Characterization of Primary Cilia in Osteoblasts Isolated From Patients With ADPKD and CKD

Renata C Pereira,1 Berenice Y Gitomer,2 Michel Chonchol,2 Peter C Harris,3 Kathleen J Noche,1 Isidro B Salusky,1 and Lauren V Albrecht4

1Department of Pediatrics, David Geffen School of Medicine at UCL, Los Angeles, CA, USA
2Department of Medicine, Division of Renal Diseases and Hypertension, University of Colorado Anschutz Medical Campus, Aurora, CO, USA
3Division of Nephrology and Hypertension, Mayo Clinic, Rochester, MN, USA
4Department of Biological Chemistry, David Geffen School of Medicine at UCLA, Los Angeles, CA, USA

ABSTRACT
Autosomal dominant polycystic kidney disease (ADPKD) is the most common inherited cause of chronic kidney disease (CKD) and leads to a specific type of bone disease. The primary cilium is a major cellular organelle implicated in the pathophysiology of ADPKD caused by mutations in polycystin-1 (PKD1) and polycystin-2 (PKD2). In this study, for the first time, cilia were characterized in primary preosteoblasts isolated from patients with ADPKD. All patients with ADPKD had low bone turnover and primary osteoblasts were obtained from patients with non-ADPKD CKD with low bone turnover. Image-based immunofluorescence assays analyzed cilia using standard markers, pericentrin, and acetylated-α-tubulin, where cilia induction and elongation were chosen as relevant endpoints for these initial investigations. Osteoblastic activity was examined by measuring alkaline phosphatase levels and mineralized matrix deposition rates. It was found that primary cilia can be visualized in patient-derived osteoblasts and respond to elongation treatments. Compared with control cells, ADPKD osteoblasts displayed abnormal cilia elongation that was significantly more responsive in cells with PKD2 nontruncating mutations and PKD1 mutations. In contrast, non-ADPKD CKD osteoblasts were unresponsive and had shorter cilia. Finally, ADPKD osteoblasts showed increased rates of mineralized matrix deposition compared with non-ADPKD CKD. This work represents the first study of cilia in primary human-derived osteoblasts from patients with CKD and patients with ADPKD who have normal kidney function, offering new insights as bone disease phenotypes are not well recapitulated in animal models. These data support a model whereby altered cilia occurs in ADPKD and has lower fracture rates.(8,9) The molecular mechanisms, underpinning how bone disease is propagated in patients who have CKD with PKD mutations versus patients who have non-ADPKD CKD, remain to be completely understood.

The primary cilium plays a central role in bone biology where PKD1/2-encoded polycystin (PC) proteins are expressed in both osteoblasts and osteocytes. Bone formation and resorption is orchestrated by bone cell activities and responses to mechanical stress through complex cellular mechanisms. Cilia-related signaling through the PC1/2 proteins has been linked to the regulation of fluid-flow mechanosensation. Many cilia-related}

This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.
Received in original form September 8, 2020; revised form January 2, 2021; accepted January 12, 2021. Accepted manuscript online January 22, 2021.
Address correspondence to: Lauren V Albrecht, PhD, 675 Charles E Young Drive, 5-712 MRL, Los Angeles, CA 90095. Email: lalbrecht@mednet.ucla.edu

JBMR Plus published by Wiley Periodicals LLC. on behalf of American Society for Bone and Mineral Research.
signaling pathways are also implicated in ADPKD such as canonical Wnt signaling, which plays an essential role in bone formation and is thought to activate PC1/2 heterodimerization in cilia. Animal models further underscore the importance of cilia in bone development as PKD1 osteoblast-specific conditional knockout models result in abnormal bone morphology, osteopenia, and low BMD. However, such PKD animal models differed from those observed in patients with ADPKD. Differences between both models are undefined, and whether the low bone turnover in ADPKD arises directly from altered cilia-related signal transduction remains an open area of research.

ADPKD has been classified as a ciliopathy whereby the primary cilium operates as the major organelle associated with disease. The primary cilium is a transient structure that forms during a process called ciliogenesis. Dynamic regulation of cilium length is required for cilia-related tissue activity and cellular signaling. Cilia elongation promotes mechanotransductive responses to strain through integrin signaling, whereas short cilia decrease intracellular cAMP and ion channel activity, which results in impaired fluid shear stress. Further, longer cilia were linked to tissue regeneration during injury whereby cilia-emanating signaling was associated with enhanced proliferation. Intriguingly, cystic ADPKD kidney cells have also been reported to increase rates of proliferation and growth. Thus, it has even been proposed that cilia length may operate as an anticystogenic repair program that becomes aberrant in ADPKD and contributes cystic pathology.

Here, we utilized primary preosteoblast cells derived from patients with ADPKD and patients who have non-ADPKD CKD to investigate for the first time primary cilia in human bone. Patients with non-ADPKD CKD were chosen based on similar adynamic bone disease phenotypes as compared with ADPKD. Low bone turnover in ADPKD arises directly from altered cilia-related signal transduction remains an open area of research.

Patient data

Patients with ADPKD were selected from a cohort with glomerular filtration rates (GFRs) above 90 mL/min/1.73 m². The bone formation rates (BFRs) of patients with ADPKD were consistent with low bone turnover or adynamic bone disease and were compared with patients who have non-ADPKD CKD with similar bone turnover in early and advanced CKD stages as previously reported. Patient bone samples with mutations in either PKD1 or PKD2 (four patients) were chosen from both male and female subjects aged between 23 and 44 years. Importantly, PKD mutations included both truncating and nontruncating variations as mutation type has been linked to the rate of kidney disease progression. The BFRs/BSs (μm²/μm²/y) of the patients with ADPKD were PKD1-truncating (T; 0), PKD1-nontruncating (NT; 4.5), PKD2-T (4.0), and PKD2-NT (5.4). The patients with early-stage non-ADPKD CKD had BFRs/BSs (μm²/μm²/y) of 5.4 and 0.001, whereas those of healthy controls were 21.8 and 10.2. Replicate analyses were performed using cells derived from both male and female patients as healthy subjects. The total number of patients used for these analyses included two healthy controls, three patients with early-stage non-ADPKD CKD, two patients with late-stage non-ADPKD, and four patients with ADPKD with normal GFRs of 80 and 110 (mL/min/1.73 m²).

Primary osteoblast cell isolation

Primary patient-derived osteoblastic cells were cultured from bone biopsies as performed previously. Bone biopsies isolated from healthy volunteers were used as controls. Bone tissues were obtained from the anterior iliac crest using a bone biopsy trephine needle as previously described. Primary preosteoblasts were isolated from small fragments of cancellous bone and were cultured in DMEM supplemented with 20% fetal calf serum, 100-mU/mL penicillin, 100-mU/mL streptomycin, 250-ng/mL amphotericin B (Invitrogen), and 100-μg/mL ascorbic acid. Bone fragments were washed with fresh medium containing antibiotic–antimycotic solution, and fragments were minced into very small pieces, which were divided into six culture plates and cultured under standard 5% CO₂ at 37°C. Upon cell attachment to plates and sufficient growth, the medium was changed twice weekly until 10–20 cell colonies were observed in each plate, which occurred within 1–4 weeks. Next, cell colonies were trypsinized, pooled, and subcultured. Upon reaching 80% confluence, the contents of the 19 plates were frozen in 20 mL of freezing medium composed of 10% DMSO and 30% FBS and stored in liquid nitrogen. Cells were harvested at equal confluences, and fed 24 hours prior to harvest to ensure reproducibility.

Immunofluorescence, cell signaling, and cilia quantification

ADPKD, non-ADPKD CKD, and healthy subject osteoblasts were plated at the same density and cultured until confluent. Osteoblast phenotypes were confirmed by staining with standard markers as previously performed. Cells were first incubated in serum-free medium for 24 hours prior to treatments to induce growth arrest as previously performed. For cilia elongation experiments, cells were treated with either 10 or 25mM LiCl or no treatment (control) for 24 hours. Finally, cells were fixed in 4% paraformaldehyde and blocked with standard blocking buffer. Primary cilia were visualized using the antibodies acetylated tubulin and pericentrin; pericentrin marks the cilium base (Abcam). Cilia length was measured using pericentrin and acetylated tubulin. Cilia with lengths <1 μm were discarded. The number of ciliated cells were counted using CellSens software package (Olympus Corp); lengths measured per condition included 100–200 cells.

Bone cell activity and mineralization assays

Biochemical parameters were determined using standard bioanalytical approaches. PKD mutation type in patients with ADPKD
was determined either at the University of Colorado (Aurora, CO) or at the Mayo Clinic (Rochester, MN) by PCH. ALP activity in cells was measured by ELISA (Abbexa Ltd). The rate of mineralized matrix deposition of osteoblast cells was determined using differentiation cocktails and was assessed weekly by staining with 1% (w/v) Alizarin Red S (Sigma-Aldrich).

|                | PKD1-Truncated (T) | PKD1-Nontruncated (NT) |
|----------------|--------------------|------------------------|
| Polycystin-1 (PC1) (462.5 kDa) | c.3114             | c.8791 +5G>C Gly2931fs |

|                          | PKD2-Truncated (T) | PKD2-Nontruncated (NT) |
|--------------------------|--------------------|------------------------|
| Polycystin-2 (PC2) (109.7 kDa) | c.1249>C>T (Arg417*) | c.1320-5T>G (Leu441fs) |

Fig 1. Primary cilia visualized in preosteoblasts derived from patients with autosomal dominant polycystic kidney disease (ADPKD). (A) Diagram of PKD1 and PKD2 genes encoding polycystin-1 (PC-1; 462.5 kDa) and polycystin-2 (PC-2; 109.7 kDa), respectively. The patients with ADPKD chosen were identified to contain truncating or nontruncating variations of both genes. PKD1 mutations include truncation (navy) at codon 3114 or a nontruncated form (aqua) with mutations at codon 8791. PKD2 mutations include truncation (dark green) at codon 1249 and nontruncated (light green) at codon 1320. (B) Cartoon depiction of primary cilia including PC-1 and PC-2 in the primary cilia, acetylated tubulin (green), and the basal body with pericentrin (red) at the base of the cilium. (C) Primary cilia (arrows) marked by pericentrin (red) and acetylated tubulin (green) in primary cultured osteoblasts from patients with ADPKD (PKD1) and healthy controls in 10% serum or serum-starved conditions (24 hours), assessed by immunofluorescence staining; the nuclei were marked by DAPI (blue). Scale bar represents 50 μm. acetylTub = acetylated-α-tubulin; DAPI = 4,6-diamidino-2-phenylindole.
Results

Primary cilia length and responsiveness is altered in ADPKD preosteoblastic cells

Given the distinct differences in the bone of patients with ADPKD and PKD animal bone models, we examined whether preosteoblast cells derived from patients with ADPKD could serve as a model to study cilia in human bone. Preosteoblast cells were isolated from bone biopsies obtained from patients with ADPKD and patients with non-ADPKD CKD with adynamic bone as defined histologically by low BFRs and biochemically by low-circulating bone ALP levels. Patients with ADPKD showed...
confirmed mutations in either PKD1 or PKD2.\(^{40-42}\) Disease progression of ADPKD is associated with the type of PKD mutation present. PKD1 mutations have been associated with worse prognosis and earlier mortality than patients with PKD2 mutations.\(^{40-42}\) Therefore, we selected both PKD1 and PKD2 with either truncating or nontruncating mutations (Fig. 1A). Patients with ADPKD were selected from a cohort with stage 1 CKD, defined by GFRs above 90 ml/min/1.73m², to reduce potential off-target effects of renal dysfunction for these initial bone analyses.

Primary cilia have been thoroughly examined in transformed cell lines and in animal models, but never in primary osteoblasts isolated from human patients. To test whether cilia form properly, cultured osteoblasts were assessed in a series of immunofluorescence-based assays. To first examine cilia induction, cells were cultured in complete medium and starved of serum for 24 hours as performed previously.\(^{43}\) To visualize cilia, cells were fixed and stained with antibodies targeting standard cilia markers of microtubules, with acetylated α-tubulin and pericentrin (Fig. 1B,C).\(^{27,44}\) Normal cell morphology was observed for osteoblasts of healthy cells and PKD mutant cells (Fig. 1C). Importantly, cilia induction by serum starvation was observed in all conditions (Fig. 1C,D), consistent with what has been well-described in immortalized cell lines.\(^{14}\) Even in basal conditions prior to any treatment, PKD mutant cells showed longer cilia when compared with control cells.

As dynamic regulation of cilia length is essential for cellular and mechanotransductive signaling,\(^{30}\) we next assessed cilia elongation as an initial indicator of cilia responsiveness. Cells were treated with LiCl as it induces cilia elongation in mice osteoblasts isolated from human patients. To test whether cilia form in all conditions (Fig. 1C,D), consistent with what has been well-described in immortalized cell lines.\(^{14}\) Even in basal conditions prior to any treatment, PKD mutant cells showed longer cilia when compared with control cells.

As dynamic regulation of cilia length is essential for cellular and mechanotransductive signaling,\(^{30}\) we next assessed cilia elongation as an initial indicator of cilia responsiveness. Cells were treated with LiCl as it induces cilia elongation in mice osteoblasts.\(^{45}\) Notably, LiCl activates canonical Wnt signaling, a pathway previously implicated in cyst expansion of polycystic kidney disease in animal models, through the inhibition of glycogen synthase kinase 3.\(^{45}\) Following serum starvation, cells were treated with 10mM or 25mM LiCl for 24 hours and processed for immunofluorescence analyses. Cilia length was measured using pericentrin to mark the cilium base acetylated α-tubulin as microtubules extend to the tip of the organelle. Ciliated cell numbers were unchanged in all conditions. In control conditions, cilia elongation was initiated as expected, increasing in a dose-responsive manner by 0.9 μm and 2.4 μm in 10mM and 25mM LiCl treatments, respectively (Fig. 2A,B). Similar baseline cilia lengths were observed across ADPKD-derived osteoblasts; however, levels of cilia responsiveness were highly varied across different PKD mutations. Nontruncating PKD2 mutant osteoblasts were not significantly increased in response to LiCl treatments, increasing by only 0.7 μm with 25mM LiCl (not significant; Fig. 2B). Strikingly, in osteoblasts with truncating PKD2 and PKD1 mutations, cilia showed significantly higher responses to LiCl, where cilia length increased by 3.75 μm and 5.49 μm, respectively (Fig. 2A,B). Osteoblasts with nontruncating PKD1 mutations also increased significantly more than controls: by 4.5 μm with 25mM LiCl (Fig. 2A,B). It is interesting to note that the highest levels of abnormal cilia elongation occur with truncating mutations in PKD1 as these have been correlated with the worse prognosis in patients.

Given the similarities of adynamic bone defects and histology, we next assessed cilia responsiveness in non-ADPKD CKD patient osteoblasts. In contrast, cilia elongation was not significantly increased in non-ADPKD CKD cells where cilia length was only increased by 1.1 μm following treatment with 25mM LiCl (Fig. 2B). That cilia elongation was significantly higher in ADPKD relative to osteoblasts from other types of CKDs suggests that ciliary dysregulation is related to PKD mutations and supports a model whereby cilia defects may be linked to molecular mechanisms underlying the distinct low bone turnover in ADPKD.

ADPKD osteoblasts accelerate mineralized matrix deposition when compared with osteoblasts from patients with non-ADPKD CKD and adynamic bone disease

Low bone turnover in CKD is associated with higher fracture rates and decreased mineralization.\(^{38}\) Circulating levels of bone ALP is an indicator of osteoblastic activity and has been reported to be low in patients with ADPKD.\(^{38,39}\) Consistent with this, healthy controls had circulating bone ALP levels of 20 (μg/L) and total circulating ALP levels of 70 (IU/L). In contrast, ADPKD bone ALP levels were lower in all four patients with PKD1-T (9.6 μg/L), PKD1-NT (6.8 μg/L), PKD2-T (8.4 μg/L), and PKD2-NT (9.6 μg/L). Total ALP levels across this ADPKD cohort were PKD1-T (42 IU/L), PKD1-NT (35 IU/L), PKD2-T (44 IU/L), and PKD2-NT (40 IU/L). Next, we assessed osteoblast activity by monitoring ALP in cultured cells from our ADPKD cohort with low bone ALP. Compared with healthy control cells, ADPKD osteoblasts showed lowered ALP activity as in previous reports of low ALP levels in cultured osteoblasts from non-ADPKD CKD (Fig. 3A).\(^{46}\) Next, we examined Alizarin Red in cultured ADPKD osteoblasts as a marker of mineralized matrix deposition. Cells were plated at equal density and maintained in an osteoblast differentiation cocktail of ascorbic acid, β-glycerol phosphate, and dexamethasone as previously performed.\(^{38}\) Time-course analyses, which assessed Alizarin Red S staining over the course of 21 days, revealed that healthy control cells began to show visible mineralization after 14 days and continued to increase through 21 days (Fig. 3B-D). Osteoblasts with truncating PKD1/2

---

**FIG 2.** Primary cilia elongation by lithium chloride (LiCl) is enhanced in autosomal dominant polycystic kidney disease (ADPKD) osteoblasts. (A) Primary cultured osteoblasts from patients with ADPKD, non-ADPKD chronic kidney disease (CKD), and healthy controls treated with 10mM LiCl or control medium and stained with the antibodies pericentrin (red) and acetylated tubulin (green) for immunofluorescence analyses. Nuclei were marked by 4,6-diamidino-2-phenylindole (DAPI; blue). White-dashed boxes mark representative cilia from each condition: healthy controls, PKD1-truncated (PKD1-T), PKD1-nontruncated (PKD1-NT), PKD2-truncated (PKD2-T), and PKD2-nontruncated (PKD2-NT). (B) Scatter-plot quantification of cilia length following 10mM LiCl, 25mM LiCl, and control treatments in serum-starved osteoblasts from patients with ADPKD and non-ADPKD CKD, and healthy controls. Healthy cilia increased from 2.6 ± 0.1 μm in serum-starved cells to 3.5 ± 0.2 μm with 10mM LiCl and 4.95 ± 0.3 μm with 25mM LiCl (**p < 0.001). PKD1-T cilia increased from 3.9 ± 0.2 μm in control treatments to 5.4 ± 0.3 μm with 10mM LiCl and 9.4 ± 0.5 μm with 25mM LiCl (**p < 0.001). PKD1-NT cilia increased from 4.2 ± 0.2 μm to 5.5 ± 0.3 μm and 8.7 ± 0.7 μm (**p < 0.001). PKD2-T cilia increased from 3.7 ± 0.2 μm in control treatments to 7.3 ± 0.4 μm with 10mM LiCl and 7.4 ± 0.4 μm with 25mM LiCl (**p < 0.001). PKD2-NT cilia did not significantly increase following LiCl treatment (4.4 ± 0.2 μm in controls; 4.3 ± 0.2 μm with 10mM LiCl; 5.1 ± 0.2 μm with 25mM LiCl; **p < 0.001). Error bars indicate mean ± SEM (n > 200). p Values were calculated by Bonferroni corrected pairwise comparisons. Scale bar = 50 μm.
mutations showed significantly higher levels of Alizarin Red S staining at both the 14- and 21-day time points, exceeding levels of control cells (Fig. 3B–D). Increased mineralized nodules were also apparent in PKD1/2 truncated osteoblasts, visualized under light microscopy (Fig. 3C). In contrast, osteoblasts with PKD2 non-truncating mutations showed lower staining by Alizarin Red S.
throughout all time points. Similarly, osteoblasts derived from patients with non-ADPKD CKD also showed lower levels of Alizarin Red S staining compared with healthy controls, consistent with previous reports (Fig. 3E). Thus, these data suggest that the PKD mutation type may be correlated with specific rates of mineralized matrix deposition, as was the case with cilia responsiveness. Furthermore, though both patients with ADPKD and non-ADPKD CKD have low bone turnover, important differences in bone quality such as fractures exist in both groups.(9) Whether primary cilia are responsible for the differences in bone from patients with CKD with or without PKD mutations remains an open area of research that warrants larger-scale studies with more patients.

Discussion

In sum, we report that osteoblasts derived from patients with ADPKD represent a key resource for studies of cilia and PKD mutations, where bone phenotypes are not well-represented in animal models.(8,9) Low-circulating ALP levels in patients with ADPKD were recapitulated in cultured primary osteoblasts, and cilia induction was similar to that seen with immortalized cell lines, supporting the utility of this system for future large-scale analyses. Specifically, patient-derived osteoblasts could provide an important way into the investigation of responses of human cells to new cilia-targeted treatments that reduced cyst formation in mice.(47,48) Using this approach, we uncovered that different PKD mutation types resulted in distinct cilia responsiveness and accelerated mineralized matrix deposition when compared with osteoblasts derived from patients with non-ADPKD CKD with adynamic bone (Fig. 4). Further, the defective cilia signaling in ADPKD cells showed increased Alizarin Red staining, and mineralized nodules suggest that PKD1 mutations and PKD2-T mutations correlate with increased mineralized matrix deposition. However, future studies on the relative contributions of preosteoblast differentiation into mature osteoblasts in relation to mineralization could uncover new insights.

Genotype–phenotype studies of ADPKD have the potential to advance diagnostics and therapeutic strategies. For example, mutations in PKD1 are more prevalent and lead to more severe kidney disease when compared with PKD2 mutations. Furthermore, truncations of PKD1 are associated with greater deterioration of kidney function than nontruncating variations of PKD1. Our data support a model whereby truncating or nontruncating mutations in PKD1 and PKD2 lead to differential responsiveness of cilia in osteoblasts. We report that PKD2 nontruncating mutations showed the least responsive cilia in the presence of LiCl when compared with truncating PKD2 or PKD1 mutations. Wnt signaling activity is linked to kidney disease in ADPKD.(19,20) As LiCl is a known activator of Wnt signaling, our data support a model whereby PKD1-truncating mutations, which are associated with more severe kidney disease, lead to osteoblasts with the most-responsive cilia and accelerated mineralized matrix deposition rates. In contrast, the PKD2 nontruncating mutations that are associated with the mildest kidney disease would not exhibit the exaggerated sensitivity to Wnt inhibition. Future work comparing the molecular signaling cascades across tissues with truncating PKD mutations, such as frameshifting indels, nonsense, and splicing; or nontruncating PKD mutations from missense or in-frame indels could lead to new insights into genetic-based prognosis and targeted therapy.

Studies including an increased number of patients will be required to definitively assess whether PKD mutation types globally determine cilia functions in bone. Because bone defects in ADPKD are already present with normal renal function,(9) future work on the molecular players linking cilia and bone health may help identify future therapies. The sensitivity of PKD-mutated osteoblasts to LiCl implicates the Wnt signaling pathway, which is consistent with reports of Wnt signaling activity...
in ADPKD kidneys. Finally, we propose that the use of primary non-ADPKD CKD osteoblasts also provides an approach for understanding the bone disease across CKD etiologies, which could have important implications for the underlying mechanisms of crosstalk between the kidney and bone.

Acknowledgments

This study was funded by the National Institute of Diabetes and Digestive and Kidney Diseases (NIH/NIDDK; R01 DK094796). LVA was funded through the A.P. Gianninni Award. The content is solely the responsibility of the authors and does not necessarily represent the official views of the NIH, the U.S. Department of Veterans Affairs, or the United States Government.

Authors’ roles: Experiment performance: RCP and KN. Genetic analyses: BYG, MC, and PCH. Data analysis: RCP, IBS, and LVA. Manuscript writing, revision, and edit: LVA, RCP, and IBS.

Conflict of Interest

Dr. Isidro B Salusky reports a grant from the National Institute of Diabetes and Digestive and Kidney Diseases (NIH/NIDDK; R01 DK35423, U01 DK 119950) and grants from Abbvie and Amgen outside the submitted work. Dr. Michel Choncol reports grants from the National Institute of Diabetes and Digestive and Kidney Diseases and from the Department of Defense during the conduct of the study, as well as grants from Otsuka, Sanogi, and Kadmon outside the submitted work. Dr. Peter C Harris reports grants from Otsuka Pharmaceuticals and other support from Otsuka Pharmaceuticals, as well as other support from Mitobridge, Regulus, and Vertex Pharmaceuticals outside the submitted work.

AUTHOR CONTRIBUTIONS

Renata Pereira: Conceptualization; data curation; formal analysis; investigation; methodology; writing-review & editing. Berenice Gitomer: Resources. Michel Choncol: Resources. Peter Harris: Resources. Kathleen Noche: Data curation. Isidro Salusky: Conceptualization; investigation; project administration; resources; supervision; writing-review & editing. Lauren Albrecht: Formal analysis; writing-original draft; writing-review & editing.

PEER REVIEW

The peer review history for this article is available at https://publons.com/publon/10.1002/jbm4.10464.

References

1. Grantham J. Mechanisms of progression in autosomal dominant polycystic kidney disease. Kidney Int Suppl. 1997;63:593-597.
2. Gall E, Torres V, Harris P. Genetic complexity of autosomal dominant polycystic kidney and liver diseases. J Am Soc Nephrol. 2018;29:13-23.
3. Veldhuisen B, Saris J, Haji S, et al. A spectrum of mutations in the second gene for autosomal dominant polycystic kidney disease (PKD2). Am J Hum Genet. 1997;61:547-555.
4. Hughes J, Ward C, Peral B, et al. The polycystic kidney disease 1 (PKD1) gene encodes a novel protein with multiple cell recognition domains. Nat Genet. 1995;10:151-160.
5. Miller P. Chronic kidney disease and the skeleton. Bone Res. 2014;2:14044.
6. Hygum K, Starup-Linde J, Harsløf T, Vestergaard P, Langdahl B. Mechanisms in endocrinology: diabetes mellitus, a state of low bone turnover – a systematic review and meta-analysis. Eur J Endocrinol. 2017;176:R137-R157.
7. Rechter S, Bachetta J, Godlefroid N, et al. Evidence for bone and mineral metabolism alterations in children with autosomal dominant polycystic kidney disease. J Clin Endocrinol Metab. 2017;102:4210-4217.
8. Evenepoel P, Claes K, Cavalier E, et al. A distinct bone phenotype in ADPKD patients with end-stage renal disease. Kidney Int. 2019;95:412-419.
9. Gitomer B, Pereira R, Salusky IB, et al. Mineral bone disease in autosomal dominant polycystic kidney disease. Kidney Int. 2020;90:30958-30961.
10. Yuan X, Yang S. Primary cilia and intraflagellar transport proteins in bone and cartilage. J Dent Res. 2016;95:1341-1349.
11. Nguyen A, Jacobs C. Emerging role of primary cilia as mechanosensors in osteocytes. Bone. 2013;54:196-204.
12. Bonewald L, Johnson M. Osteocytes, mechanosensing and Wnt signaling. Bone. 2008;42:606-615.
13. Bonewald L. The amazing osteocyte. J Bone Miner Res. 2011;26:229-238.
14. Ma M, Tian X, Igarashi P, Pazour G, Somlo S. Loss of cilia suppresses cyst growth in genetic models of autosomal dominant polycystic kidney disease. Nat Genet. 2013;45:1004-1012.
15. Besschetnova T, Kolpakova-Hart E, Guan Y, Zhou J, Olsen B, Shah J. Identification of signaling pathways regulating primary cilium length and flow-mediated adaptation. Curr Biol. 2010;20:182-187.
16. Hopp K, Ward C, Hommerding C, et al. Functional polycystin-1 dosage governs autosomal dominant polycystic kidney disease severity. J Clin Invest. 2012;122:4257-4273.
17. Xiao Z, Zhang S, Mahlós J, et al. Cilia-like structures and polycystin-1 in osteoblasts/osteocytes and associated abnormalities in skeletalogenesis and Runx2 expression. J Biol Chem. 2006;281:30884-30895.
18. Xiao Z, Dallas M, Qiu N, et al. Conditional deletion of Pkd1 in osteocytes disrupts skeletal mechanosensing in mice. FASEB J. 2011;25:2418-2432.
19. Miyazaki A, Sugimoto A, Yoshizaki K, et al. Coordination of WNT signaling and cilogenesis during odontogenesis by piezo type mechanosensitive ion channel component 1. Sci Rep. 2019;9:14762.
20. Pala R, Alomari N, Nauli S. Primary cilium-dependent signaling mechanisms. Int J Mol Sci. 2017;18:2272.
21. Thoma C, Frew I, Hoerner C, Montani M, Moch H, Krek W. pVHL and GSK3beta are components of a primary cilia-maintenance signaling network. Nat Cell Biol. 2007;9:588-595.
22. Xiao Z, Zhang S, Cao L, Qiu N, David V, Quarles L. Conditional disruption of Pkd1 in osteoblasts results in osteopenia due to direct impairment of bone formation. J Biol Chem. 2010;285:1177-1187.
23. Xiao Z, Quarles L. Role of the polycystin-primary cilia complex in bone development and mechanosensing. Ann NY Acad Sci. 2010;1192:410-421.
24. Qiu N, Xiao Z, Cao L, David V, Quarles L. Conditional mesenchymal disruption of Pkd1 results in osteopenia and polycystic kidney disease. Plos One. 2012;7:e46038.
25. Xiao Z, Cao L, Liang Y, et al. Osteoblast-specific deletion of Pkd2 leads to low-turnover osteopenia and reduced bone marrow adiposity. Plos One. 2014;9:e114198.
26. Lu W, Shen X, Pavlova A, et al. Comparison of Pkd1-targeted mutants reveals that loss of polycystin-1 causes cystogenesis and bone defects. Hum Mol Genet. 2001;10:2385-2396.
27. Sharma N, Berbaki N, Yoder B. Ciliary dysfunction in developmental abnormalities and diseases. Curr Top Dev Biol. 2008;85:371-427.
28. Gerdes J, Davis E, Katsanis N. The vertebrate primary cilium in development, homeostasis, and disease. Cell. 2009;137:32-45.
29. Schwartz R, Hildebrandt F, Benzing T, Katsanis N. Ciliopathies. N Engl J Med. 2011;364:1533-1543.
30. Nakamura T, Asano-Hoshino A, Suzuki T, et al. The elongation of primary cilia via the acetylation of α-tubulin by the treatment with lithium chloride in human fibroblast KD cells. Med Mol Morphol. 2015;48:44-53.
31. Vaughan T, Mullen C, Verbruggen S, McNamara L. Bone cell mechanosensation of fluid flow stimulation: a fluid–structure interaction
model characterising the role integrin attachments and primary cilia. Biomech Model Mechanobiol. 2015;14:703-718.

32. Harris P, Hopp K. The mutation, a key determinant of phenotype in ADPKD. J Am Soc Nephrol. 2013;24:868-870.

33. Heyer C, Sundsbak J, Abebe K, et al. Predicted mutation strength of nontruncating PKD1 mutations aids genotype-phenotype correlations in autosomal dominant polycystic kidney disease. J Am Soc Nephrol. 2016;27:2872-2884.

34. Ma M, Gallagher A, Somlo S. Ciliary mechanisms of cyst formation in polycystic kidney disease. Cold Spring Harb Perspect Biol. 2017;9: a028209.

35. Patel V, Li L, Cobo-Stark P, et al. Acute kidney injury and aberrant planar cell polarity induce cyst formation in mice lacking renal cilia. Hum Mol Genet. 2008;17:1578-1590.

36. Leonhard W, Happe H, Peters D. Variable cyst development in autosomal dominant polycystic kidney disease: the biologic context. J Am Soc Nephrol. 2016;27:3530-3538.

37. Takakura A, Contrino L, Zhou X, et al. Renal injury is a third hit promoting rapid development of adult polycystic kidney disease. Hum Mol Genet. 2009;18:2523-2531.

38. Pereira R, Delany A, Khouzam N, et al. Primary osteoblast-like cells from patients with end-stage kidney disease reflect gene expression, proliferation, and mineralization characteristics ex vivo. Kidney Int. 2015;87:593-601.

39. Reed B, McFann K, Kimberling W, et al. Presence of de novo mutations in autosomal dominant polycystic kidney disease patients without family history. Am J Kidney Dis. 2008;52:1042-1050.

40. Rossetti S, Kubly V, Consugar M, et al. Incompletely penetrant PKD1 alleles suggest a role for gene dosage in cyst initiation in polycystic kidney disease. Kidney Int. 2009;75:848-855.

41. Gout A, Consortium A, Ravine D, et al. Analysis of published PKD1 gene sequence variants. Nat Genet. 2007;39:427-428.

42. Gall E, Alam A, Perrone R. Autosomal dominant polycystic kidney disease. Lancet. 2019;393:919-933.

43. Wheatley D. Cilia in cell-cultured fibroblasts. 3. Relationship between mitotic activity and cilium frequency in mouse 3T6 fibroblasts. J Anat. 1971;110:367-382.

44. Lee S, Somlo S. Cyst growth, polycystins, and primary cilia in autosomal dominant polycystic kidney disease. Kidney Res Clin Pract. 2014;33:73-78.

45. Zhou J, Gao Y, Zhu B, et al. Sinusoidal electromagnetic fields increase peak bone mass in rats by activating Wnt10b/β-catenin in primary cilia of osteoblasts. J Bone Miner Res. 2019;34:1336-1351.

46. Pereira RC, Salusky IB, Bowen RE, Freymiller EG, Wesseling-Perry K. Vitamin D sterols increase FGF23 expression by stimulating osteoblast and osteocyte maturation in CKD bone. Bone. 2019;127:626-634.

47. Shimada H, Lu Q, Insinna-Kettenhofen C, et al. In vitro modeling using ciliopathy-patient-derived cells reveals distinct cilia dysfunctions caused by CEP290 mutations. Cell Rep. 2017;20:384-396.

48. Nikonova A, Deneka A, Kiseleva A, et al. Ganetespib limits ciliation and cystogenesis in autosomal-dominant polycystic kidney disease (ADPKD). Faseb J. 2018;32:2735-2746.