca\(^{2+}\)]\(_i\) elevations in CD cells of mice.\(^{30}\) To probe whether TRPV4 is also critical for mechanosensitive properties in rat CD cells, we took advantage of a potent and selective TRPV4 antagonist, HC067047. Acute treatment with HC067047 (4 \(\mu M\)) nearly abolished elevations in [Ca\(^{2+}\)]\(_i\) to the mechanical stimulation (Figure 2B). Pretreatment with a less selective TRPV4 antagonist, Ruthenium red (2 \(\mu M\)) yielded similar results (Figure 2C). Inhibition of TRPV4 with HC067047 also significantly decreased basal [Ca\(^{2+}\)]\(_i\) levels (Figure 2B). These results suggest a key role of TRPV4 in mediating flow-dependent Ca\(^{2+}\) responses and shaping basal [Ca\(^{2+}\)]\(_i\) in rat CD cells. Importantly, pharmacologic inhibition of TRPV4 recapitulates the state of compromised [Ca\(^{2+}\)]\(_i\) signalization observed in cyst cells (Figure 2, A and C).

Lack of flow-mediated Ca\(^{2+}\) responses and diminished basal [Ca\(^{2+}\)]\(_i\) levels in ARPKD cyst cells indicate TRPV4 dysfunction. Indeed, stimulation of TRPV4 with a highly selective activator, GSK1016790A (30 nM), caused [Ca\(^{2+}\)]\(_i\) elevations in control CDs that were four times greater than in cyst cells (Figure 2D). GSK1016790A had no effect when TRPV4 was inhibited with Ruthenium red (Supplemental Figure 2). We concluded that functional activity of TRPV4 is drastically diminished in ARPKD cyst cells contributing to the impaired flow-mediated Ca\(^{2+}\) signalization.

TRPV4 dysfunction in cyst monolayers may result from decreased levels of the protein. Figure 3A shows a representative Western blot monitoring TRPV4 expression in whole kidney homogenates from S/D and PCK453 rats. As expected, TRPV4 antibodies recognized both nonglycosylated (lower) and glycosylated (upper) bands around 95 kD in S/D rats. Renal TRPV4 levels were modestly but significantly reduced in PCK453 rats. Strikingly, the glycosylated form of TRPV4 was virtually absent in these animals (Figure 3). We concluded that post-translational modification of TRPV4 is compromised during ARPKD.

We next probed subcellular TRPV4 distribution in control split-opened CDs from S/D rats and CD-derived cyst monolayers from PCK453 rats with immunofluorescence. TRPV4 is distributed in subapical cytosolic compartments and at the apical plasma membrane, with stronger expression in AQP2-positive cells (Figure 4A; see Supplemental Figure 3 for a lower magnification). Surprisingly, we detected a striking shift of the TRPV4-reporting fluorescent signal toward the apical membrane in CD-derived cyst monolayers (Figure 4B; see Supplemental Figure 3 for a lower magnification). We also observed AQP2 targeting to the apical side, which is consistent with reported upregulation of cAMP levels in ARPKD cells.\(^{42,46}\) For quantitative estimation, we utilized line-scan analysis (Figure 4C) of TRPV4-reporting fluorescent signal distribution along the z-axis in cross-sections of three-dimensional stacks similar to that shown in Figure 4, A and B. Cyst cells clearly exhibit a leftward narrowing of the bell-shape distribution of TRPV4 along the z-axis toward the apical plasma membrane. As summarized in Figure 4D, the half-width was significantly reduced from 2.63 ± 0.12 \(\mu M\) in the control to 1.89 ± 0.03 \(\mu M\) in CD and cyst cells, respectively.

To exclude the possibility that isolation of CD-derived cysts might potentially cause subcellular TRPV4 redistribution,
we monitored TRPV4 localization in kidney sections from S/D and PCK453 rats (Figure 5). TRPV4 was preferentially localized to the AQP2-positive distal nephrons and cysts (respective top rows in Figure 5). Higher-resolution images of the same respective kidney sections revealed that TRPV4- and AQP2-reporting fluorescent signals are restricted to the apical/subapical domain of cyst cells (Figure 5B, middle row). This is consistent with the results obtained in freshly isolated cyst fragments (Figure 4B). In contrast, we detected diffuse subcellular distribution of TRPV4 and AQP2 in nondilated CDs (Figure 5B, bottom row). These expression patterns are clearly different from observed in cyst cells and are reminiscent of those observed in control CD cells from S/D rats (Figure 5A).

Different subcellular TRPV4 distribution in nondilated CD cells and cyst cells might indicate altered TRPV4 functionality. Thus, we assessed mechanosensitive Ca\textsuperscript{2+} signalization and functional TRPV4 status in nondilated CDs from PCK453 rats. We detected prominent [Ca\textsuperscript{2+}]\textsubscript{i} response to elevations in luminal flow and unchanged basol [Ca\textsuperscript{2+}]\textsubscript{v} values (Figure 6A and B). Activation of TRPV4 with GSK1016790A also produced a similar rise in [Ca\textsuperscript{2+}]\textsubscript{i} in control CD cells from Sprague-Dawley rats (Figure 6C and Supplemental Figure 4). Consistently, subcellular TRPV4 and AQP2 distribution in freshly isolated nondilated CDs was not different from that in control CDs as visualized with immunofluorescence (Figure 6D). Therefore, the results in Figure 6 strongly argue for unchanged mechanosensitivity in nondilated CDs.

Substantial experimental evidence supports a critical role of paracrine ATP release in facilitating [Ca\textsuperscript{2+}]\textsubscript{i} responses in the CD.\textsuperscript{21} Development of ARPKD results in a virtually closed environment inside renal cysts, creating favorable conditions for amplification of the purinergic signal.\textsuperscript{14,47,48} Thus, we next characterized the functional status of purinergic cascade in CD-derived cyst monolayers and nondilated CDs from PCK453 rats. Exogenous ATP (10 μM) induced a transient [Ca\textsuperscript{2+}]\textsubscript{i} peak followed by a sustained plateau (Figure 7). The magnitude of the initial peak was indeed markedly increased in cyst cells compared with the control CD cells from S/D rats (Figure 7). However, the plateau phase was nearly absent in ARPKD cells. In nondilated CDs from PCK453 rats, ATP produced Ca\textsuperscript{2+} response undistinguishable from that observed in control CDs (Figure 7A). We recently identified a dominant role of TRPV4 in generating ATP-induced Ca\textsuperscript{2+} plateau in murine CD cells.\textsuperscript{18} The absence of the plateau phase in cyst cells supports the concept of TRPV4 dysfunction in ARPKD. These results suggest malfunctioning of purinergic signaling in CD-derived cysts but not in nondilated CDs.

Our results point to a severe TRPV4 dysfunction in CD-derived cysts of ARPKD. Physiologic stimuli, including flow-mediated shear stress and activation of purinergic cascade, fail to effectively stimulate TRPV4 to elicit proper mechanosensitive response. Moreover, impaired TRPV4 functional status contributes to the decreased resting [Ca\textsuperscript{2+}]\textsubscript{i} levels (Figures 2 and 7). However, TRPV4 can still be activated to some extent by strong pharmacologic inputs, such as GSK1016790A (Figure 2D). We hypothesized that augmentation of the mechanosensitivity on the TRPV4 axis will interfere with ARPKD progression. Indeed, a recent report provided initial evidence that prolonged injection of GSK1016790A to PCK453 rats decreases renal cystic area.\textsuperscript{49} Therefore, we next tested if GSK1016790A treatment interferes with ARPKD progression by restoring mechanosensitive properties of the CD-derived cyst cells.

Systemic administration of GSK1016790A (3 μg/kg body weight per day) for 1 and 2 months (Figure 8, A and B) to 1-month-old PCK453 rats diminished renal cyst development and growth compared with the control vehicle-treated group. Specifically, we detected a significant reduction of relative cyst area (Figure 8C; also see low-powered section slices in Supplemental Figure 5) at both time points. We still observed the disease progression in GSK1016790A-treated animals between 1 and 2 months of treatment, although at a slower rate compared with that in vehicle-treated animals. The kidney/total body index was also reduced by GSK1016790A treatment (Figure 8D). In contrast, GSK1016790A (1 month) had no significant effect on kidney/total body index in control S/D rats (0.89 ± 0.08 versus 0.86 ± 0.04 for treated and untreated animals, respectively).

We next examined the ramifications of systemic GSK1016790A administration on [Ca\textsuperscript{2+}]\textsubscript{i} signaling in cells from residual CD-derived cysts. We detected gradual restoration of the flow-mediated [Ca\textsuperscript{2+}]\textsubscript{i} responses and return of resting [Ca\textsuperscript{2+}]\textsubscript{i} levels to control values after 1 and 2 months...
of treatment, respectively (Figure 9A). We also found progressive augmentation of \([\text{Ca}^{2+}]_i\) response to an acute application of GSK1016790A (Figure 9B). This suggests enhancement of functional TRPV4 status in GSK1016790A-treated animals, likely due to TRPV4 sensitization. Consistently, we detected appearance of the ATP-induced Ca\(^{2+}\) plateau (Figures 9C).

Overall, we concluded that prolonged systemic administration of GSK1016790A recovers the bulk of mechanosensitive \([\text{Ca}^{2+}]_i\) signal in GiN-derived cyst cells.

We next subjected CD-derived cyst monolayers of GSK1016790A-treated PCK453 rats to immunofluorescence. Systemic GSK1016790A treatment for 1 month greatly reversed the apical TRPV4 translocation observed in cysts from untreated animals of the same age (Figure 10; see Figure 4B for comparison). Specifically, we detected a shift of the TRPV4-reporting fluorescent signal from apical membrane to cytosol. We also detected redistribution of AQP2 from the apical membrane, indicating a possible decrease in cellular cAMP levels. Treatment with GSK1016790A for 2 months yielded similar results (Supplemental Figure 6). However, GSK1016790A fails to restore TRPV4 glycosylation in the kidney of PCK453 rats (Supplemental Figure 7). Total TRPV4 levels were also not altered in GSK1016790A-treated PCK453 rats (87\%\(\pm\)3\% versus 100\%\(\pm\)2\% for treated and untreated rats, respectively). We concluded that TRPV4 distribution in cyst cells of GSK1016790A-treated animals mainly recapitulates TRPV4 localization in CD cells of control Sprague-Dawley rats (Figure 4A). This occurs independently of significant changes in renal TRPV4 levels and TRPV4 glycosylation status.

**DISCUSSION**

This study provides two major advances in our understanding of the relation between mechanosensitivity and cystogenesis in the CD. First, we demonstrate that the defect in the fibrocystin gene in PCK453 rats does not lead to immediate disruption of flow sensitivity and compromised TRPV4 activity in epithelial cells of nondilated CDs. Thus, the intact mechanosensitive \([\text{Ca}^{2+}]_i\) signaling might have a permissive role for normal tubular function and its disruption specifically occurs before, or as an early event in, cyst development. Second, we provide evidence that restoration of mechanosensitivity and TRPV4 function in cyst cells drastically diminishes ARPKD progression. This study provides two major advances in our understanding of the relation between mechanosensitivity and cystogenesis in the CD. First, we demonstrate that the defect in the fibrocystin gene in PCK453 rats does not lead to immediate disruption of flow sensitivity and compromised TRPV4 activity in epithelial cells of nondilated CDs. Thus, the intact mechanosensitive \([\text{Ca}^{2+}]_i\) signaling might have a permissive role for normal tubular function and its disruption specifically occurs before, or as an early event in, cyst development. Second, we provide evidence that restoration of mechanosensitivity and TRPV4 function in cyst cells drastically diminishes ARPKD progression. This strongly supports the concept that flow-dependent \([\text{Ca}^{2+}]_i\) signaling plays an important modulatory role in renal cyst development. However, it does not seem that the disruption of mechanosensitivity per se initiates cystogenesis because TRPV4−/− mice\(^{50}\) and zebrafish\(^{29}\) do not have renal cysts.

We consistently observed diminished basal levels of \([\text{Ca}^{2+}]_i\) in native CD-derived cyst cells of ARPKD rats (Figures 2 and
BASIC RESEARCH

Increased [Ca\textsuperscript{2+}]i values during PKD may contribute to the development of familial digital arthropathy-brachydactyly. \textsuperscript{54} We demonstrate that TRPV4 functionality determines the status of flow-mediated [Ca\textsuperscript{2+}]i signaling in the CD. Diminished TRPV4 activity is associated with the lack of flow-sensitive [Ca\textsuperscript{2+}]i elevations in freshly isolated CD-derived cyst monolayers, whereas TRPV4 sensitization after systemic GSK1016790A treatment rejuvenates [Ca\textsuperscript{2+}]i responses to elevated flow in cyst cells. This enables us to correlate changes in mechanosensitivity at the cellular level with systemic progression of ARPKD. We found that TRPV4 dysfunction in cyst cells is not likely associated with decreased TRPV4 protein levels but rather with impaired ability of the channel to be activated (Figure 2D). Our results point to dramatically reduced glycosylation of TRPV4 in PCK453 rats (Figure 3). Interestingly, a recent study demonstrated that mutations in TRPV4 associated with diminished glycosylation abolished responses to environmental stimuli (such as hypotonicity) and decreased responses to GSK1016790A.\textsuperscript{54} Such impaired post-translational TRPV4 processing has been implicated in the development of familial digital arthropathy-brachydactyly.\textsuperscript{54}

Interestingly, the beneficial effects of TRPV4 activation during ARPKD might not be limited to the kidney. A recent study identifies an antiproliferative role for TRPV4 in cholangiocytes from ARPKD rats and TRPV4 activation has a tendency to decrease liver cysts.\textsuperscript{49} This study also provides initial observation that injection of GSK1016790A diminishes renal cyst area. In our study, we manage, for the first time, isolation of monolayers of CD-derived cysts to correlate TRPV4 function, mechanosensitivity of native CD cells, and renal cystogenesis at the timescale of ARPKD progression using \textit{per os} administration of the TRPV4 activator, GSK1016790A. However, it should be noted that excessive activation of TRPV4 by an acute administration of high doses of GSK1016790A increases permeability of the pulmonary microvascular barrier leading to circulatory collapse.\textsuperscript{55} In contrast, gradual ingestion of small doses of the drug for up to 2 months, as used in this study, is well tolerated and does not produce any detrimental side effects.

This investigation also addresses the purinergic aspect of mechanosensitivity and its relation to TRPV4 function. ATP is due to augmentation of basal [Ca\textsuperscript{2+}]i (Figures 4 and 10). This explains, at least in part, the anticystogenic effect of GSK1016790A (Figure 8 and Supplemental Figure 5). The observation that systemic GSK1016790A treatment enhances TRPV4 activity requires a comment. Unlike many other Ca\textsuperscript{2+}-permeable channels, TRPV4 can be activated by modest elevations in [Ca\textsuperscript{2+}]i via mechanisms involving direct binding of calmodulin to the channel.\textsuperscript{52} Initial small activation of TRPV4 by GSK1016790A provides a source for such increases in [Ca\textsuperscript{2+}]i, leading to sustained potentiation of TRPV4 activity. On a longer timescale, elevations of [Ca\textsuperscript{2+}]i will also promote phosphorylation of TRPV4 by Ca\textsuperscript{2+}-dependent protein kinase C isoforms further activating the channel.\textsuperscript{53} In contrast, an excessive increase in [Ca\textsuperscript{2+}]i will lead to inhibition/desensitization of TRPV4 via a negative feedback mechanism\textsuperscript{52} protecting cells from the cytotoxic effects of elevated [Ca\textsuperscript{2+}]i.

We demonstrate that TRPV4 functionality determines the status of flow-mediated [Ca\textsuperscript{2+}]i signaling in the CD. Diminished TRPV4 activity is associated with the lack of flow-sensitive [Ca\textsuperscript{2+}]i elevations in freshly isolated CD-derived cyst monolayers, whereas TRPV4 sensitization after systemic GSK1016790A treatment rejuvenates [Ca\textsuperscript{2+}]i responses to elevated flow in cyst cells. This enables us to correlate changes in mechanosensitivity at the cellular level with systemic progression of ARPKD. We found that TRPV4 dysfunction in cyst cells is not likely associated with decreased TRPV4 protein levels but rather with impaired ability of the channel to be activated (Figure 2D). Our results point to dramatically reduced glycosylation of TRPV4 in PCK453 rats (Figure 3). Interestingly, a recent study demonstrated that mutations in TRPV4 associated with diminished glycosylation abolished responses to environmental stimuli (such as hypotonicity) and decreased responses to GSK1016790A.\textsuperscript{54} Such impaired post-translational TRPV4 processing has been implicated in the development of familial digital arthropathy-brachydactyly.\textsuperscript{54}

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7). Similar decreases in resting [Ca\textsuperscript{2+}]i values were reported for cultured human ADPKD and ARPKD cells.\textsuperscript{45} We further provide strong experimental support to the conception that TRPV4 activity is a critical determinant of basal [Ca\textsuperscript{2+}]i. Inhibition of TRPV4 with HC067047 decreases basal [Ca\textsuperscript{2+}]i, recapitulating the values observed in cyst cells that have marked TRPV4 dysfunction (Figure 2, B–D). Restoration of functional TRPV4 status in GSK1016790A-treated animals brings [Ca\textsuperscript{2+}]i back to the levels detected in normal CD cells. Decreased [Ca\textsuperscript{2+}]i values during PKD may contribute to the abnormally elevated cAMP signaling.\textsuperscript{42,44,45} This is known to stimulate Cl\textsuperscript{−} secretion inside the cyst lumen and to promote proliferation of cyst cells.\textsuperscript{51} By monitoring changes in AQP2 subcellular distribution patterns, we provide evidence that pharmacologic activation of TRPV4 blunts cAMP levels likely

![Figure 5. Subcellular TRPV4 localization is different in cysts and nondilated CDs in PCK453 rats. (A) Immunofluorescence of a kidney section from Sprague-Dawley (S/D) rat. A representative low-magnification 10-μm-thick kidney section (top row) shows staining patterns for TRPV4 (pseudocolor green), AQP2 (pseudocolor red), and the combined image. The area defined by a rectangle is shown below with a higher magnification. (B) Immunofluorescence of kidney section from a PCK453 rat. A representative low-magnification 10-μm-thick kidney section (top row) shows staining patterns for TRPV4 (pseudocolor green), AQP2 (pseudocolor red), and the combined image. Areas defined by rectangles and containing a cyst (middle) and a nondilated CD (bottom) are shown below with a higher magnification.](image-url)
constitutively released from CD cells and this process is strongly potentiated by mechanical stimuli. It becomes increasingly appreciated that paracrine purinergic signaling is an important facilitator of mechanosensitivity in the CD. Indeed, we demonstrated that genetic deletion of the major purinergic receptor, P2Y2, markedly blunts cellular responses to elevated flow and hypotonicity. Remodeling kidney tissue during development of PKD creates a virtually closed environment inside cysts that can trap ATP. Several reports suggest a marked augmentation of purinergic cascade and an increased ATP concentration inside the cyst lumen in PKD. This potentiation was shown to have detrimental renal effects by promoting Cl− secretion and cyst growth. Pharmacologic inhibition of purinergic signaling could be a putative pharmacologic target to manage PKD. Despite the fact that paracrine ATP release might be an important component of mechanosensitivity in normal CDs, this paradoxical upregulation of purinergic signaling in PKD pathology does not improve mechanosensitive properties of cyst cells. We found that the initial transient ATP-induced [Ca2+]i peak was elevated in CD-derived cyst cells. In contrast, the sustained plateau is nearly absent in cyst cells. This suggests that the chronic potentiation of purinergic cascade in ARPKD will have a minimal effect on [Ca2+]i levels. We identified a dominant role of TRPV4 in the generation of the ATP-induced [Ca2+]i plateau and found that genetic deletion of TRPV4 nearly abolishes the sustained elevation of [Ca2+]i in response to ATP. The absence of the plateau in cyst cells is consistent with TRPV4
dysfunction and impaired mechanical perception and vice versa, restoration of TRPV4 functionality in GSK1016790A-treated animals causes appearance of ATP-induced Ca\(^{2+}\) plateau (Figure 9C). Therefore, our results allow consolidating the upregulation of purinergic signaling inside cyst lumen and the loss of mechanosensitivity in ARPKD.

In summary, this study directly demonstrates the relation between TRPV4 activity, mechanosensitivity in CD cells, and renal cyst development in ARPKD. By identifying TRPV4 dysfunction as an underlying cause for disrupted flow perception by the cyst cells, we provide evidence that augmentation of TRPV4 status limits renal ARPKD manifestations. It is also tempting to speculate that pharmacologic targeting of signaling pathways that possess stimulatory actions on TRPV4 would yield similar results. Finally, although our investigation focuses exclusively on ARPKD, we anticipate that systemic enhancement of TRPV4 function will also be beneficial for treatment of ADPKD because both pathologies are associated with disrupted mechanosensitive [Ca\(^{2+}\)]\(_i\) signalization.

CONCISE METHODS

Materials and Animals
All chemicals and materials were from Sigma (St. Louis, MO), VWR (Radnor, PA), and Tocris (Ellisville, MO) unless noted otherwise and were of reagent grade. Animal use and welfare adhered to the National Institutes of Health Guide for the Care and Use of Laboratory Animals following a protocol reviewed and approved by the Institutional Laboratory Animal Care and Use Committee of the University of Texas Health Science Center at Houston. For experiments, 1- to 3-month-old control Sprague-Dawley (S/D) rats and ARPKD PCK453 rats (Charles River Laboratories, Wilmington, MA) were used. Animals were maintained on standard rodent regimen (Purina #5001) and had free access to tap water. For some experiments, 1-month-old PCK453 rats received either tap water or tap water containing GSK1016790A (3 μg/kg body weight per day) for either 1 or 2 months. This concentration is 100 times lower than the reported lethal dose for an acute intravenous injection of the substance. We did not observe any noticeable morbidity in the GSK1016790A-treated groups during the time course of the drug administration. After the end of the treatment, rats were sacrificed by CO\(_2\) administration followed by cervical dislocation and the kidneys were immediately removed and weighted.

Tissue Isolation
The procedure for isolation of the CDs from Sprague-Dawley rats and nondilated CDs from PCK453 rats suitable for Ca\(^{2+}\) imaging and immunofluorescence closely follows our previously reported rat and mouse protocols. Kidneys were cut into thin slices (<1 mm), with slices placed into ice-cold physiologic saline solution buffered with HEPES (pH 7.4). The CD was visually identified by its morphologic features (pale color, coarse surface, and, in some cases, bifurcations) and was mechanically isolated from kidney slices by microdissection using watchmaker forceps under a stereomicroscope. Isolated CDs were attached to 5x3 mm coverglass coated with poly-L-lysine. A coverglass containing CD was placed in a perfusion chamber mounted on an inverted Nikon Eclipse Ti microscope and perfused with room temperature HEPES-buffered (pH 7.4) saline solution. CDs were split-opened with two sharpened micropipettes, controlled with different micromanipulators, to gain access to the apical membrane. The nephrons were used within 1–2 hours of isolation.

Using a similar approach, CD-derived cyst monolayers were gently teased from open cyst cavities in kidney slices from PCK453 rats using watchmaker forceps under a stereomicroscope. ARPKD cyst
monolayers were further mechanically separated from surrounding tissue/nephrons and attached with the basal side to 5 μm coverglass coated with poly-L-lysine.

**[Ca^{2+}]** Measurements

Unless otherwise noted, 2-month-old Sprague-Dawley and PCK453 rats were used for experiments. Intracellular calcium levels were measured in individual cells of the split-opened CDs and CD-derived cyst monolayers using Fura-2 fluorescence radiometric imaging as described previously.\(^{18,34,38}\) Split-opened CDs and cysts fragments were loaded with Fura-2 by incubation with 2 μM Fura-2/AM in a bath solution for 45 minutes at room temperature. Subsequently, the tissue samples were washed and incubated for an additional 10–15 minutes before experimentation. The CDs and cyst monolayers were then placed in an open-top imaging study chamber (Warner RC-10) with a bottom coverslip viewing window and the chamber attached to the microscope stage of an InCa Imaging Workstation (Intracellular Imaging Inc, Cincinnati, OH). Cells were imaged with a 20x Nikon Super Fluor objective and regions of interest drawn for individual cells. The Fura-2 fluorescence intensity ratio was determined by excitation (an average for approximately 300 ms) at 340 nm and 380 nm and calculating the ratio of the emission intensities at 511 nm in the usual manner every 5 seconds. We observed no significant Fura-2 bleaching and minimal Fura-2 leakage at both wavelengths during experiments. The changes in the ratio are converted to intracellular Ca\(^{2+}\) concentrations using the calibration methods as previously described\(^{18,59,62}\). At least three individual CDs or cysts from three rats were used for each experimental set.

**Western Blot Analyses**

Immediately after dissection, kidneys were placed on ice, decapsulated, and homogenized in 3 volumes of ice-cold hypotonic lysis buffer containing 50 mM Tris, 1% Triton X-100, 5 mM EDTA (pH 7.4) supplemented with 1 mM PMSF and 2 mg/ml protease inhibitor cocktail (Complete Mini; Roche Diagnostics, Indianapolis, IN). Protein concentration was determined with a Bradford assay using IgG as a standard. The samples were diluted with hypotonic lysis buffer, denatured, and reduced in Laemmli buffer supplemented with 10 mM of dithiothreitol at +75°C for 10 minutes to obtain the final protein concentration of 4 mg/ml. The samples (10 μg/lane) were separated on 9% polyacrylamide gels at 150 V for 1.75 hours and transferred to nitrocellulose membrane for 1.5 hours at 100 V. Subsequently, the nitrocellulose membrane was incubated with anti-TRPV4 (1:500; Alomone Labs, Jerusalem, Israel) and anti-actin (1:1000; Abcam, Cambridge, UK) primary antibodies for 2 hours at room temperature. We previously tested specificity of the TRPV4 antibodies in expression systems, cultured M1 cells,\(^{34}\) and TRPV4\(^{−/−}\) mice.\(^{30}\) Upon washout (three times for 10 minutes in TBS-Tween), the membrane was incubated with peroxidase-conjugated goat anti-rabbit secondary antibodies (1:30,000; Bio-Rad, Hercules, CA) for 1 hour at room temperature. Blots were quantified using ImageJ 1.47 software (National Institutes of Health, Bethesda, MD). The intensities for TRPV4 protein bands were normalized to the intensities of the corresponding actin bands, used as a loading control. All experiments were repeated three times.

**Immunofluorescence and Renal Histology**

Kidneys from PCK453 rats were briefly fixed in situ via intracardiac perfusion and subjected to standard immunohistochemical procedures as done before.\(^{30}\) Briefly, animals were anesthetized by
isoflurane inhalation and kidneys prepared by intracardiac perfusion by a fixation solution (20 ml, 4% paraformaldehyde and 0.1 M cacodylate buffer in ice-cold PBS, pH 7.4). Kidneys were removed and placed in 4% paraformaldehyde overnight, and then mounted in tissue freezing medium (Tissue Tek; Sakura Finetek Inc., Torrance, CA) and frozen at −30°C. Sagittal sections (10 μm thick) were obtained, using an OTF 5000 cryostat (Bright Instrument, Huntingdon, UK). Sections were allowed to warm to room temperature and washed in PBS. Sections were then blocked with 1% donkey serum, and incubated overnight at 4°C with primary antibody: anti-TRPV4 (1:500; Alomone Labs) and anti-AQP2 tagged with ATTO 550 (1:200; Alomone). Sections were subsequently washed and incubated at room temperature for 3 hours with secondary antibody Cy2 anti-rabbit (1:500; Jackson ImmunoResearch, West Grove, PA). The tissue was then mounted in ProLong Gold antifade reagent with 4′,6-diamidino-2-phenylindole (Invitrogen, Carlsbad, CA).

For histologic assessment of renal ARPKD manifestation, kidney sections (30 μM) from untreated and GSK106790A-treated PCK453 rats were obtained as described above. Sections were stained with hematoxylin and eosin with standard procedures and used to quantify cystic area. Image analysis was performed using NIS elements 4.00 software (Nikon, Nakville, NY). At least four different sections from three different animals were used for the assessment.

Freshly isolated CD-derived cyst monolayers and split-opened CDs were fixed with 4% paraformaldehyde in PBS (pH 7.4) for 20 minutes at room temperature (RT). After fixation the samples were permeabilized by addition of 0.1% Triton in PBS for 5 minutes and washed in PBS three times for 5 minutes. Nonspecific staining was blocked with 10% normal goat serum (NGS; Jackson ImmunoResearch) in PBS for 30 minutes at RT. After washing with PBS (three
times for 5 minutes) the samples were incubated for 3 hours at RT in dark with the mix of anti-TRPV4 (1:300; Alomone Labs), anti-AQP2 tagged with ATTO 550 (1:100; Alomone Labs) in 1% serum + 0.1% Triton in PBS. Subsequently, samples were washed three times with PBS and incubated for 1.5 hours at RT in dark with goat anti-rabbit IgG labeled with Alexa Fluor 488 (1:400 dilution; Invitrogen) in 1% NGS + 0.1% Triton in PBS. After washing with PBS (three times for 5 min) the samples were stained with 4',6-diamidino-2-phenylindole (300 nM concentration; Calbiochem, San Diego, CA) to visualize nuclei. Subsequently the samples were dehydrated, and mounted with permanent mounting media (Thermo Scientific, Pittsburg, PA). Labeled tissue samples were examined with an inverted Nikon Eclipse Ti fluorescent microscope using a 40× Plan-Fluor oil-immersion (1.3 NA) objective. Samples were excited with 405, 488, and 561 nm laser diodes and emission captured with a 16-bit Cool SNAP HQ2 camera (Photometrics, Tucson, AZ) interfaced to a PC running NIS 4.0 elements software.

**Solutions**

Typical bath solution was as follows: 150 mM NaCl, 5 mM KCl, 1 mM CaCl₂, 2 mM MgCl₂, 5 mM glucose, and 10 mM HEPES (pH 7.4). All reagents were applied by perfusing the experimental chamber at 1.5 ml/min. To test the effect of elevated shear stress, perfusion was instantly increased from 1.5 ml/min (approximately 80 mm H₂O) to 15 ml/min (approximately 450 mm H₂O). Using a parallel plate chamber, we recently estimated that this maneuver produces shear stress of approximately 3 dyn/cm². This value fits well within the physiologic range of shear stress present in the rat and mouse CD as assessed previously. Statistical Analyses

All summarized data are reported as mean ± SEM. Data were compared using the t test or an one-way ANOVA as appropriate. P ≤ 0.05 was considered significant.

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**DISCLOSURES**

None.

**REFERENCES**


