Curcumin inhibits cystogenesis by simultaneous interference of multiple signaling pathways: in vivo evidence from a Pkd1-deletion model

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Leonhard WN, van der Wal A, Novalic Z, Kunnen SJ, Gansevoort RT, Breuning MH, de Heer E, Peters DJ. Curcumin inhibits cystogenesis by simultaneous interference of multiple signaling pathways: in vivo evidence from a Pkd1-deletion model. Am J Physiol Renal Physiol 300: F1193–F1202, 2011. First published February 23, 2011; doi:10.1152/ajprenal.00419.2010.—Autosomal dominant polycystic kidney disease (ADPKD) caused by mutations in either the PKD1 or PKD2 gene is a major cause of end-stage renal failure. A number of compounds targeting specific signaling pathways were able to inhibit cystogenesis in rodent models and are currently being tested in clinical trials. However, given the complex signaling in ADPKD, an ideal therapy would likely have to comprise several pathways at once. Therefore, multitarget compounds may provide promising therapeutic interventions for the treatment of ADPKD. To test this hypothesis, we treated Pkd1-deletion mice with diferuloylmethane (curcumin), a compound without appreciable side effects and known to modulate several pathways that are also altered in ADPKD, e.g., mammalian target of rapamycin (mTOR) and Wnt signaling. After conditional inactivation of Pkd1, mTOR signaling was indeed elevated in cystic kidneys. Interestingly, also activation of signal transducers and activator of transcription 3 (STAT3) strongly correlated with cyst progression. Both pathways were effectively inhibited in vitro by curcumin. Importantly, Pkd1-deletion mice that were treated with curcumin and killed at an early stage of PKD displayed improved renal histology and reduced STAT3 activation, proliferation index, cystic index, and kidney weight/body weight ratios. In addition, renal failure was significantly postponed in mice with severe PKD. These data suggest that multitarget compounds hold promising potential for safe and effective treatment of ADPKD.

The polyphenol diferuloylmethane (curcumin) is a yellow spice derived from the rhizome of the plant Curcuma Longa (26). Its anti-oxidant, anti-inflammatory, and anti-proliferative properties initiated many studies for the treatment of several diseases. Curcumin has been shown to be safe and effective in treating several inflammatory and malignant diseases in animal models and, although preliminary, also in a number of early phase clinical trials (3, 12, 16, 22, 24, 58, 59). In addition, promising developments regarding curcumin analogs and other multitarget compounds will likely improve their utility in clinical settings (1, 20, 21, 25, 36, 39, 40, 57). The complex mechanisms by which curcumin exerts its beneficial effects involve a wide variety of pathways (26, 58). Inhibition has been reported for the transcription factors activator protein-1, nuclear factor-κB (NF-κB), Wnt/β-catenin signaling, TNF-α, MAPKs, early growth response gene-1, hypoxia inducible factor-1, notch-1, and also the mTOR-regulated signaling (4, 5, 2300 RC Leiden, The Netherlands (e-mail: d.j.m.peters@lumc.nl).
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23, 26, 32, 58). Many of these pathways are altered in ADPKD (27, 31, 35, 55, 56, 61, 69). In addition, curcumin acts as a free radical scavenger and may further attenuate several forms of tubular injury by induction of NF-E2-related factor-2 (Nrf2) and subsequent increase of several cytoprotective enzymes (2). The ability of curcumin to inhibit signal transducer and activator of transcription 3 (STAT3) may also be relevant to ADPKD since we show here that STAT3 is activated in iKsp-Pkd1<sup>del</sup> mice (10, 26).

The similarities between altered signaling pathways in ADPKD and pathways that are modulated by curcumin and the ongoing developments regarding curcumin analogs and compounds with similar modes of action prompted us to investigate whether this multitarget drug might have therapeutic potential in the iKsp-Pkd1<sup>del</sup> model.

MATERIALS AND METHODS

**Mice.** Tamoxifen was administered orally at PN days 40–42 to inducible kidney-specific Pkd1-deletion mice (tam-KspCad-CreERT2:Pkd1<sup>lox/lox</sup>-CreER<sup>+</sup>) or in short iKsp-Pkd1<sup>del</sup> as described previously (29, 30). Curcumin (~70% purity) was obtained from Sigma (Zwijndrecht, The Netherlands) and RM3(P) pellets with 1% curcumin or without curcumin were prepared by Special Diets Services (Witham, UK). This dose and way of administration were chosen in analogy to a recent experimental study that studied the effectiveness of curcumin to inhibit prostate carcinoma (3). Blood samples and blood urea (BU) measurements were performed as described previously (18). Local animal experimental committee of the Leiden University Medical Center and the Commission Biotechnology in Animals of the Dutch Ministry of Agriculture approved the experiments performed.

**Immunohistochemistry.** After removal, kidneys were fixed O/N in buffered 4% formaldehyde solution and embedded in paraffin. Kidney sections (4 μm) were stained with standard hematoxylin and eosin or Periodic acid-Schiff (PAS). For immunohistochemical analysis, sections were deparaffinized and subjected to heat-mediated antigen retrieval [1 mM EDTA, pH 8.0 for anti-phospho-Stat3 (Tyr705) and 0.01 M citrate buffer, pH 6.0 for anti-phospho-ribosomal protein S6 (Ser240/244); Cell Signaling, no. 9145 and no. 2215S, respectively]. Sections were blocked with 0.1% H<sub>2</sub>O<sub>2</sub> for 20 min for endogenous peroxidase activity and preincubated for 1 h with 5% normal goat serum in 1% BSA in PBS. Next, the sections were incubated O/N with 0, 2.5, 5, or 10 μM curcumin (Sigma) in the absence of medium supplements. Then, the cells were incubated with the same curcumin concentrations in the presence of medium supplements for an additional 6 h. Protein lysates were prepared and analyzed by Western blot analysis as described below.

**Western blot and densitometric analysis.** Throughout the lysis procedure, 2 mM NaF, 1 mM Na<sub>2</sub>VO<sub>4</sub>, and protease inhibitor cocktail (Roche) were used to inhibit phosphatase and protease activity. Kidneys were homogenized in 600 μl RIPA without detergents (50 mM Tris·HCl, pH 7.4, 150 mM NaCl, 1 mM EDTA) using Magnafluor in animals of the Dutch Ministry of Agriculture approved the experiments performed.

Protein concentrations were determined using BCA protein assay (Thermo Fisher Scientific, Ettten-Leur, The Netherlands). Ten micrograms of cell lysates and 80–120 μg of kidney lysates were used for SDS-PAGE and immunoblotting. Membranes were blocked for 1 h in 25% sea block blocking buffer (Thermo Fisher Scientific/Pierce, no. 35727, Ettten-Leur, The Netherlands) in PBS and incubated O/N with anti-pSTAT3 (1:1,000), anti-p-pS6 (1:1,000), rabbit anti-Ki-67 (1:3,000; Nova Castra), rabbit anti-megalin (1:500; Pathology LUMC, Leiden, the Netherlands) (11), goat anti-Tamm-Horsfall protein (urodulin, 1:500; Organon Teknika-Cappel, Turnhout, Belgium), or rabbit anti-aquaporin-2 (1:4,000; Calbiochem, Amsterdam, The Netherlands) in 1% BSA in PBS. Following incubation with rabbit envision horseradish peroxidase (HRP; DakoCytonation, Glostrup, Denmark) or rabbit-anti-goat HRP (1:300, Dako), immune reactions were revealed using diaminobenzidine as a chromogen and counterstained with hematoxylin, dehydrated, and mounted.

**Cystic and proliferation index.** For each mouse, the percentage of cystic area relative to total tissue was determined by Image J software (public domain software, National Institutes of Health) on 10 x 40 magnified photographs throughout the cortex on kidney sections stained with PAS. For the proliferation index, sections were stained for Ki-67 to visualize dividing cells. At least a thousand nuclei were counted per mouse and the proliferation index was calculated as the percentage of Ki-67-positive nuclei.

**Cell culture and characterization.** SV40 large T-antigen immortalized murine renal epithelial cells, derived from a Pkd1<sup>lox/lox</sup> mouse, were cultured in DMEM/F12 with glutamine and 15 mM HEPES, 100 U penicillin/streptomycin (Invitrogen, Breda, The Netherlands) supplemented with 2% Ultroser G (Pall BioSepsra, Cergy, France), 2.5 μg/l prostaglandin E<sub>1</sub>, 5 μg/l insulin-transferrin-sodium selenite, and 30 μg/l hydrocortisone obtained from Sigma. To obtain Pkd1<sup>del/del</sup> cells, Pkd1<sup>lox/lox</sup> cells were transfected with a Cre plasmid containing a neomycin resistance cassette using Lipofectamin 2000 Transfection Reagent (Invitrogen) according to the manufacturer’s protocol. After transfection, the cells were cultured in selection medium [similar to medium described above but with 400 μg/ml geneticin (Invitrogen)] and diluted to obtain single clones that were tested for the Pkd1 deletion by PCR (detailed information about the Cre construct and primer sequences can be provided by the corresponding author).

RNA and cDNA were obtained using Tri-Reagent (Sigma) and the Transcriptor First Strand cDNA Synthesis Kit (Roche, Almere, the Netherlands) according to the manufacturer’s protocol. PCR was performed using primers for the proximal tubule marker Cublin (Cubn) and for the collecting duct marker arginine vasopressin receptor 2 (Avpr2) and analyzed on a 2.5% agarose gel. Immunofluorescence using antibodies against β-catenin (1:100; BD Biosciences, Breda, The Netherlands) or α-tubulin (1:200; Sigma) was performed as described previously (29).

At 50% confluence, cells were incubated O/N with 0, 2.5, 5, or 10 μM curcumin (Sigma) in the absence of medium supplements. Then, the cells were cultured with the same curcumin concentrations in the presence of medium supplements for an additional 6 h. Protein lysates were prepared and analyzed by Western blot analysis as described below.

Detection and densitometric analysis. Throughout the lysis procedure, 2 mM NaF, 1 mM Na<sub>2</sub>VO<sub>4</sub>, and protease inhibitor cocktail (Roche) were used to inhibit phosphatase and protease activity. Kidneys were homogenized in 600 μl RIPA without detergents (50 mM Tris·HCl, pH 7.4, 150 mM NaCl, 1 mM EDTA) using Magnafluor (per manufacturer’s protocol). After centrifugation, the cells were cultured in selection medium [similar to medium described above but with 400 μl RIPA without detergents followed by three 5-s pulses of sonification. Next, an equal amount of RIPA with detergents (50 mM Tris·HCl, pH 7.4, 150 mM NaCl, 1 mM EDTA) using Magnafluor technology (Roche) followed by three 5-s pulses of sonification. The rest of the procedure was similar to the kidney lysis preparation. Protein concentrations were determined using BCA protein assay (Thermo Fisher Scientific, Ettten-Leur, The Netherlands). Ten micrograms of cell lysates and 80–120 μg of kidney lysates were used for SDS-PAGE and immunoblotting. Membranes were blocked for 1 h in 25% sea block blocking buffer (Thermo Fisher Scientific/Pierce, no. 35727, Ettten-Leur, The Netherlands) in PBS and incubated O/N with anti-pSTAT3 (1:1,000), anti-p-pS6 (1:1,000); Cell Signaling, no. 9139), anti-p-pS6 (1:2,000), anti-p-pS6 (Ser240/244; 1:1,000; Cell Signaling, no. 2317), or anti-β-actin (1:5,000; Cell Signaling, no. 4967) in 5% BSA-TBS (Twen 20, 0.1%). Goat-anti-rabbit (1:10,000) DyLight 800 (Perbio Science, no. 35571) was used as secondary antibody for the detection of anti-pSTAT3, anti-p-pS6, and anti-β-actin. Goat-anti-mouse DyLight 680 (1:10,000; Perbio Science, no. 35518) was used as secondary antibody for the detection of anti-pSTAT3 and anti-p-pS6. Detection and densitometric analysis were carried out with the Odyssey Infrared Imaging System (Westburg, Leusden, The Netherlands). After detection, the same membrane was used to detect the next protein; i.e., after pSTAT3 detection, the same membrane was used to detect total STAT3 and then β-actin. Because total STAT3 and total pS6 were detected with mouse antibodies (detected at 680 nm) and pSTAT3 and p-pS6 with rabbit antibodies (detected at 800 nm), the total pool of protein could be easily discriminated from the phosphorylated variant. pSTAT3 nicely overlapped with total STAT3 and p-pS6 overlapped with total pS6. From each blot, protein levels were quantified using pSTAT3/STAT3 and p-pS6/pS6 integrated intensity ratios. β-Actin was used as a loading control.
Statistical analysis. Differences in Western blot/densitometric analysis of p-rpS6/rpS6 and pSTAT3/STAT3 ratios between cystic and control samples were tested by Student’s t-tests. Differences in two-kidney weight/body weight (2KW/BW%) ratios, cystic and proliferation indexes, and Western blot/densitometric analysis of p-rpS6/rpS6 and pSTAT3/STAT3 ratios between untreated and curcumin-treated mice at the 11-wk time point were tested using Mann-Whitney U-tests. The Kaplan-Meier method was used to analyze overall survival of curcumin-treated and -untreated mice that were followed until the onset of renal failure and the significance of differences between those groups was analyzed by the Log-Rank (Mantel-Cox) test adjusted for litter effects.

RESULTS

Curcumin and PKD-related pathways. The ability of curcumin to inhibit mTOR and Wnt signaling is likely to be relevant for ADPKD since these signaling routes are altered during cystogenesis (4, 5, 18, 23, 27, 32, 55). Previously, we and others showed that cystogenesis is accompanied by up-regulation of several canonical Wnt targets, including cyclin D1 and survivin (18, 27). Since these genes are also direct targets of STAT3, a transcription factor that is frequently activated in several types of tumors, we investigated possible STAT3 activation in iKsp- Pkd1^del mice (8, 17, 33, 49, 50, 52). In these mice, Pkd1 inactivation by tamoxifen treatment at days 40–42 results in end-stage PKD ~16 wk later (Fig. 1B). Immunohistochemical analysis of renal sections demonstrated numerous regions with strong nuclear accumulation of phosphorylated-(Tyr705)-STAT3 (pSTAT3) in cyst-lining epithelial cells and in interstitial cells, while in kidneys of control mice only few pSTAT3-positive nuclei were observed (Fig. 1, A and D). To further quantify STAT3 activity, total and phosphorylated STAT3 levels were analyzed by Western blot and densitometric analysis. Whereas total STAT3 increased slightly, which did not reach statistical significance, the pSTAT3/STAT3 ratio was strongly elevated in cystic mice and thus revealed that STAT3 was strongly activated in cystic kidneys, thereby confirming the immunohistochemical data (P < 0.01; Fig. 2, A and C).

The relevance of mTOR signaling in ADPKD has previously been demonstrated in a number of ADPKD models where inhibition of mTOR by rapamycin resulted in reduced cyst progression (55, 56, 61). To confirm mTOR activation, ribosomal protein S6 (rpS6) that acts downstream from the mTOR/S6K pathway was analyzed. Numerous cysts displayed intense p-rpS6 levels in cyst-lining epithelial cells, whereas in control mice only negative and weak-to-moderate p-rpS6 expression could be observed (Fig. 1, E and F). Also, Western blot analysis indicated strong activation of rpS6 in cystic kidneys, whereas total levels were not different (P > 0.01; Fig. 2, B and D). Sequential sections stained with antibodies against pSTAT3, p-rpS6, and markers for proximal tubules (megalin), distal tubules (Tamm-Horsfall protein), and collecting ducts (aquaporin-2) revealed that STAT3 and rpS6 activation occurred in all tubular segments (Fig. 1, G-M). These data suggest that mTOR but also STAT3 is activated in PKD.

Interestingly, activation of these proteins is known to be inhibited by curcumin (4, 5, 10, 26). To confirm the effect of curcumin on STAT3 and mTOR activation, we performed in vitro experiments on immortalized renal tubular epithelial cells that were derived from Pkd1^loxp/lox mice. These cells likely originate from the proximal epithelium since they express the proximal marker Cubn but not the collecting duct marker Avpr2 (Supplementary Fig. S1C; the online version of this article contains supplemental data). They were also able to form cysts, as detected by α-tubulin, and expressed β-catenin in the plasma membrane (Supplementary Fig. S1, A and B). A subclone was transfected with a Cre plasmid to generate Pkd1^del,del cells and a pcr was performed to confirm the deletion (Supplementary Fig. S1D). Both Pkd1^lox,lox and Pkd1^del,del cells were incubated with different curcumin concentrations (0, 2.5, 5, and 10 μM) and analyzed for pSTAT3/STAT3 and p-rpS6/rpS6 ratios. β-Actin was used as a loading control. Both cell lines demonstrated a comparable dose-dependent reduction of STAT3 and rpS6 activation, indicating that curcumin inhibits both pathways regardless of Pkd1 expression (Fig. 3, A–D).

Curcumin inhibits cystogenesis in vivo. Given the apparent overlap between curcumin-targeted and PKD-related pathways, we set out to investigate the therapeutic potential of curcumin in our previously developed iKsp- Pkd1^del mouse model (29, 30). An overview of the treatment strategy is given in Fig. 4 and was performed as follows: at PN days 40–42, we administered tamoxifen to male iKsp- Pkd1^del mice to inactivate Pkd1. Seven days after gene disruption, each litter was divided into a control and a curcumin group. Control mice were put on a standard diet and curcumin mice received the same diet supplemented with 1% curcumin. The mice were followed in time and killed after ascertainment of renal failure where a BU concentration of 20 mmol/l or higher was used as criterion (long-treatment group). In addition, a subgroup of mice was killed 11 wk after gene disruption (short-treatment group). For both groups, there was no difference in total body weight between curcumin-treated and -untreated mice (Supplementary Fig. S2A). Control mice that did not receive tamoxifen had normal kidneys (Fig. 1A). Mice that did receive tamoxifen and killed 11 wk later (n = 5 from 2 litters) displayed tubular dilations and small cysts accompanied by increased 2KW/BW% (median 2.6%), cystic index (CI; median 44%), and proliferation index (PI; median 4.2%; Figs. 5A and 6). These parameters were significantly reduced in curcumin-treated littermates (n = 5; 2KW/BW% = 1.9%, P < 0.01; CI = 34%, P < 0.05; and PI = 3%, P < 0.01), which also corresponded to improved histology (Figs. 5, A and B, and 6). To assess whether besides proliferation also apoptosis took part in the curcumin-mediated reduction in cystogenesis, we stained kidney sections from curcumin-treated and -untreated mice for cleaved caspase-3 but found only very few apoptotic cells, indicating that apoptosis did not play a significant role (data not shown).

From mice that were followed until the onset of renal failure (long treatment), BU was measured on a weekly basis. When BU exceeded 15 mmol/l, the frequency of the measurements was increased to pinpoint the moment of renal failure. Both curcumin-treated and -untreated mice reached renal failure. However, curcumin treatment postponed renal failure in all five litters tested (Fig. 7A; also shown combined in Supplementary Fig. S2B). The median survival after tamoxifen treatment for all curcumin-treated mice (n = 12) was 119 days and for all untreated mice (n = 11) 105 days. Kaplan-Meier analysis showed that both groups significantly differed from each other (P < 0.001; Fig. 7B). Taken together, these data
Fig. 1. Expression of phosphorylated signal transducers and activator of transcription 3 (STAT3) and ribosomal protein S6 (rpS6) in cystic kidneys from iKsp-Pkd1<sup>del</sup> mice. A–F: iKsp-Pkd1<sup>del</sup> mice untreated (A, C, and E) or treated (B, D, and F–M) with tamoxifen and killed ~16 wk later (for mice that received tamoxifen, this means that they developed renal failure). A and B: hematoxylin and eosin (HE) staining indicating normal histology in mice without tamoxifen treatment and severe cystic disease 4 mo after tamoxifen administration. C and D: phosphorylated-(Tyr705)-STAT3 (pSTAT3) staining could hardly be observed in control mice but regions with intense staining were observed in mice with polycystic kidney disease (PKD); examples of positive cystic epithelial and interstitial nuclei are indicated by open and closed arrowheads, respectively. E and F: mosaic staining pattern for phosphorylated-rpS6 (p-rpS6) in control and cystic mice showing more areas with increased intensity in cystic kidneys; indicated by arrowheads. G–J: sequential sections of a cystic area stained for the proximal marker megalin (G), the distal marker Tamm-Horsfall protein (H), p-rpS6 (I), and pSTAT3 (J). Overlap between megalin, pSTAT3, and p-rpS6 is indicated by * and overlap between Tamm-Horsfall protein, pSTAT3, and p-rpS6 by #. K–M: sequential sections of a cystic area stained for the collecting duct marker aquaporin-2 (K), p-rpS6 (L), and pSTAT3 (M). Overlap between aquaporin-2, pSTAT3, and p-rpS6 is indicated by π. Magnifications were ×25 for A and B, ×200 for C–F, and ×400 for G–M.
demonstrate that curcumin treatment reduces proliferation resulting in inhibited cystogenesis and delayed renal failure.

**Involvement of pSTAT3 and p-rpS6.** To investigate whether STAT3 and mTOR signaling contributed to the curcumin-mediated inhibition of cystogenesis, we performed immunohistochemical and Western blot analysis on the short-treatment group.

Immunohistochemical staining of pSTAT3 revealed that the reduction of the cystic phenotype by curcumin treatment correlated with fewer nuclei expressing active STAT3 (Fig. 5, C and D). Also, Western blot analysis on the short-treatment groups showed clear activation of STAT3 in untreated mice that was significantly reduced in curcumin-treated mice ($P < 0.01$; Fig. 8, A and C). In kidneys from untreated and curcumin-treated mice that had been followed until renal failure (long treatment), pSTAT3 levels did not differ but were higher compared with the short-treatment group (Supplementary Fig. S3, A and C).

Also, rpS6 was analyzed in the short-treatment group. Although expression of p-rpS6 appeared slightly reduced in curcumin-treated mice (Fig. 5, E and F), the reduction observed by Western blotting did not reach statistical significance (Fig. 8, B and D). Also, the activation of rpS6 compared with controls was not significant, indicating that the role of mTOR at this stage of the disease is limited. RpS6 was clearly activated in the long-treatment groups and curcumin-treated mice displayed a similar reduced yet statistically insignificant trend (Supplementary Fig. S3, B and D).
These data indicate that STAT3 is already activated at a relatively early stage of PKD and suggest that the curcumin-mediated inhibition of cyst formation at least in part acts through STAT3 inhibition. However, mTOR was only significantly activated at end-stage PKD and although curcumin treatment resulted in reduced activation in both the short- and long-treatment groups, this effect was not significant.

**DISCUSSION**

In ADPKD, cyst development is preceded by a critical cellular drop in PC1 and/or PC2 levels resulting in deregulation of many signaling pathways that control proliferation, differentiation, PCP, cell-cell and cell/matrix communication, and fluid transport (13, 43, 44, 67). Altered expression of PCP-related genes such as Fjx1 and Fat4 has been linked to cystogenesis possibly by causing disturbed centrosome positioning of newly formed cells after cell division (18, 51). We observed downregulation of Fjx1 in Pkd1-deletion mice during tissue repair when normally its expression is required (18). This apparent defect in PCP signaling, which is regulated by noncanonical Wnt signaling, was accompanied by increased expression of canonical Wnt targets (18). Moreover, their expression has also been demonstrated to be elevated in human cystic samples (27). Together with the ability of the COOH-terminal tail of PC1 to physically interact with and thereby suppress the activity of β-catenin, these data suggest important links between Wnt signaling and cystogenesis (27).

Expression of canonical Wnt targets, however, may not be exclusively accountable to Wnt signaling. For example, cyclin D1 and survivin are also known targets of STAT3, a transcription factor that, when activated in several types of tumors, is associated with poor prognosis (8, 17, 33, 50, 52). Although STAT3 promotes cell survival and proliferation, it has been demonstrated that overexpression of PKD1 in vitro could activate a STAT3 luciferase reporter construct (7, 49).

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**Fig. 4. In vivo experimental setup.** Tamoxifen is administered to iKsp-Pkd1<sup>del</sup> mice at postnatal (PN) days 40–42 (t = 0) to induce conversion of the Pkd1<sup>lox</sup> to the inactive Pkd1<sup>del</sup> allele. One week later, mice either continued to receive normal food or food supplemented with 1% curcumin. One group was killed 11 wk after tamoxifen (short treatment). Another group was followed until renal failure [blood urea (BU) > 20 mmol/l], meaning that in this group, the time at which mice are killed is the read-out (long treatment).

**DISCUSSION**

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**Fig. 5.** Effect of curcumin treatment on iKsp-Pkd1<sup>del</sup> mice. A–F: most severely affected iKsp-Pkd1<sup>del</sup> mice in the short-treatment group without (A, C, and E) or with (B, D, and F) curcumin treatment. After curcumin treatment, HE staining indicated improved histology (A and B), decreased nuclear pSTAT3 staining (C and D), and to a lesser extent decreased p-rpS6 staining (E and F); arrowheads indicate examples of positive pSTAT3 nuclei. Both untreated (G; 93 days after tamoxifen) and curcumin-treated (H; 117 days after tamoxifen) reached renal failure as a consequence of severe PKD. Magnifications were ×25 for HE stainings and ×200 for pSTAT3 and p-rpS6 stainings.
creased phosphorylation levels, however, were not detected (7). In the same study, the closely linked but counteracting STAT1 was analyzed. Overexpression of PKD1 in Madin-Darby canine kidney (MDCK) cells resulted in enhanced STAT1 activation and subsequent p21waf-induced G0/G1 growth arrest. In line with these results, Pkd1−/− embryos indeed exhibited reduced pSTAT1 and p21waf levels (7). Although the exact contribution remains to be elucidated, our observations indicate that STAT3 activation strongly correlates with increased cyst progression.

Fig. 6. Effect of curcumin treatment on 2-kidney weight/body weight (2KW/BW) ratios, cystic and proliferation indexes. Curcumin-treated iKsp-Pkd1+/− mice that were killed 11 wk after Pkd1 disruption showed significant lower 2KW/BW ratios (1.9%) compared with their untreated littermates (2.6%; P < 0.01). In line with the 2KW/BW ratios, the cystic area and the number of proliferating cells, as determined by Ki-67 staining, were lower in the curcumin-treated mice (44 vs. 34%, P < 0.05 and 4.2 vs. 3.0%, P < 0.01, respectively). These parameters increased in mice that were followed until renal failure but there was no difference between untreated and curcumin-treated mice other than the time frame within renal failure developed.

Fig. 7. Curcumin treatment delays renal failure in iKsp-Pkd1+/− mice in the long-treatment groups. A: from 11 wk after Pkd1 disruption, BU was measured on a weekly basis in curcumin-treated and -untreated iKsp-Pkd1+/− mice until renal failure (BU > 20 mmol/l) was ascertained. Curcumin treatment delayed renal failure in all 5 litters tested. B: Kaplan-Meier analysis of all litters combined revealed a significant difference between the untreated and curcumin-treated mice (P < 0.001, Log-Rank test adjusted for litter). The median survival of untreated mice (n = 11) was 105 days and of curcumin-treated mice (n = 12) 119 days after Pkd1 disruption.
Another crucial modulator of proliferation that is activated in ADPKD is mTOR. It has been shown that, together with tuberin, the COOH-terminal tail of PC1 forms a complex with mTOR, thereby reducing its activity (55). The ability of mTOR inhibitors to greatly reduce cystogenesis in several animal models further supports its role in ADPKD (55, 56, 61). However, the promising results in rodents seem to contrast the data obtained in recent clinical trials that at best demonstrate mild benefits (45, 53, 66). We observed a dose-dependent effect of an mTOR inhibitor on mTOR signaling and on renal cystic disease in our iKsp-Pkd1del mice (Novanic Z, unpublished data). Of note, it has also been shown that the serum concentration of mTOR inhibitors that can be used in clinical practice is too low to inhibit tissue mTOR (9).

Levels of p-rpS6 were not yet significantly elevated in untreated mice at the 11-wk time point. The small and statistically insignificant effect of curcumin on mTOR signaling at this stage is therefore not a major contributor for the reduced cystic phenotype by curcumin treatment and reduced STAT3 signaling supports the involvement of STAT3.

Curcumin was not able to prevent cystogenesis entirely, which may be a consequence of its limited bioavailability. Many studies addressed this issue, leading to curcumin analogs with improved pharmacological profiles and new delivery methods for curcumin (1, 21, 25, 36, 40, 57). In addition, the generation of a series of synthetic derivatives of oleanolic acid, a triterpenoid that can be found in olives, may be an interesting candidate for ADPKD treatment (20, 39). These synthetic oleanane triterpenoids (SO) act on similar pathways as curcumin possibly by the so-called Michael acceptor properties that both these molecules possess (39).

One such SO, CDDO-Me, seems to be active at lower concentrations than curcumin, has successfully been tested in rodent models against several malignant diseases, and is currently being tested in phase I and phase II clinical trials (37–39). An additional property of curcumin-like molecules and SO that was not tested in this study but may be of relevance to ADPKD is their ability to reduce kidney damage by acting as antioxidants and by pharyological profiles and new delivery methods for curcumin (1, 21, 25, 36, 40, 57). In addition, the generation of a series of synthetic derivatives of oleanolic acid, a triterpenoid that can be found in olives, may be an interesting candidate for ADPKD treatment (20, 39). These synthetic oleanane triterpenoids (SO) act on similar pathways as curcumin possibly by the so-called Michael acceptor properties that both these molecules possess (39).

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inducing a series of cytoprotective genes through Nrf2 activation (2, 16, 22). Since we and others showed that cyst progression is greatly accelerated by these kind of stress factors, it would be interesting to see whether alleviating kidney damage by this strategy could slow down the observed acceleration in cyst progression (18, 60).

In conclusion, this study provides evidence that multitarget compounds like curcumin are able to inhibit cyst progression without apparent side effects. Current developments in the field of curcumin analogs and SO are highly promising and should be further evaluated as possible therapeutics for ADPKD.

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DISCLOSURES

No conflicts of interest, financial or otherwise, are declared by the author(s).

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