



**PKD FOUNDATION**  
Polycystic Kidney Disease

## PKD Foundation Work Samples of Logo in Use

### Exhibit B

2. Print ad 2014
3. Print ad 2015
4. PKD Banner
5. PKD Folder
6. Information Packet Contents  
Collage
7. About PKD
8. Promotional Items with New Logo



# Help Us Bring an End to PKD.



Polycystic kidney disease (PKD) is one of the most common genetic diseases – it often devastates entire families. It is the fourth leading cause of kidney failure and there is no treatment or cure.

**The PKD Foundation leads the fight** against PKD by funding research to find treatments and a cure, and providing information and support to people affected by PKD.

Help us bring an end to this life-altering and deadly disease...please join us as we unite to fight.

**Help us:  
Fund research  
Educate and inform  
Raise awareness**



**PKD FOUNDATION**  
Polycystic Kidney Disease

**Visit [pkdcure.org](http://pkdcure.org).**





# PKD FOUNDATION

Polycystic Kidney Disease

Print Ad 2015

1-800-pkd-cure  
[www.pkdcure.org](http://www.pkdcure.org)

Polycystic kidney disease (PKD) is one of the most common, life-threatening genetic diseases. Fluid-filled cysts develop and enlarge in both kidneys, eventually leading to kidney failure.



The Foundation is the only organization in the U.S. solely dedicated to finding treatments and a cure for PKD. Our vision is that one day, no one will suffer the full effects of PKD.





# Help Us Bring an End to PKD.

Polycystic kidney disease (PKD) is one of the most common genetic diseases – it often devastates entire families. It is the fourth leading cause of kidney failure and there is no treatment or cure. Once a person has kidney failure, dialysis or a transplant are their only options.

**The PKD Foundation leads the fight** against PKD by funding research to find treatments and a cure, and providing information and support to people affected by PKD.

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Polycystic Kidney Disease

**Visit [pkdcure.org](http://pkdcure.org).**





## MISSION

Our Mission: Promote programs of research, advocacy, education, support and awareness in order to discover treatments and a cure for polycystic kidney disease and improve the lives of all it affects.

## VISION

Our Vision: One day, no one will suffer the full effects of polycystic kidney disease.



**PKD FOUNDATION**  
Polycystic Kidney Disease

8330 Ward Parkway, Suite 510  
Kansas City, MO 64114  
1.800.PKD.CURE

[pkdcure.org](http://pkdcure.org)



**PKD FOUNDATION**  
Polycystic Kidney Disease



**PKD FOUNDATION**  
Polycystic Kidney Disease

**John Smith**  
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johns@pkdcure.org | pkdcure.org

**pkdcure.org**

**POLYCYSTIC KIDNEY DISEASE (PKD)**  
**is one of the most common, life-**  
**threatening genetic diseases. The PKD**  
**Foundation funds research to find**  
**treatments and a cure, and provides**  
**information and support to people**  
**affected by PKD.**



## PKD FOUNDATION CHAPTER NETWORK



### What is a Chapter?

Chapters are groups of volunteers - PKD patients, family members and friends just like you - who want to learn, connect and take action with other concerned individuals. In fact, chapters have been forming to fight polycystic kidney disease since the 1980s. Finding research, telling others about the disease and providing one another with help and hope - it's a true sense of community.

### About Our Chapters

Each year, Chapter volunteers in more than 30 states across the United States hold events to raise awareness of PKD and money for the PKD Foundation's programs and services, as well as offer education and support. These volunteers bring to life the mission of the organization by ensuring that no one faces this disease alone.

### Chapter activities focus on three key areas:

- **Education and Support:** Chapters host education meetings throughout the year, bringing in medical experts to discuss diagnosis, management, treatment issues and other important topics, and promote educational opportunities offered by the PKD Foundation. Chapters provide opportunities for patients to connect and share, helping each other cope with the emotional aspects of a PKD diagnosis, while finding the local resources they need.
- **Event Support:** Regular Chapter fundraising events include the Walk for PKD, as well as other PKD awareness events at garage sales, lemonade stands and on the sidelines of sports events. Additionally, Chapter support programs are also for PKD, workplace giving and volunteerism. Chapter volunteers also help us identify individuals, organizations and foundations interested in financially supporting the work of the PKD Foundation.
- **Advocacy:** Chapter volunteers help raise awareness of PKD by exhibiting at local health fairs, speaking at community events and participating in the Walk for PKD.

Each Chapter aims to have four key individuals: a Chapter Coordinator, a Walk for PKD Coordinator, a Fundraising Coordinator and a Volunteer Coordinator.

For more information or to connect to a Chapter near you, please visit [pkdcure.org/chapters](http://pkdcure.org/chapters) or call 1-800-PKD-CURE (753-2873).



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## PKD PATIENT HANDBOOK

Understanding and living with autosomal  
dominant polycystic kidney disease



**PKD FOUNDATION**  
Polycystic Kidney Disease

## PKD Progress

A magazine to inform and educate PKD patients and families,  
Foundation supporters, health professionals and researchers

Spring 2015

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PKD FOUNDATION'S NEW  
CHIEF SCIENTIFIC OFFICER

4 MAYO CLINIC'S ADPKD  
MUTATION DATABASE  
PROVIDES INVALUABLE  
INFORMATION ABOUT PKD

8 JINARC™ (TOLVAPTAN)  
APPROVED IN CANADA AND  
RECOMMENDED IN EUROPE

9 ST. AUBINS REMEMBER  
THEIR DAUGHTER THROUGH  
HELPING OTHERS



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### ABOUT PKD

#### PKD FACTS

Polycystic kidney disease (PKD) is one of the most common, life-threatening genetic diseases. It affects thousands in America and millions worldwide, who are in urgent need of treatment and a cure.

• In autosomal dominant PKD (ADPKD), fluid-filled cysts develop and enlarge in both kidneys, eventually leading to kidney failure.

• PKD is the fourth leading cause of kidney failure. More than 50 percent of people with PKD will develop kidney failure by age 50.

• Once a person has kidney failure, dialysis or a transplant are the only options to treat the damage the disease has caused.

• It is a painful disease that impacts quality of life. The average size of a normal kidney is a human fist. Polycystic kidneys can get much larger, some growing as large as a football, and weighing up to 30 pounds each.

• Parents have a 50 percent chance of passing the disease to each of their children. Unlike some genetic diseases, it does not skip a generation, because it can be passed from generation to generation. PKD often affects many people in one family.

• Approximately 10 percent of the people diagnosed with PKD have no family history of the disease, with PKD developing as a spontaneous (new) mutation. Once they have it, there is a 50 percent chance of passing it on to each of their children.

• PKD equally affects people of all ages, genders, nationalities, geographic locations and income levels.

#### Symptoms of PKD

People living with PKD can experience these common symptoms:

- High blood pressure (hypertension)
- Frequent urinary tract infections
- Blood in urine (hematuria)
- Protein in urine (proteinuria)
- Abnormal kidney enlargement
- Back/flank pain
- Kidney stones
- Enlarged kidneys
- Depression and anxiety (due to stress and emotional impact)

#### ADPKD FACTS

Autosomal recessive polycystic kidney disease (ARPKD) is a relatively rare form of PKD, affecting approximately 1 in 20,000 children. It often causes death in the first month of life.

• For ARPKD children who survive the newborn period (about 50 percent), approximately one-third will need dialysis or transplant by age 10.

• Previously thought to be a fatal condition, the prognosis for children with ARPKD has improved dramatically. Twenty years ago, only half of the children born with the disease survived to their 10th birthday, but now that percentage has increased to 85 percent.



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# ABOUT THE PKD FOUNDATION

The PKD Foundation is the only organization in the U.S. solely dedicated to finding treatments and a cure for PKD to improve the lives of those it affects. We do this through promoting research, education, advocacy, support and awareness on a national level, along with direct services to local communities across the country.

Our vision is that one day no one will suffer the full effects of PKD.

## Research

Since its founding in 1982, the PKD Foundation has invested more than \$38 million in research, clinical and scientific grants, as well as fellowships and scientific meetings, making us the second largest funder of PKD research after the National Institutes of Health (NIH). This has led to new discoveries about PKD, including identifying the genes responsible for PKD, which enables researchers to investigate possible treatments.

## Education

The Foundation provides in-depth resources and education about living with PKD to empower people to manage their health. Offerings include webinars, videos, a multi-faceted website ([pkdcure.org/learn](http://pkdcure.org/learn)), online communities and print materials. The biannual PKD National Convention is the Foundation's largest education event.

## Advocacy

The Foundation plays a key role in legislative advocacy to support PKD-related initiatives. The PKD Advocacy Action Center ([pkdcure.org/advocate](http://pkdcure.org/advocate)) provides resources and ways for people to advocate, and the Foundation sends Advocacy Alerts for legislative and public policy issues impacting PKD patients and families.

## Support

The PKD Connection is a blog for people to be inspired and empowered, and to connect with others affected by PKD. Discussion boards provide a forum for PKD patients, family and friends to make connections, ask questions and share stories and experiences.

## Awareness

The Foundation raises awareness through marketing and public relations so people know what PKD is, about the Foundation and how to donate. Marketing materials include PKD Progress magazine and PKDnews monthly email newsletter. Voices of PKD ([pkdcure.org/voicesofpkd](http://pkdcure.org/voicesofpkd)) features testimonials and stories about people's experiences with the disease.

## Walk for PKD

The Walk for PKD ([walkforpkd.org](http://walkforpkd.org)) is the Foundation's signature event to raise funds and awareness. Walks take place in more than 50 cities across the nation each year, with more than 11,000 walkers. The event has raised nearly \$24 million since 2000.

## Chapters

The Foundation provides local services through its more than 60 volunteer-run Chapters across the country. These volunteers bring to life the mission of the organization by ensuring that no one faces this disease alone.

- ▶ Education meetings provide valuable information from health care professionals, while support activities range from lending a listening ear to organized group outings.
- ▶ Chapters raise funds for the Foundation through events like the Walk for PKD and Cocktails for a Cure.

## How You Can Help

- ▶ Donate: The research, advocacy, education, support and awareness the PKD Foundation promotes would not be possible without you.
- ▶ Get involved: Volunteer, advocate, join a Chapter and spread the word.

## Join

- ▶ [walkforpkd.org](http://walkforpkd.org)
- ▶ [pkdcure.org/chapters](http://pkdcure.org/chapters)

## Connect

- ▶ Visit [pkdcure.org](http://pkdcure.org).
- ▶ For parents of children with ARPKD or ADPKD, email [pkdparents@pkdcure.org](mailto:pkdparents@pkdcure.org).
- ▶ Call 1.800.PKD.CURE (753.2873).
- ▶ Follow us on Facebook and Twitter (@PKDFoundation).
- ▶ Visit Voices of PKD for stories about people affected by PKD: [pkdcure.org/voicesofpkd](http://pkdcure.org/voicesofpkd).







## ABOUT PKD

### PKD Facts

Polycystic kidney disease (PKD) is one of the most common, life-threatening genetic diseases. It affects thousands in America and millions worldwide, who are in urgent need of treatments and a cure.

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- ▶ PKD is the fourth leading cause of kidney failure. More than 50 percent of people with PKD will develop kidney failure by age 50.
- ▶ Once a person has kidney failure, dialysis or a transplant are the only options to treat the damage the disease has caused.
- ▶ It is a painful disease that impacts quality of life. The average size of a normal kidney is a human fist. Polycystic kidneys can get much larger, some growing as large as a football, and weighing up to 30 pounds each.
- ▶ Parents have a 50 percent chance of passing the disease to each of their children. Unlike some genetic diseases, it does not skip a generation. Because it can be passed from generation to generation, PKD often affects many people in one family.
- ▶ Approximately 10 percent of the people diagnosed with PKD have no family history of the disease, with PKD developing as a spontaneous (new) mutation. Once they have it, there is a 50 percent chance of passing it on to each of their children.
- ▶ PKD equally affects people of all races, genders, nationalities, geographic locations and income levels.

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People living with PKD can experience these common symptoms:

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- ▶ Blood in urine (hematuria)
- ▶ Protein in urine (proteinuria)
- ▶ Mitral valve prolapse
- ▶ Hernia
- ▶ Back/flank pain
- ▶ Kidney stones
- ▶ Enlarged kidneys
- ▶ Depression and anxiety (due to stress and emotional impact)

### ARPKD Facts

Autosomal recessive polycystic kidney disease (ARPKD) is a relatively rare form of PKD, affecting approximately 1 in 20,000 children. It often causes death in the first month of life.

- ▶ For ARPKD children who survive the newborn period (about 70 percent), approximately one-third will need dialysis or transplantation by age 10.
- ▶ Previously thought to be a fatal condition, the prognosis for children with ARPKD has improved dramatically. Twenty years ago, only half of the children born with the disease survived to their 10th birthday, but now that percentage has increased to 85 percent.







**PKD FOUNDATION**

Polycystic Kidney Disease

8. Promotional  
Items  
with New Logo



CART

0 Items

CHECKOUT

[My Account](#)

[Home](#)

[Apparel](#)

[Business](#)

[Promotional](#)

[Drinkware](#)

[View All](#)

***Thank You For Visiting The PKD Foundation Online Store!***  
*By purchasing items from the PKD Store, you are helping to raise awareness in order to discover treatments and a cure for polycystic kidney disease.*





## Promotional

Sort Results By:

Creation Date(newest first) ▼



NEW

Awareness Ribbon  
#PKD2033  
On Hand: 35  
\$2.25



NEW

Lapel Pin 1 1/4"  
#PKD2031  
On Hand: 66  
\$3.70



NEW

Embroider Patch  
#PKD2040  
On Hand: 89  
\$3.50



NEW

Twist Silicone Bracelet  
#PKD2035  
\$1.25



Auto Open  
Compact Umbrella  
#PKD3302  
On Hand: 10  
\$8.99



Power Bank Charger  
#PKD4510  
On Hand: 94  
\$11.75



Color Pop Journal Book  
#PKD4100  
On Hand: 46  
\$8.25



Temporary Tattoo  
1.5" x 1.5" - 100 PK  
#PKD2028  
On Hand: 2  
\$20.26



Stylus Ballpoint Pen  
Silver/Purple - 10 PK  
#PKD4006  
On Hand: 57  
\$8.16



END PKD Awareness  
Bracelet - Teal  
#PKD2030  
On Hand: 74  
\$1.32



4" Round Static Cling  
#PKD2025  
On Hand: 103  
\$3.16



2 Color Static Cling  
4" Round  
#PKD2026  
On Hand: 144  
\$2.11

