



Polycystic Kidney Disease (PKD)

Causes, Symptoms, Diagnosis, and Treatment



## **Definition of PKD**

- A genetic disorder causing fluid-filled cysts in the kidneys
- Can lead to kidney failure and other complications







### **Types of PKD**

- Autosomal Dominant PKD (ADPKD)
- More common
- Symptoms appear in adulthood
- Autosomal Recessive PKD (ARPKD)
- Rarer but more severe
- Symptoms appear in infancy or childhood





#### **Causes and Risk Factors**

#### **Genetic Mutations**

- ADPKD: PKD1 and PKD2 gene mutations
- ARPKD: PKHD1 gene mutation

Hereditary Nature – Runs in families

Other Risk Factors:
High blood pressure,
lifestyle factors







#### Slide 5: Symptoms of PKD

- High blood pressure
- Back or side pain
- Blood in urine
- Frequent kidney infections
- Enlarged kidneys
- Kidney stones









## **Complications of PKD**

- Chronic kidney disease (CKD) → Kidney failure
- Liver cysts
- Heart valve abnormalities
- Aneurysms (brain blood vessel swelling)
- Pregnancy complications





## Diagnosis of PKD

- Imaging Tests:
- Ultrasound
- CT scan
- MRI
- Genetic Testing: Identifies gene mutations





# Treatment and Management

No cure, but symptoms can be managed

- Medications:
  - Blood pressure control (ACE inhibitors, ARBs)
  - Pain management
  - Tolvaptan (for slowing cyst growth)
- Dialysis or Kidney Transplant: For advanced cases





## Lifestyle and Preventative Measures

- Low-sodium, kidney-friendly diet
- Hydration (but avoiding excessive fluid intake)
- Regular exercise
- Avoiding smoking and alcoho
- Managing blood pressure







### Conclusion

- PKD is a serious genetic condition affecting kidney function
- Early diagnosis and management can slow progression
- · Ongoing research aims to find better treatments





#### References and Resources

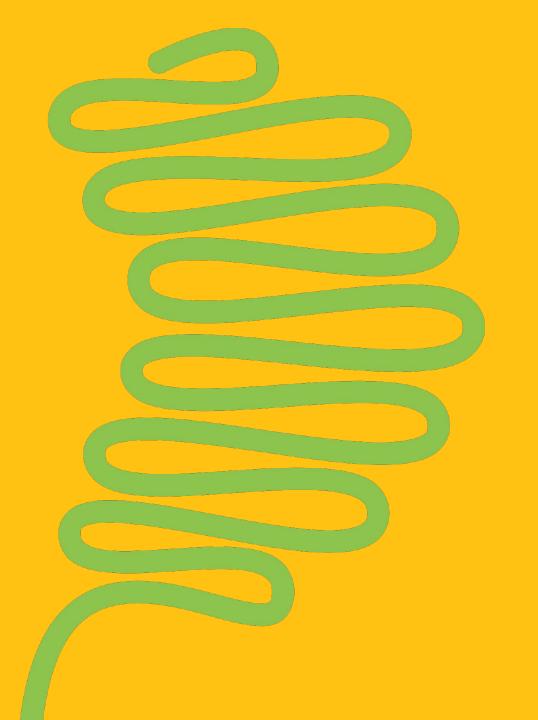


- https://www.kidney.org/kidney-topics/polycystic-kidney-disease
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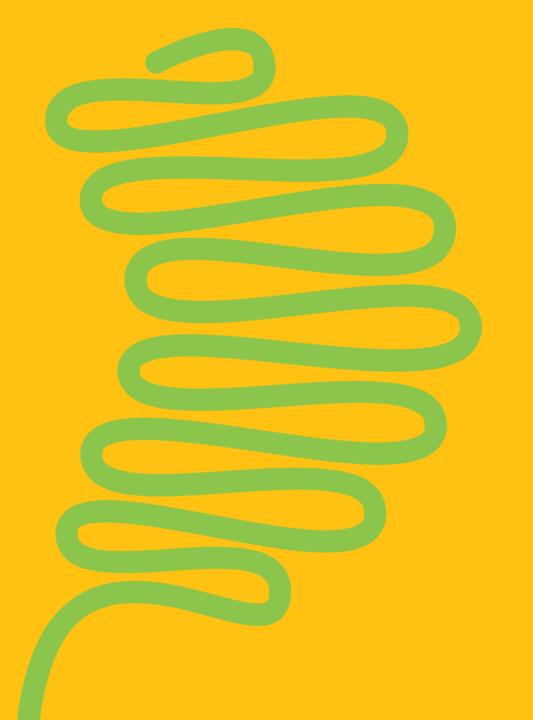






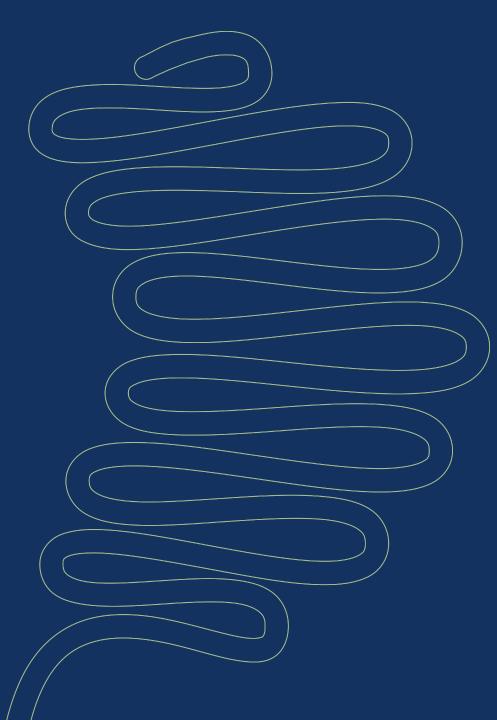












# **QUESTIONS?**Contact Us

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