Metabolic Reprogramming and Reconstruction: Integration of Experimental and Computational Studies to Set the Path Forward in ADPKD

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Metabolic reprogramming is a key feature of Autosomal Dominant Polycystic Kidney Disease (ADPKD) characterized by changes in cellular pathways occurring in response to the pathological cell conditions. In ADPKD, a broad range of dysregulated pathways have been found. The studies supporting alterations in cell metabolism have shown that the metabolic preference for abnormal cystic growth is to utilize aerobic glycolysis, increasing glutamine uptake and reducing oxidative phosphorylation, consequently resulting in ADPKD cells shifting their energy to alternative energetic pathways. The mechanism behind the role of the polycystin proteins and how it leads to disease remains unclear, despite the identification of numerous signaling pathways. The integration of computational data analysis that accompanies experimental findings was pivotal in the identification of metabolic reprogramming in ADPKD. Here, we summarize the important results and argue that their exploitation may give further insights into the regulatory mechanisms driving metabolic reprogramming in ADPKD. The aim of this review is to provide a comprehensive overview on metabolic focused studies and potential targets for treatment, and to propose that computational approaches could be instrumental in advancing this field of research.

Keywords: metabolism, systems biology, systems medicine, therapeutics, drug repositioning, ADPKD

INTRODUCTION

Autosomal Dominant Polycystic Kidney Disease (ADPKD) is a common monogenic disorder characterized by bilateral renal cyst formation and extra-renal manifestations. These manifestations include cysts in other organs, such as the liver and pancreas, as well as abnormalities including intracranial aneurysms. Worldwide around 12.5 million people suffer from ADPKD, and it is a common cause of end-stage renal disease (1). ADPKD is caused by mutations in either PKD1 (polycystin-1) or PKD2 (polycystin-2) genes, which account for almost 80 and 15% of ADPKD cases, respectively. More recently, a third gene has been described, GANAB (which encodes neutral alpha-glucosidase AB), which results in a milder renal phenotype and a more severe liver phenotype (2).

The polycystin1 (PC1) protein contains a large extracellular N-terminal region (3074 aa), 11 transmembrane domains (1032 aa), and a short intracellular C-terminal region (198 aa) (3, 4). PC1 interacts with PC2 through a coiled-coil domain in the C-terminal...
portions (5). The PC1 and PC2 complexes are detected in the ER membrane, the plasma membrane, exosomes, and the primary cilia, where they have been suggested to mediate calcium influx in response to fluid flow (6). Notwithstanding, the function of polycystin proteins in cilia and their role in calcium fluxes has been intensely debated (7).

The polycystin proteins have been shown to modulate numerous signaling pathways [please refer to this review for evidence of altered signaling pathways (8)] however their function remains unclear. In the last decade, research aimed at elucidating metabolic-related pathways in ADPKD has grown exponentially, and metabolic reprogramming is now considered a main feature of ADPKD (9–12). The number of researchers using multiple-omics (metabolomics, transcriptomics, genomics, and lipidomics) approaches is still growing and these studies have been pivotal in gaining substantial insights into the pathogenesis of PKD. There is a need for the scientific community to establish principles for data management, which can be challenging to interpret, unless efficiently analyzed. This huge amount of data creates more confusion and blurring than understanding.

One solution to overcome this problem is through the integration of Systems Biology approaches, which offer powerful abstraction tools to analyze biological reactions in a cell, providing a snapshot that is similar to its observable phenotype. Computational tools can be used not only to explore comprehensive molecular mechanisms but also to discover and test novel therapeutic targets. For example, they were recently applied to identify novel drug candidates for the treatment of ADPKD (13, 14). A possible System Biology workflow for the development of possible therapeutic targets that integrates wet-lab experiments, statistical and computational analysis, and System Medicine, is proposed in Figure 1. Systems Medicine represents a novel, more personalized, precise, and systematic, interdisciplinary approach in medicine. It involves the implementation of computational approach in medical concepts, research, and clinical practice, through iterative and feedback exchanges between clinicians, biologists, pharmacologists, bioinformaticians, and mathematicians.

**METABOLIC CHANGES IN ADPKD**

A broad range of metabolic pathways in ADPKD were described including increased glycolysis, reduced fatty acid oxidation (FAO) and altered glutamine utilization with defective mitochondrial function. We describe the current biological findings and how mathematical modeling applied to biological data can be useful to gain further insights into the metabolic alterations.

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**FIGURE 1** | The systems biology approach in ADPKD. Steps of the workflow for studying possible new therapeutic targets for ADPKD. The first step concerns the formulation of an initial hypothesis on possible changes in a metabolic network due to the disorder. In the second step, wet-lab experiments are employed to obtain omics data. The third step involves statistical analysis of the experimental data that will be used in the fourth phase to develop mathematical models for in-silico simulations. In the fifth step, in order to validate the initial hypothesis, the simulation results are mapped to the metabolic network. Finally, systems medicine is employed to propose new possible therapeutic targets.
### Table 1 | Characteristics of the included clinical trials.

| Drug                  | ClinicalTrials.gov Identifier | Publications indexed to study | Population                  | Outcome       |
|-----------------------|-------------------------------|-------------------------------|-----------------------------|---------------|
| Metformin             | NCT02656017                  | (19)                          | Adult 18 to 60 years        | Completed     |
| Pravastatin           | NCT00456365                  | (20-25)                       | Child, Adult 8 to 22 years  | Completed     |
| Pravastatin           | NCT03273413                  | (26)                          | Adult 25 to 60 years        | Recruiting   |
| Lisinopril, Telmisartan | NCT00283686           | (27-30)                       | Child, Adult 15 to 64 years | Completed     |
| Lisinopril, Telmisartan | NCT01885559               | (31)                          | Child, Adult 15 to 64 years | Completed     |
| Sirolimus             | NCT01223755                  | (32)                          | Adult, Older Adult 18 to 80 years | Completed     |
| Sirolimus             | NCT00346918                  | (33-37)                       | Adult 18 to 40 years        | Completed     |
| Rapamune              | NCT00286156                  |                               | Adult, Older Adult 18 to 75 years | Completed     |
| Everolimus            | NCT00414440                  | (38)                          | Adult 18 to 50 years        | Completed     |
| Rapamycin (sirolimus) | NCT00090309                  |                               | Adult, Older Adult 18 to 70 years | Terminated   |
| Lisinopril, Telmisartan | NCT01885559               | (31)                          | Child, Adult 15 to 64 years | Completed     |
| Pravastatin           | NCT00456365                  | (20-25)                       | Child, Adult 8 to 22 years  | Completed     |
| Lisinopril, Telmisartan | NCT00283686           | (27-30)                       | Child, Adult 15 to 64 Years | Completed     |
| Lisinopril, Telmisartan | NCT01885559               | (31)                          | Child, Adult 15 to 64 Years | Completed     |
| RGLS4326              | NCT04536688                  |                               | Adult, Older Adult 18 to 70 years | Recruiting   |
| Fasting/ketogenic*    | NCT04472624                  | (39)                          | Adult 18 to 60 years        | Active, not recruiting |
| Tolvaptan             | NCT00428948                  | (40-49)                       | Adult 18 to 50 years        | Completed     |

*Order indicated as in the review. * intervention/treatment.

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**Lactate as an Energy Source**

One of the common features of cystic cells derived from murine models of ADPKD is that they exhibit increased glucose uptake and lactate accumulation (9). In Pkd1<sup>−/−</sup> cells, glucose is converted to pyruvate which is then reduced to lactate instead of being transported into mitochondria (9, 12). Metabolic tracing with <sup>13</sup>C<sub>6</sub>-glucose by NMR revealed a significant increase in lactate production in cystic kidneys compared to age-matched controls derived from an aggressive model (Ksp-Cre:Pkd1<sup>flox/−</sup>) and a slowly progressive murine model (Tm-Cre:Pkd1<sup>flox/−</sup>) of ADPKD (9). Increased glucose uptake was also found in Pkd1 mutant mouse embryonic kidney (MEK) cells compared with Pkd1 WT MEK cells (15). Further, an upregulation of glycolytic markers was observed (9, 15, 16), and the glycolytic signature was confirmed in a subset of human microarray data and cystic kidneys derived from hypomorphic Pkd1 mutant mice (9, 12, 17). A recent study investigated the cross-sectional association of urinary metabolic biomarkers with ADPKD (18) severity among the TAME-PKD trial participants at baseline (19) (Please refer to Table 1 for mentioned clinical trials). This study is the first evidence of human urine samples supporting that increasing ADPKD severity is associated with a metabolic shift toward increased aerobic glycolysis. Other studies on the analysis of biomarkers in ADPKD should follow as to determine the ability to detect if a particular biomarker changes with time and if the correlations may be associated with cellular damage rather than metabolic changes per se. The enhanced glycolysis, the so-called Warburg effect reminiscent of cancer cells (49), contributes to the metabolic re-wiring in ADPKD and is considered a prominent feature of this disease.

**Mitochondrial Abnormalities and ER Cross-Talk**

Mitochondria are the central organelles for metabolism and are the key players for energy production in the cells. Mitochondrial...
respiration has been reported to be altered in Pkd1−/− proximal
tubule cells (11). Indeed, the investigators demonstrated that
oxygen levels regulate the localization and activity of PCI by a
prolyl hydroxylase domain-containing protein 3 ( EGLN3).
Furthermore, the study reported that the polycystins proteins
localize in the mitochondria-associated membranes (MAMs),
where they can mediate calcium signaling from the ER lumen
to the mitochondrial matrix (11). PC2 depletion alters calcium
transfer and decreases mitochondrial movement. This decrease
is thought to be the result of an increased expression of the
ER-mitochondrial tethering protein, mitofusin 2 ( MFN2), which
ultimately affect calcium-mediated mitochondrial bioenergetics
that contribute to cell proliferation (50). These findings
collectively suggest that the polycystins are important regulators
of mitochondrial function, but whether mitochondrial calcium
and dynamics are disrupted in PKD1 mutations warrants further
investigation. A cleaved carboxyl-terminal fragment of PC1
has been proposed to accumulate in mitochondria, indicating
that PC1 may directly modulate mitochondrial function (51).
However, how this cleavage is involved in cyst formation or
mitochondrial function is still unknown.

Morphologically, abnormal mitochondria were described
in cyst-lining cells in an ADPKD mouse model ( Ksp-
Cre: Pkd1fl/−/− ) and in rats (Han:SPRD Cy/+ ) (52). This
was accompanied by a reduction in mitochondrial number
suggesting a decrease in mitochondrial mass in the cystic
epithelia (52). In another study, the decrease in mitochondrial
mass was found to be correlated with a reduced activity of
mitochondrial respiratory enzymes and fragmentation of
the mitochondrial network in the cystic epithelia of Ksp-
Cre: Pkd1fl/−/− kidneys (53). The inhibition of mitochondrial
function was shown to be mediated by miR-17- Pparα axis (54).
The miR-17 inhibits OXPHOS and fatty acid oxidation, and its
suppression improves mitochondrial function of cyst epithelia
(54). PPARγ, another member of the PPAR subfamily has been
shown to be implicated in ADPKD pathogenesis. Moreover,
treatment with PPARγ agonist rescues proliferation and retards
cyst growth in rodent PKD models (55-57).

The master regulator of mitochondrial biogenesis, peroxisome
proliferator-activated receptor γ coactivator 1 α (PGC-1α),
was downregulated in kidney tissues from Pkd1fl/−/− and 7-
week-old Cy/+ rats (52). PPARA and its target genes were
shown to belong to a down regulated pathway in human
ADPKD cysts (17). Interestingly, the mitochondria from PC2
knock-down (KD) cyst-modeling cells exhibit altered metabolic
capacity and show a high degree of fragmentation, which they
have in common with Pkdi; however, an increased level in
PCG-1α was shown in PC2 KD cells (50). The reasons for
these different phenotypes between cells lacking PC1 and PC2
proteins still remains unclear, but the idea that mitochondria-
ER communication is a common mechanism that contributes to
altered metabolism in ADPKD is well-evidenced.

The major pathway for the degradation of fatty acids
inside the mitochondria is mitochondrial fatty acid β-oxidation
(FAO), which is the preferred energy source for highly
metabolic cells and can contribute to mitochondrial dysfunction.
Under physiological conditions, FAO provides more ATP than

Glutaminolysis and Amino Acid Metabolism

Glutamine is another important metabolite that sustains the
supply of carbons and nitrogen to fuel biosynthesis of the TCA
cycle metabolites (59). In standard culture, many cancer cell lines
fuel the TCA cycle using acetyl-CoA produced from glucose and
most of the anaeroplosis is supplied by glutamine (60). Glutamine
addiction in ADPKD was first reported in the study by Hwang
et al. In this study, a non-targeted metabolomics approach was
investigated in autosomal recessive polycystic kidney disease
(ARPKD) kidneys, and cpk (congenital polycystic kidneys) cells.
The results were then validated via targeted analysis with changes in
human ARPKD tissue and unaffected controls (61). Cpk cells
are addicted to glutamine, which is required for the production
of the oncometabolite glutamine-sourced 2-hydroxylglutarate (2-
HG) (61). Accumulation of 2-HG has been described in renal
cell carcinoma and it is thought to contribute to enhanced
proliferation through epigenetic modifications (62). This striking
parallel between cancer and ADPKD was firstly evidenced in this
study (61) and subsequently associated with profound metabolic
reprogramming that is found in tumors. However, ADPKD does
not predispose to malignancy and the reader can find further
details in the following review (63).

The glutamine addiction in ADPKD opened novel therapeutic
opportunities, and subsequent studies have shown that inhibition
of glutaminase 1 (GLS1) via BPTES or CB-839, tested in a
less aggressive model of cystic disease (Aqp2-Cre: Pkd1fl/−/−
and Pkhd1-Cre: Pkd1fl/−/− ) (64), affected proliferation and cyst
growth. The inhibition of glutamine with CB-839 resulted in
Glutamine revealed that Pkd1 gene, in cystic kidneys derived from Ksp-Cre: Pkd1<sup>flox/flox</sup>−/− mice, and in a subset of microarrays from humans and murine kidneys when compared to the corresponding controls (12). To evaluate the relevance of ASNS upregulation in PKD, silencing Asns in Pkd1−/− cells completely rescued the accumulation of α-ketoglutarate (α-KG), specifically reducing the contribution to glutamine-derived α-KG (12). One possible mechanism for the increased in ASNS is that asparagine synthesis supplies the availability of the TCA cycle intermediates and supplies the reduced nitrogen to maintain the pool of non-essential amino acid synthesis (66) but this needs to be investigated in ADPKD. In a metabolic profiling performed in children and young adults with ADPKD, Plasma samples were collected analysed with mass spectrometry (MS) and the most influential metabolite in ranked based area was asparagine, which was more than threefold higher in patients with ADPKD (67). In another study, glutamine addiction in PKD was investigated by an alternative glutamine pathway, via arginine synthetic enzyme argininosuccinate synthase1 (ASS1) directed to urea production (68). In both mouse models and human ADPKD tissue, the authors demonstrated that ASS1 expression was reduced and arginine depletion resulted in decreased cyst formation (68). Targeting arginine, via arginine-degrading enzymes, as used in clinical trials, and reviewed in Patil et al. (69), could be used to reduce circulating arginine in PKD. Collectively, these results suggest that ADPKD depends on glutamine for growth, and its inhibition reduces cells growth (12).

Glutamine is also a precursor for the major antioxidant system compromised of glutathione (GSH) and oxidized glutathione (GSSG) (70). In another study by Flowers et al., the investigators found that embryonic Pkd1 mCCD are also dependent on glutamine, which is required for glutathione synthesis (71). The changes in glutathione metabolism and the vulnerability of mutant PKD cells to reactive oxygen species (ROS) was found at a very early stage of PKD, which was reflected by increased 8-hydroxydeoxyguanosine (8-OHdG) that enhanced disease progression (72). The oxidative stress biomarker 8-OHdG was increased in the kidney cyst-lining cells from Ksp-Cre: Pkd1<sup>flox/flox</sup>−/− mice and Cy/+ rats, indicating that mitochondrial abnormalities also incite oxidative stress in these cells (52). There has been a link between oxidative stress and impaired endothelial-dependent relaxation (endothelial dysfunction) in progression of kidney injury with ADPKD, as observed in Nowak et al. (73) and Klawitter et al. (74). The PC1/PC2 complex is necessary for normal vasculature development, since it is required for endothelial cilia to sense fluid shear stress through complex biochemical cascades involving many factors including nitric oxide (NO) (74). NO is produced by endothelial cells and is a critical regulator of this balance, such that endothelial dysfunction is defined as reduced capacity for NO production (75). Endothelial dysfunction and decreased endothelial NO synthetase activity

cystic amelioration in Aqp2-Cre: Pkd1<sup>flox/flox</sup>−/− but not in Pkd1<sup>−/−</sup> mice. The investigators identified differences in signaling pathways due to mTORC1 and ERK activation, which might explain the likely failed inhibition, suggesting the upregulation of an adaptive metabolic pathway is able to bypass GLS1 inhibition. In another study by Podrini et al., tracing studies with [13C<sub>5</sub>N<sub>2</sub>] glutamine revealed that Pkd1<sup>−/−</sup> cells uptake more glutamine by utilizing the enzyme asparagine synthetase (ASNS) (12) (please refer to Figure 2). ASNS is a transamidase that converts aspartate into asparagine while deaminating glutamine to form glutamate (65). ASNS was found to be upregulated in cells lacking the Pkd1 gene, in cystic kidneys derived from Ksp-Cre: Pkd1<sup>flox/flox</sup>−/−, and in a subset of microarrays from humans and murine kidneys when compared to the corresponding controls (12). To evaluate the relevance of ASNS upregulation in PKD, silencing Asns in Pkd1−/− cells completely rescued the accumulation of α-ketoglutarate (α-KG), specifically reducing the contribution to glutamine-derived α-KG (12). One possible mechanism for the increased in ASNS is that asparagine synthesis supplies the availability of the TCA cycle intermediates and supplies the reduced nitrogen to maintain the pool of non-essential amino acid synthesis (66) but this needs to be investigated in ADPKD. In a metabolic profiling performed in children and young adults with ADPKD, Plasma samples were collected analysed with mass spectrometry (MS) and the most influential metabolite in ranked based area was asparagine, which was more than threefold higher in patients with ADPKD (67). In another study, glutamine addiction in PKD was investigated by an alternative glutamine pathway, via arginine synthetic enzyme argininosuccinate synthase1 (ASS1) directed to urea production (68). In both mouse models and human ADPKD tissue, the authors demonstrated that ASS1 expression was reduced and arginine depletion resulted in decreased cyst formation (68). Targeting arginine, via arginine-degrading enzymes, as used in clinical trials, and reviewed in Patil et al. (69), could be used to reduce circulating arginine in PKD. Collectively, these results suggest that ADPKD depends on glutamine for growth, and its inhibition reduces cells growth (12).

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![Figure 2](image-url)
are observed in patients (74) and young adults with ADPKD (73). Because statins have anti-inflammatory and anti-oxidant effects (76), there have been studies for the prevention of acute kidney injury after cardiac surgery (77) which predisposes to endothelial dysfunction. In a pediatric ADPKD trial, plasma levels of oxidative stress biomarkers declined during pravastatin treatment but not with placebo (78). There is also an ongoing clinical trial in adults (Pravastatin in adult phase 4, ClinicalTrials.gov Identifier: NCT03273413). However, a post hoc analysis was performed on adults in the HALT-PKD trials on statins resulting in no benefit from statin therapy (79). Whether oxidative stress is a driver of disease still needs further investigation.

**Mathematical Modeling of Metabolism: Application to ADPKD**

At the genome-scale level, metabolism is the result of thousands of compounds that are linked by reactions, which are continuously interacting with each other. The mathematical modeling process of metabolic networks can be broken in two steps. First, the stoichiometry of the system, which is the set of reactions representing the species connectivity, is established, and then rate laws are assigned to each reaction. For specific pathways, kinetic models based on sets of ordinary differential equations (ODEs) are broadly used to describe the metabolites' dynamic along time (80, 81). The foundation of this approach is the theory of molecular collisions that represents the basis of the mass action law. According to this theory, the algebraic expression for the ODEs depends on the reaction kinetics under consideration. There are several approaches for deriving kinetic functions: mass action kinetics (82), the Michaelis-Menten type and allosteric kinetics (83, 84), and the power-law approximation (85). The major drawback in using ODEs is the scarcity of experimental data on rate constants, especially at the genome-scale level. Genome-scale metabolic models require a large number of kinetic constants, many of which have not yet been experimentally measured. Moreover, the value of these parameters are subject to high uncertainty, resulting in erroneous simulations. Stoichiometric models address these problems by assuming that the changes in the compound concentrations are the result of the topological structure of the metabolic reactions. Starting from this assumption, by applying Flux Balance Analysis (FBA), a set of ODEs is transformed into a system of linear equations, and its rates can be obtained by solving a linear programming problem that optimizes specific objective functions (86). Further constraints are added to narrow the solution space. Although these types of models do not depend on the mathematical formalization of the rate laws, their analysis can lead to general results about the reactions properties. In one study (12), a computational systems-level approach based on FBA, called Differential Flux-balance Analysis (DFA) (87, 88), was employed to gather a broader understanding of the metabolic changes observed in Pkd1 mutant cells. By employing tracing experiments with $^{13}$C-labeled glucose, the investigators found an 1.6 fold increase in the uptake of glucose in Pkd1$^{−/−}$ cells compared to controls. This data was used as mathematical constraint for DFA. The simulations suggested that an increased uptake of glucose is able to drive a statistically significant change of the bioenergetics mechanisms, such as the TCA cycle, oxidative phosphorylation, the pentose phosphate pathway, and fatty acid synthesis and oxidation. The simulations have been followed by an extensive study of the observed metabolic reprogramming by integrating in vitro and in vivo analysis with metabolomics and transcriptomics data. In a subsequent work (89), DFA has been applied to test whether these alterations are causally interconnected or if they might occur simultaneously without interfering with each other. Particularly, DFA has been integrated with a Compounds-Pathways Enrichment Analysis to simulate the metabolic changes of bioenergetics pathways observed in Podrini et al. (12): (i) increased glucose uptake; (ii) increased mitochondrial glutamine uptake; (iii) increased fatty acids synthesis; (iv) reduced fatty acids oxidation. They found that simulations of these alterations result in important overall metabolic changes in pathways that are very similar in the different conditions, suggesting that they are all interconnected by key genes and specific chemical reactions. However, slightly different alterations were observed in the different simulations and only increased glycolysis appears to be the alteration that better recapitulates simultaneously all changes observed in Pkd1 mutant cells, while none of the other simulations can recapitulate the overall picture, suggesting an important possible causal link and directionality of the alterations observed. In addition, DFA showed that an increased flux of citrate from mitochondria to cytosol modulates the switch toward alternative bioenergetics pathways, hence, this metabolite could be one of the key factors in the regulation of the metabolic alterations observed in ADPKD.

Computational analysis may contribute to basic research on ADPKD, but the success of this approach will certainly require new modeling and simulation tools because our current understanding of how regulations are carried out is probably still missing several significant pieces. More experimental work is needed, and these experimental results must be incorporated in improved mathematical models, which would benefit from the creation of a common modeling framework that takes into account different entities, such as genes, proteins, and metabolites, and relationships, like metabolic reactions, interactions, regulations, and transports. Several problems and requirements arise toward this strategy, such as how to deal with incomplete information, how to manipulate large models, how to extract valuable information about the regulative mechanisms, how to analyze these models, and how to infer suitable models from experimental data.

**SYSTEMS BIOLOGY OF ADPKD**

The goal of Systems Biology is to try to understand global dynamics and properties of biological processes on the basis of microscopic factors depending on the scale of the phenomenon under study. The unifying framework is that biological systems are the result of a high number of interacting elements that are organized, at different levels of complexity, in functional networks. Particularly, the availability of high-throughput data
about multiple omics allows mapping cellular ADPKD processes at the level of genes, proteins, and metabolites.

The work of Song et al. (17) represents the first application of a Systems Biology approach in ADPKD. By employing microarray expression profiling from cysts of different size and from minimally cystic tissue from five PKD1 human polycystic kidneys, the authors identified that a cross-talk of specific signaling pathways modulates renal cyst expansion. They inferred downregulation of several gene sets associated with metabolic pathways like amino acids, fatty acids, urea cycle and ATP metabolism, and other kidney specific gene sets associated with ciliary functions, and renal cystic diseases. In contrast, sets of signaling genes resulted upregulated. The database developed by Song et al. still represents an invaluable resource for researchers interested in studying molecular pathobiology of ADPKD, and it was employed in several computational approaches related to ADPKD. This dataset has been analyzed in Rahimmanesh and Fatehi (90) by employing differentially expressed genes (DEGs) to construct protein-protein interactions (PPIs) networks. Enrichment analysis applied to protein clusters found in networks resulted in the detection of critical metabolic pathways such as oxidative phosphorylation and energy metabolism. The inferred networks also demonstrated that the interactions among proteins become more complex with the disease progression, while clustering revealed that EDN1, a vasoconstrictor that may promote tumor growth and for which serum levels have been associated with disease progression in ADPKD (91, 92), seems to be a seed gene in the early stage of the disorder. In addition, miRNAs and transcription factors enriched for dysregulated genes have been determined, such as hepatocyte nuclear factor alpha (HNF4A). In another study (93), the Song’s database has been integrated with a mice microarrays datasets (17, 94), and a set of DEGs was found in both the datasets. The downregulated genes were associated with carbon metabolism, fatty acids, and amino acid metabolism. Pathways involved in cancer, PI3K-Akt signaling pathway, focal adhesion, complement, and coagulation cascades resulted enriched for the upregulated DEGs. Starting from the DEGs, the authors filtered out a PPI network containing two significant modules. One module contained proteins related to mitochondrion, degradation of valine, lysine, leucine, fatty acids, ATP binding, and metabolism of carbon, propanoate, and tryptophan. The second one was associated with protein binding, extracellular region, extracellular space, platelet degranulation, extracellular exosome, plasma membrane, and aging (93).

With the aim of revealing the gene expression signatures related to progression of ADPKD, to infer signaling pathways associated with dysregulation of gene expression, and to study the roles of miRNAs in regulating the differences between normal and cystic kidneys, Pandey et al. investigated global microarray expression profiling (95). Kidneys from Pkd1<sup>−/−</sup> littermates at the embryonic ages 14 (E14.5) and 17 (E17.5) were investigated and even if mutants at E14.5 do not exhibit cysts, specific changes in gene expression related to signaling were observed. This result indicates that gene expression changes at E14.5 could provide an indication of signaling pathways that cause cysts rather than being a consequence of cyst formation. Most of the deregulated gene sets in Pkd1<sup>−/−</sup> mutants compared to WT at E14.5 and E17.5 were metabolic pathways and others involved in kidney development and regeneration. However, mutants at E17.5 days showed a broad change of genes associated with development, differentiation of specific nephron segments, and cyst formation and growth. Particularly, about 50% of the dysregulated pathways were in common with the deregulated ones observed in human ADPKD by Song et al. (17). In order to understand if miRNAs are directly involved in cyst formation and growth, miRNA microarrays using RNA from the kidneys of control and Ksp-Cre-Klfα<sup>fl/fl</sup> mice was performed (96). The main goal of this work was the identification of differentially expressed miRNAs. Patel and colleagues showed that the oncogenic miRNA cluster miR-17~92 is upregulated in a mouse model of PKD. This miRNA drives cyst epithelial proliferation and inhibits the post-transcriptional expression of Pkd1, Pkd2, and hepatocyte nuclear factor 1β (Hnf-1β). An integrated analysis of miRNA and RNA-sequencing was employed in (94) to investigate the miRNA profiles correlated with the severity of cyst development in two ADPKD models, kidney-specific Pkd1 or Pkd2 KO mice, at three time points, P1, P3, and P7. Woo et al. discovered differentially expressed miRNAs that were common in both mouse models. Particularly, a small subset of these miRNAs have been associated with cyotogenesis-related pathways. The role of miRNAs in ADPKD has also been recently studied in (97) by employing a transcriptional profiling of orthologous mouse models and cystic kidneys from humans with ADPKD. Lakhia et al. (97) showed that miRNAs represent molecular signals in the crosstalk between cyst microenvironment and immune cells. Particularly, miR-214 is upregulated in both mice and human PKD, while miR-214 deletion worsened cyst enlargement in the Pkd2 KO murine model of ADPKD and increased cyst-related inflammation. This result suggests that miR-214 is a protective signal against cyst development and formation.

Since it has been observed that the kinetics of cyst formation after Pkd1 inactivation in mice is sensitive to the age, specifically between days 12 and 14 (98), a network analysis has been applied to transcriptomics data from 36 mutants and 34 controls kidneys in order to study the changes during the P12-P14 interval that have such a drastic modifier effect in ADPKD (99). They inferred a cluster of DEGs enriched for several gene ontology categories related to metabolic pathways, development, cell differentiation, and anatomical structure morphogenesis. A gene ontology analysis suggested that metabolic pathways represent a key element in post-natal kidney development and the first step of cyst formation. The alteration of metabolic pathways has been confirmed by a metabolomics analysis of urine from 84 mice, which also showed an impairment of tyrosine metabolism, oxidative phosphorylation, purine metabolism, and aminoacyl-tRNA biosynthesis.

Finally, we would like to point out that long non-coding RNAs (lncRNAs) have emerged as epigenetic regulators of disorders and development. However, their role in ADPKD is not very clear yet. Therefore, RNA-sequencing analysis has been performed (100) with the aim to identify lncRNAs deregulated in ADPKD. Several lncRNAs were deregulated in two kidneys specific Pkd1 and Pkd2 mutant mice. One of these, Hoxb3os, regulates a network of genes and signaling pathways that are commonly
dysregulated in ADPKD. The ortholog HOXB3-AS1 has been found to be downregulated in cystic kidneys from ADPKD patients. Several genes were differentially expressed between Hoxb3os KO and control cells. Pathway analysis suggested that Hoxb3os regulates mTOR pathway and metabolism in mouse kidney ADPKD cells. Specifically, Hoxb3os could represent a negative regulator of mTORC1 activity and mitochondrial respiration, suggesting that re-expression of Hoxb3os could be used as a therapeutic strategy in ADPKD.

In conclusion, it is important to highlight that the Systems Biology tools developed in the last few years represent powerful methods to generate systematic views of the ADPKD, with the aim of exploring the comprehensive molecular mechanisms of this disease.

Emerging Therapeutics and Systems Medicine in ADPKD

The pathophysiology behind ADPKD is not completely understood, but a “threshold model” of cytogenesis is supported by previous studies (101–103). In this model, reduction of cellular levels of PC1 can influence cystic disease severity by a complexity of signaling pathways (104). However, re-establishing the levels of PC1 to restore normal cystic tissue remains an elusive approach. Targeting altered signaling pathways that drive cytogenesis (104) may be the preferred strategic approach to ameliorate disease progression in ADPKD. As a consequence of vasopressin V2-receptor activation, dysregulated signaling pathways in ADPKD such as increased cyclic adenosine monophosphate (cAMP) concentration, resulted in the identification of tolvaptan as a first treatment for ADPKD (105, 106). The activity of key energy metabolic enzymes was shown to be central in the metabolic shift of ADPKD. For example, AMP-activated protein kinase (AMPK) decreased (9, 107), while mTORC1 and ERK increased (108, 109). It was proposed that the decreased phosphorylation of AMPK is due to regulation of the liver kinase B1 (LKB1)/AMPK axis (9, 16). In another study, it was reported that the activation of the mitogen-activated protein kinase (MAPK)-ERK pathway was observed in an acute perinatal Pkd model; however, an inhibitor against ERK was not able to prevent cyst formation and may not be the basis for cyst expansion (110). Inhibitors of the mTORC1 complex, including sirolimus and everolimus, have been tested in five clinical trials. The results of the clinical trials showed no beneficial effects on the rate of change of total kidney volume (TKV) (111).

A novel therapeutic approach that targets mRNA expression is to utilize the action of an antisense oligonucleotide (ASO), a single-stranded deoxyribonucleotide that is complimentary to the mRNA target. The goal of the ASO is the down regulation of a specific molecular target (112–114). Therapeutic use of ASOs has been approved by the FDA for various pathologies (115). An ASO target strategy has been tested by chronic administration of mTORC1 ASO in Pkd2\textsuperscript{W25S−/−} mice, a homologous model of human PKD caused by a mutation in the Pkd2 gene, and resulted in improvement of the renal cystic phenotype (116). ADPKD also effects endothelial dysfunction (117) and circulating bioactive lipid mediators (118) which can contribute to metabolic disorders. From a clinical point of view, there is a higher incidence of new onset of diabetes mellitus after transplantation in recipients with ADPKD vs. other primary kidney diseases (119). To tackle a broad range of metabolic disorders in relation to vascular damage in ADPKD, clinical trials have been conducted with angiotensin-converting enzyme (ACE) inhibitors in ADPKD patients (31, 67). These trials have not been successful in decreasing renal function (120). Alternatively, the use of ASO against angiotensin (AGT) has been investigated in pre-clinical studies (116, 121, 122). The ASOs resulted in an improved cystic phenotype in the Pkd2\textsuperscript{W25S} model (123) and in the global (121) and in the tamoxifen-inducible knockout model of PKD (122). This suggests that targeting AGT with ASO may be more efficient than ACE inhibitors, and could open novel therapeutic opportunities for targeting abnormal signaling cyst formation in ADPKD.

The inhibition of mitochondrial function by anti-miR\textsubscript{17}-92 attenuates disease progression in ADPKD mouse models irrespective of the mutated gene (Pkd1 or Pkd2), the type of mutation (null or hypomorphic) or the dynamics of cyst growth (54). An anti-miR\textsubscript{17} oligonucleotide has proved to be efficient in reducing cyst formation in PKD mouse models (54, 124). Based on these results, anti-miR\textsubscript{17} oligonucleotides were designed and screened with favorable pharmacological properties (125). In fact, a phase 1 trial with an anti-miR\textsubscript{17} drug (RGLS4326) is currently being performed in ADPKD patients (ClinicalTrials.gov Identifier: NCT04536688). The miR\textsubscript{17} family is a promising drug target for ADPKD. Recent evidence shows that large clusters of transcriptional enhancers, super-enhancers (SE), undergo extensive remodeling during cytogenesis and that SE-associated transcripts are most enriched for metabolic genes in cystic cells (126). AMP deaminase 3 (AMPD3), a key enzyme involved in AMP metabolism and AMPK activation, was found to be SE-driven in cyst development. Inhibiting AMPD3 by pentostatin, or cyclic dependent kinase 7 (CDK7), one of the controlled metabolic target genes, decreased cell proliferation and apoptosis in ADPKD cells and delayed disease progression in tamoxifen-inducible Pkd1 mice with slowly progressing ADPKD (126). These findings suggest that CDK7-containing SEs are critical in ADPKD. As for therapy, inhibiting AMPD3, could be beneficial to target before disease progression and can provide a greater efficacy than targeting the downstream effectors in ADPKD (please refer to Figure 3 for an overview of therapeutic targets).

Recent preclinical data have generated promising results for targeting cellular bio-energetics in ADPKD. The high rate of glycolysis encountered in ADPKD has been targeted with glucose analog 2-DG in orthologous mouse models of ADPKD (9, 107, 127) with effective cyst amelioration. A phase 1 clinical trial has been designed with 2-DG in an ADPKD cohort (128). In addition, the dependence on glucose in cystic cells opened new venues to target disease by modifying diet for cystic amelioration. In fact, 2-DG can be compared to calorie restriction without reducing food intake given its capability to reduce metabolic rate addiction to glucose (129). Thus, ketogenic diets that decrease glucose availability have been tested in ADPKD, with the hypothesis that ketogenesis may
be beneficial in preventing cyst formation. This hypothesis was evaluated in an ADPKD rat model and in a spontaneous feline model of ADPKD, where a high ketogenic diet resulted in strong inhibition of PKD progression (39). The increase in ketosis was also investigated by acute fasting in mouse, rat, and feline models of PKD and significantly reduced renal cystic burden in all three models (39). Currently, there is an ongoing clinical trial on the short term induction of ketosis in ADPKD patients (ClinicalTrials.gov Identifier: NCT04472624) and also the patent-pending supplements are also being formulated and will be tested in the near future in clinical trials (130). Furthermore, prior to testing the ketogenic diet calorie restriction was found to reduce cyst growth (131, 132). There is growing evidence that diet manipulation can be a modifier of ADPKD disease and that the plethora of metabolic dysregulation observed in ADPKD may support the proliferating cyst epithelia. One possible mechanism is through the impact of one-carbon and methionine metabolism which generate the essential methyl groups for which altered nutrient utilization is facilitated. Ramalingam et al. found a higher methionine-S-adenosyl methionine (SAM) levels and methyltransferase-like 3 (Mettl3) upregulation in multiple ADPKD mouse models and that methionine-SAM supplementation or Mettl3 overexpression induced cyst growth (133). Dietary methionine restriction or Mettl3 ablation attenuated PKD, which may be regulated through epitranscriptomic mechanism through c-Myc and cAMP signaling activation (133). As for therapy, a vegan-based diet could also be used to ameliorate cyst formation, as methionine is found in meat and fish; alternatively, METTL3 inhibitors may also be targetable for ADPKD but warrants further investigation.

Tolvaptan, as already mentioned, is the unique drug that has been approved, in several countries, for the treatment of ADPKD (134). Frequent liver chemistry monitoring is suggested for patients using Tolvaptan, as it caused liver enzyme elevations in two pivotal Phase III clinical trials in ADPKD patients (42, 106). Taking into account this information, there is an urgent need for new compounds that can be used to stop the disorder progression.

A novel approach for identifying new drugs for ADPKD has been investigated by Malas et al. (14). Their approach combines integrated computational tools, repurposing analysis, cheminformatics data, and RNA sequencing of Pkd1 KO mice at different disease stages in order to identify new drugs for ADPKD. Results from this approach indicated that three compounds (i.e., Meclofenamic acid, Gamolenic acid, and Birinapant) are able to inhibit cyst swelling in vitro without toxicity, and could be further investigated for the treatment of ADPKD. Meclofenamic acid targets the aldo-ketoreductase family 1, which is involved in steroid metabolism (135), a pathway known to be a player in cyst development in a PKD rat model (136). Gamolenic acid has been selected based on PPARδ, a nuclear hormone receptor that controls a set of metabolic genes belonging to glucose metabolism, and fatty acid synthesis, storage, mobilization, and catabolism. Members of the PPAR family have been studied in preclinical trials for PKD (55, 137). Birinapant represents a second mitochondrial-derived activator of caspases (SMAC) mimic that modulates apoptosis (138). It has been found that the SMAC mimic GT13072 reduces PKD progression in a Pkd1 mouse model (139).

Lixivaptan, another vasopressin V2 receptor antagonist, was proposed for ADPKD treatment. This compound was initially developed for treating hyponetremia (140), and it has been evaluated by applying quantitative systems toxicology modeling (141). Their results showed that when compared to Tolvaptan, Lixivaptan had a significantly lower risk of hepatotoxicity at their respective doses for ADPKD treatment; therefore, it is a good candidate for continued study in the treatment of this disorder. The difference in the toxicity mechanisms between Tolvaptan and Lixivaptan demonstrates
the fact that even compounds within the same class can vary widely in terms of potential liability and potential mechanisms of toxicity.

Based on drug repositioning, Asawa et al. developed a high-throughput screening platform for ADPKD (13) that employs Pkd1-null postnatal cells of proximal tubule origin, Pkd1-null mouse embryonic kidney cells of collecting duct origin (MEK-null), and Pkd1−/− human kidney epithelial cells in order to evaluate a set of 8814 approved drugs and studied compounds principally having anti-neoplastic usage. It resulted in 155 compounds that reduced the viability of mutant cells but had minimal effects on wild-type cells. The investigators inferred that 109 of these compounds reduced cyst growth in Pkd1−/− cells cultured in a 3D matrix. Furthermore, with the aim of screening out possible therapies that failed to be translated in the treatment of the human disease, the platform has been extended by introducing the evaluation of the compounds in a panel of primary ADPKD and control human kidney epithelial cells. This step allowed the repurposing of 18 compounds that affected the viability of ADPKD cells with minimal effects on control human kidney epithelial cells. These compounds are: (i) Brincilib, Prexasertib, AHPN, and IMD-0354 which have been not yet been analyzed in the context of ADPKD; (ii) Cytarabine, Ancitabine, Clofarabine, Gemcitabine, Adefovir, and Topotecan, Belotecan, and 9-Methoxycamptothecin, which inhibit the proliferation of Pkd1−/− cells directly interfering with DNA replication; (iii) Combretastatin A-4, Leptomycin B, Delanzomib, CB30865, Cervistatin, and Pitavastatin which have been previously proposed as potential therapeutic options for ADPKD (142–144).

In short, given the complexity of the altered pathways in cyst-lining epithelia, and the availability of a broad range of potential targets, powerful strategies are needed to make faster advances in this field. Systems medicine and drug repurposing for the development of new treatments can help expedite the discovery of new effective treatments. The tools and the approaches presented in this section represent valuable platforms to repurpose chemical compounds and to identify drug targets for ADPKD treatment. All the presented studies discussed are of high relevance to ADPKD patients, as Tolvaptan, is the only approved therapy but has limited treatment options. The identified compounds represent a starting point for future efficacy studies employing relevant ex vivo and in vivo ADPKD models.

**USE OF OMICS DATA AND FUTURE THERAPEUTICS IN ADPKD**

Omic data are valuable tools for elucidating key genes and pathways associated with different aspects of ADPKD. However, one of the main drawbacks is the much of the profiling data from mouse models and patients has been published on public databases (the reader can find details about the most known datasets in Table 2), but it has not yet been exhaustively analyzed with the most appropriate computational approaches. Some points have to be carefully considered: (i) how samples are organized in different groups in order to follow a differential expression analysis; (ii) how results from a computational analysis are corrected for multiple comparisons in order to have strong statistics; and (iii) how to take into consideration that data from independent studies could be limited by the fact that they were derived from single cohort and by the samples heterogeneity.

It has also been observed that genome-scale profiling analysis can lack reproducibility because a large number of hypothesis tests are based on relatively small sample sizes. This problem can be solved by executing a meta-analysis based on the integration of different studies, which can increase statistical power and lead to more robust inferences (162, 163). However, the development of reliable tools to integrate different datasets remains a challenge due to low data comparability and consistency. A cross species meta-analysis was carried out by (164) with the aim of identifying conserved pathways that could be key targets for ADPKD therapy. Their analysis was rooted on mouse, rat, and human microarray datasets and revealed statistically significant enriched dysregulated pathways and processes conserved across all the species. Processes related to genomic instability, growth factors and hormones, protein localization in ER and insulin resistance promote cysts formation that is regulated by key pathways mediating apoptosis, cell proliferation, cell adhesion, and differentiation. In contrast, cysts progression seems to be a consequence of processes associated with cell adhesion and inflammation. In addition, a causative link among immune response pathways, signaling pathways, and cellular architecture pathways seems to affect cyst development.

Regarding the drug repositioning for the ADPKD, we would like to highlight that treatments and their targets are biased toward the most studied proteins and disorders. Future functional wet-experiments are needed to infer the actual contribution of each gene involved in cyst enlargement and disease progression. Further investigations will be facilitated by the development and the continuous update of the molecules database with drug-like properties.

Of note, there is a lack of knowledge about specific biomarkers, particularly in ADPKD pediatric patients. A recent work tried to identify plasma fingerprints in children and young adult patients with preserved kidney functions (67). The goal of the study was to identify markers and associated pathways involved in the progression of pediatric ADPKD over 36 months. The authors collected plasma samples from 58 young patients during a phase III clinical trial designed to test the effectiveness of pravastatin on slowing down cyst progression. In this study, metabolomics analysis revealed that metabolites deriving from the TCA cycle were associated with ADPKD progression. This is a set example of how metabolomics can offer a powerful tool for the prediction of the patient outcomes in clinical trials. However, there are also limited therapies focusing on treating ADPKD in early childhood where the benefits could be more likely. Screening of at-risk children is controversial and not recommended (1, 163) because of its potential to cause psychological harm. There is not yet a means of stratifying patients by rapid progressors of disease or not. To achieve this, it will be fundamental to start lifelong therapy at a later stage. The
TABLE 2 | A list of datasets related to ADPKD.

| Database | Dataset series | Experiment type | Organism | References |
|----------|----------------|-----------------|----------|------------|
| GEO | GSE35831 | Expression profiling by array | Homo Sapiens | (145) |
| GEO | GSE7869 | Expression profiling by array | Homo Sapiens | (17) |
| GEO | GSE9167 | Expression profiling by array | Mus Musculus | (98) |
| GEO | GSE24352 | Expression profiling by array | Mus Musculus | (95) |
| GEO | GSE32586 | Expression profiling by array | Mus Musculus | (99) |
| GEO | GSE19460 | Expression profiling by array | Rattus Norvegicus | (148) |
| GEO | GSE11500 | Expression profiling by array | Rattus Norvegicus | (147) |
| GEO | GSE13065 | Expression profiling by array | Rattus Norvegicus | (148) |
| GEO | GSE72554 | Expression profiling by array | Mus Musculus | (10) |
| GEO | GSE86509 | Expression profiling by high throughput sequencing | Mus Musculus | (94) |

| GEO | GSE121563 | Expression profiling by array | Mus Musculus | (12) |
| GEO | GSE13452 | Expression profiling by array | Mus Musculus | (149) |
| GEO | GSE2673 | Expression profiling by array | Mus Musculus | (150) |
| GEO | GSE23079 | Expression profiling by array | Rattus Norvegicus | (151) |
| GEO | GSE74451 | Expression profiling by array | Homo Sapiens | (152) |
| GEO | GSE75578 | Expression profiling by array | Rattus Norvegicus | (16) |
| GEO | GSE101308 | Expression profiling by array | Homo Sapiens | (153) |
| GEO | GSE108864 | Expression profiling by high throughput sequencing | Mus Musculus | (100) |
| GEO | GSE131277 | Expression profiling by high throughput sequencing | Mus Musculus | (145) |
| GEO | GSE134721 | Expression profiling by high throughput sequencing | Homo Sapiens Mus Musculus | (125) |
| GEO | GSE128524 | Expression profiling by high throughput sequencing | Genome binding/occupancy profiling by high throughput sequencing | Mus Musculus | (154) |
| GEO | GSE137945 | Expression profiling by high throughput sequencing | Mus Musculus | (67) |
| GEO | GSE89764 | Expression profiling by high throughput sequencing | Mus Musculus | (54) |
| GEO | GSE108864 | Expression profiling by high throughput sequencing | Mus Musculus | (103) |
| GEO | GSE141281 | Expression profiling by high throughput sequencing | Genome binding/occupancy profiling by high throughput sequencing | Mus Musculus | (126) |
| PRIDE | PXD001075 | Proteomics | Homo Sapiens | (155) |
| MetaboLights | MTBLS677 | Metabolic Tracing Experiments | Mus Musculus | (12) |
| Podrini et al. | Suppl. Data | Metabolomics | Mus Musculus | (12) |
| Podrini et al. | Suppl. Data | Lipidomics | Mus Musculus | (12) |
| Menezes et al. | Suppl. Data | Quantile-normalized microRNA expression data | Mus Musculus | (95) |
| ArrayExpress | E-MTAB-8086 | RNA-seq of coding RNA, compound treatment design, disease state design, genetic modification design | Mus Musculus | (14) |
| ArrayExpress | E-MTAB-4188 | Transcription profiling by array, disease state design | Homo sapiens | (156) |
| ArrayExpress | E-MTAB-6640 | RNA-seq of coding RNA, stimulus or stress design | Mus Musculus | (157) |
| ArrayExpress | E-MTAB-6641 | RNA-seq of coding RNA, stimulus or stress design | Mus Musculus | (157) |

Details about the different databases can be found in the following papers: GEO (159), MetaboLights (159), ArrayExpress (159), and Pride (161).

The main burden of ADPKD is that it does not occur until adulthood and some children are usually asymptomatic. This may, as a consequence, influence their life-style. There is a significant discrepancy regarding data obtained in adults vs. children as diagnostic criteria are highly variable, leading to incomparable studies and results (166). In children with ADPKD, EU guidelines recommends to stratify high risk patients and Tolvaptan is currently being tested in EU in children with ADPKD with high risk of rapid progression (167). The result of this study could form the basis for understanding the tolerability of tolvaptan in patients who still have a good renal function and not yet compromised. What remains to be evaluated is whether the treatment could be effective if initiated at an early stage, before cysts are formed, while the kidneys are still well-preserved.

Several effective therapies are under current investigation. It is unclear why defective PKD proteins alter signaling pathways involved in bio-energetics processes. Despite the gap of knowledge, current therapeutic interventions and clinical trials are ongoing with promising results. It is still debated whether the best strategy is to target one single pathway or to target
multiple pathways simultaneously. In the future, an increase in the application of less invasive therapies, such as dietary restrictions, may be proposed to patients because, as described above, certain dietary interventions have been successful in slowing the progression of ADPKD in animal models. For example, time-restricted feeding improved kidney function and reduced cytogenesis, and a ketogenic diet also demonstrated similar beneficial effects in ameliorating cytogenesis (39). Human trials are ongoing and this will be a stepping stone to pursue non-invasive therapies which can be exploited and refined with a customized diet. Although the results seem promising, the benefits of the PKD ketogenic diet in human are unpredictable and the risk of potential side effects, such as cardiovascular problems or impaired liver function, is unknown. It is known that nutritional aspects such as high fluid and low sodium diet are beneficial for improving cysts formation and are also able to restrict phosphorus and protein intake (168); however, this intervention is not prescribed in the presence of a severe renal function. At present, perhaps the important aspect to consider would be to follow a balanced diet and to avoid a large amount of refined carbohydrates. Further impact studies are needed on this issue as to address the benefits of non-invasive therapies in ADPKD.

**AUTHOR CONTRIBUTIONS**

CP and RP wrote the manuscript, reviewed and edited the manuscript before submission. Both authors contributed to the article and approved the submitted version.

**FUNDING**

The work on metabolism and PKD was funded by the Italian Ministry of Health (GR-2016-02364851 to CP and RP).

**ACKNOWLEDGMENTS**

The authors are grateful to other members of the laboratory of Dr. A. Boletta for helpful suggestions.

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Conflict of Interest: CP is a co-inventor on the use of metabolic inhibitors in Polycystic Kidney Disease. The remaining author declares that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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