Together, we made significant strides toward finding treatments and improving the lives of those impacted by PKD.

THANK YOU FOR AN INCREDIBLE YEAR!

Your generous donations and volunteerism make our work possible. Read on to learn more about the impact you made during fiscal year 2017 (FY17).

OUR MISSION

We give hope. We fund researchers, support patients and build a community for all affected by polycystic kidney disease (PKD).
RESEARCH

Since our founding in 1982, we have invested more than $42 million in research, clinical and scientific grants, as well as fellowships and scientific meetings on an international scale. This makes us the largest private funder of PKD research in the United States.

Research Funding

Our Research Grant Program funds critical research to increase understanding of the genetic and pathological processes involved in PKD and to accelerate the development of potential treatments. PKD Foundation Fellowships build momentum in PKD research by recognizing early-career scientists whose achievements and potential identify them as rising stars in the field. In FY17, you helped us continue to fund 15 grant recipients and finish funding five research fellows.

Scientific Meetings

Collaboration among PKD researchers is imperative to make advancements in research. In FY17, we sponsored and attended 5 scientific meetings that brought PKD experts together.

CTAP

CLINICAL TRIAL AWARENESS PROGRAM

Clinical studies are a critical and required step to develop new therapies. Our Clinical Trial Awareness Program (CTAP) creates awareness among patients and families to speed up recruitment and increase participation in clinical studies. The program focuses on educating patients about current studies so they can make informed decisions about their participation. To simplify the process of finding clinical studies for PKD patients in their geographic areas, we send Accelerating Clinical Trials (ACT) Alert emails about studies that are being conducted.

In FY17, we sent 10 ACT Alerts in the interest of getting patients involved in finding treatments. These included recruiting patients to join an ARPKD database, a PKD biomarker research study, a curcumin study in children and young adults with ADPKD, and studies surrounding three of five notable potential treatments: tesevatinib, pioglitazone and metformin.
SUPPORT

A major part of our mission is to support patients and build a community for all affected by PKD. We aim to provide all patients and caregivers with the outreach and resources they need to live with this disease and understand that they are not alone in their journey.

No one should face this disease alone.

Local Chapters

Our Chapters, and the volunteers who lead them, are the backbone of the PKD community. They are comprised of PKD patients, family members and friends who want to learn, connect and take action with other committed individuals. Chapters regularly hold events to raise awareness for PKD and offer education and support. They reinforce our belief that no one has to face this disease alone.

In FY17, Chapters collectively hosted over 100 education and support meetings for their local communities.

Learn about our Chapters.
pkdcure.org/chapters
EDUCATION

Our education resources aim to help PKD patients fully understand their disease, live their best lives and empower them to educate others.

Webinars

Webinar Wednesdays, our monthly education webinars, highlight targeted topics in PKD science and research. From basic kidney anatomy to Q&As with Scientific Advisory Committee members, these are a go-to resource for PKD knowledge and education.

Online Education Conference

On March 25, 2017, PKD community members all over the country joined their local Chapters to participate in our exclusive, online education conference “What’s new in PKD: A day of learning.” The conference was organized in an effort to offer remote learning opportunities that increase accessibility to information without limiting patients’ sense of togetherness. The event featured PKD experts from Mayo Clinic in Rochester, Minn., who discussed the kidney transplantation process.

Watch the presentation on our website.

pkdcure.org/resource/whats-new-pkd-2017

Watch the webinars from FY17 on a variety of topics.

pkdcure.org/webinars

What’s New in PKD: A Day of Learning

31 CHAPTERS PARTICIPATED

28 DIFFERENT LOCATIONS

6 KIDNEY EXPERT SPEAKERS

Speakers Tammy Kozel, Dr. Patrick Dean, Margo Vandrovec, Erin Hamilton, Lisa King, Dr. Naim Issa and emcee Dr. Peter Harris
ADVOCACY

By getting involved in advocacy efforts, you can educate your elected officials about how they can help fight PKD. Constituent voices are vital to increasing PKD research dollars to bring treatments and therapies to patients, and providing the best care for those who are reliant on dialysis or a kidney transplant to stay alive. We encourage people with PKD and those that care about them to meet with their local congressional representatives because just one personal story can make a significant difference.

Advocacy Alerts

Our Advocacy Alerts empower PKD patients and members of the community to raise their voices when important legislation hangs in the balance on the Hill. Prepared call scripts and email templates make taking action quick and easy.

Just a few of the important pieces of legislation our PKD community had an impact on:

**HEALTH CARE**

When the new administration pushed the American Health Care Act as a repeal-and-replace option for the Affordable Care Act, PKD patients raised their voices for affordable insurance coverage for people with pre-existing conditions.

**21ST CENTURY CURES**

Signed into law on Dec. 13, 2016, 21st Century Cures provides $4.8 billion in new funding to the National Institutes of Health (NIH) and assures that patient voices will be included in the drug development process for treatments and therapies.

Taking action in FY17

WE SENT

10
ADVOCACY ALERTS

653
CONSTITUENTS CALLED THEIR MEMBERS OF CONGRESS

3,584
CONSTITUENTS EMAILED THEIR MEMBERS OF CONGRESS

Learn how you can advocate and receive advocacy email alerts.

pkdcure.org/advocate
Chapter and individual fundraising events are critical to raising the money that allows us to continue working towards our mission. Led and organized by a wide variety of volunteers throughout the country, we could not continue to build the funds that come from these events without our PKD community.

Fundraising Events

The top five fundraisers of FY17

Memphis/Liberty Mutual Driving Home a Cure
$118,051

Miami Kidney Casino
$115,461

Celtic Marketing Golf Tournament
$41,510

Sacramento Corks for a Cure
$21,575

Ryan’s Reason Putting Fore PKD
$18,000

FY17 Fundraisers

76
TOTAL ORGANIZERS
$304,586
TOTAL AMOUNT RAISED

Learn more about individual events.
pkdcure.org/get-involved

Sacramento Corks for a Cure attendees
Walk for PKD

The Walk for PKD is our signature fundraising and public awareness event, and the largest gathering of PKD patients and supporters. Since 2000, the Walk for PKD has raised more than $28 million. This family-oriented event creates a sense of community and provides a unique opportunity to support and honor loved ones.

### TOP 10 Walk teams

<table>
<thead>
<tr>
<th>Team Name</th>
<th>City</th>
<th>Raised</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bost Bunch</td>
<td>Pittsburgh</td>
<td>$32,030</td>
</tr>
<tr>
<td>Ivan’s Investors for a PKD Cure</td>
<td>Tampa</td>
<td>$26,266</td>
</tr>
<tr>
<td>Team Odyssey</td>
<td>New York City</td>
<td>$24,200</td>
</tr>
<tr>
<td>Going for Geigers</td>
<td>Chicago</td>
<td>$21,250</td>
</tr>
<tr>
<td>Mike &amp; Poppy’s All-Stars</td>
<td>New York City</td>
<td>$18,770</td>
</tr>
<tr>
<td>Support a Cystah</td>
<td>Seattle</td>
<td>$18,290</td>
</tr>
<tr>
<td>Team Rechter</td>
<td>South Florida</td>
<td>$18,083</td>
</tr>
<tr>
<td>Team Pinnacle</td>
<td>Nashville</td>
<td>$17,610</td>
</tr>
<tr>
<td>Three Peas in a Pod</td>
<td>New York City</td>
<td>$16,770</td>
</tr>
<tr>
<td>Team OC</td>
<td>Twin Cities</td>
<td>$14,700</td>
</tr>
</tbody>
</table>

### TOP 10 Walk events

<table>
<thead>
<tr>
<th>City</th>
<th>Raised</th>
</tr>
</thead>
<tbody>
<tr>
<td>Boston</td>
<td>$111,168</td>
</tr>
<tr>
<td>New York City</td>
<td>$110,692</td>
</tr>
<tr>
<td>North Texas</td>
<td>$79,076</td>
</tr>
<tr>
<td>Nashville</td>
<td>$64,877</td>
</tr>
<tr>
<td>Kansas City</td>
<td>$63,585</td>
</tr>
<tr>
<td>Chicago</td>
<td>$62,058</td>
</tr>
<tr>
<td>San Antonio</td>
<td>$56,075</td>
</tr>
<tr>
<td>Pittsburgh</td>
<td>$48,977</td>
</tr>
<tr>
<td>Tampa</td>
<td>$47,180</td>
</tr>
<tr>
<td>Los Angeles</td>
<td>$46,599</td>
</tr>
</tbody>
</table>

Since 2000, the Walk for PKD has raised more than $28 million.

51 WALKS

TOTAL RAISED: $1,688,050

MORE THAN 10,000 PARTICIPANTS

1,569 WALK VOLUNTEERS

Learn more about Walk for PKD.

walkforpkd.org

Las Vegas Walk for PKD participants
FINANCIALS
1 JULY 2016 – 30 JUNE 2017

Where your money went

Where our money came from

INDIVIDUALS
$4,008,670

CORPORATIONS
$1,010,153

FOUNDATIONS
$742,292

BEQUESTS
$130,375

TOTAL DONATIONS
$5,891,490

Annual revenues
FY15
$7,925,685*

FY16
$6,680,617

FY17
$6,424,607

Financial position
ASSETS
$5,218,772

LIABILITIES
$503,054

NET ASSETS
$4,715,718

* Includes a single bequest of $1 million

RESEARCH
$2,184,499

DEVELOPMENT
$763,596

AWARENESS & ADVOCACY
$1,354,957

EDUCATION & SUPPORT
$1,273,991

ADMINISTRATIVE
$971,301