

Keep hope within reach

connect.pkdcure.org · 844.PKD.HOPE · pkdconnect@pkdcure.org



PKDCONNECT

A Community of Care

A service of the PKD Foundation



PKD FOUNDATION
Polycystic Kidney Disease

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PKDCONNECT

A Community of Care



A Community of Family

Friends

Patients

Parents

Volunteers

Advocates

Professionals

Mentors

Resources

Care



**With PKD Connect,
no one faces
PKD alone.**

PKD Hope Line

From the moment of diagnosis on, there is no end to the questions patients, family and supporters have about PKD. We created the PKD Hope Line to provide answers and connections to the people and resources who can help put your mind at ease.

**Call 844.PKD.HOPE (844.753.4673)
Monday through Friday,
8 a.m. to 5 p.m. Central Time.**



Peer Mentor Program

If you, a family member or friend are impacted by PKD, sometimes you need advice, encouragement or a shoulder to lean on from someone who knows firsthand what you're going through. That's the purpose of the PKD Connect Peer Mentor Program.

Whether you or your loved one has just been diagnosed or is getting ready for dialysis or a kidney transplant, PKD Connect mentors are here to listen to concerns and share their own experiences and perspectives. PKD Connect mentors are carefully selected and trained to provide support, guidance and resources to help you manage living with PKD.

ADPKD Patient Registry

The purpose of the ADPKD Registry is to create a patient network of individuals with autosomal dominant polycystic kidney disease (ADPKD). Participants are asked to answer surveys online that can inform new research to improve ADPKD patient outcomes and quality of life, learn more about the patient journey, and discover unmet medical needs. ADPKD Registry keeps information in one place making it easier for researchers to utilize Registry information while protecting the privacy of those who take part.

Navigating a PKD diagnosis

A polycystic kidney disease (PKD) diagnosis can seem overwhelming. Questions flood your mind. Your self-esteem may suffer. Relationships can become strained. You may be filled with doubt and a feeling of isolation. But with PKD Connect, you're not alone. **Because hope is always within reach.**

Disease Management Resources

As a patient or caregiver of someone with PKD, managing the disease and maintaining a high quality of life are two of your top priorities. From nutrition tips and pain management information to kidney replacement options, we have the resources to help you throughout your PKD journey.

As we continue to enhance our resource library, we remain committed to include tools and education materials to assist patients, parents, family members and caregivers as they navigate managing PKD. Because the more you know, the better you can take care of yourself or your loved one.

Please be mindful that the PKD Hope Line is an information and support line, and is not staffed by medical professionals. The PKD Foundation does not offer medical advice.

