Learn about APOL1 kidney disease



APOL1 kidney disease is a type of **chronic kidney disease** (CKD) linked to a certain gene called the **APOL1 gene**.

People with Western and Central African ancestry have a higher chance of APOL1 kidney disease. Most people don't know they have kidney disease until it is more serious.

What is chronic kidney disease (CKD)?

Chronic kidney disease is when your kidneys get damaged and slowly stop working. This can lead to kidney failure and other health problems such as heart disease.

Signs and symptoms that your kidneys may not be working well



Foamy pee
Peeing more
often than usual



Feeling tired Feeling nauseous



appetite
Weight loss
without trying

Loss of



High blood pressure Swelling in your legs

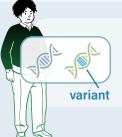
What causes APOL1 kidney disease?

APOL1 kidney disease is CKD that is linked to an abnormal version of the APOL1 gene (piece of DNA) that you may have been born with.

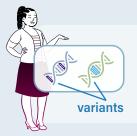


Everyone has 2 copies of the **APOL1 gene**.

Some people have an abnormal version (a variant) of the APOL1 gene that can raise their chance of developing CKD.



If someone has 1 copy of this variant, they have a **slightly higher chance of getting CKD** than people with no variant copies.



If someone has 2 copies of this variant, they have a **much higher chance of getting CKD** than people with no variant copies.



However - only **1 in 5 people** with **2 variant copies** get CKD. People with 2 variant copies may only get CKD after a **second** condition triggers it, such as high blood pressure.

Who gets APOL1 kidney disease?

APOL1 is often found in people with Western and Central African ancestry, such as people who identify as:

Black

- Afro-Caribbean
- African American
- Latina/Latino



APOL1 by the numbers

For people with Western and Central African ancestry in the U.S.:

About 4 - 5 people in 10 have at least 1 variant copy of the gene.

A little over 1 in 10 people have 2 variant copies of the gene.

How do I know if I might have APOL1 kidney disease?

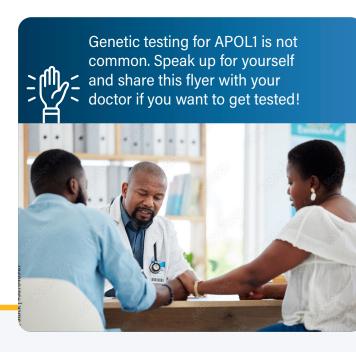


If you have Western and Central African ancestry, ask your doctor about genetic testing for the APOL1 variant.

Genetic testing is important because most people with APOL1 kidney disease:

- Get CKD before age 50
- Don't find out they have CKD until they have symptoms and their kidneys have already been damaged

There are currently no treatments for APOL1 kidney disease – so it is best to find out if you have APOL1 kidney disease through genetic testing before it damages your kidneys.



Where can I learn more?

Learn how you can keep your kidneys healthy:

NIH - Preventing Chronic Kidney Disease



National Kidney
Foundation - 6-Step
Guide to Protecting

Kidney Health



Kidney Health Initiative -APOL1 PDF





CDC - Chronic Kidney Disease Basics

Learn more about APOL1 KD:

National Kidney
Foundation's
APOL1 KD page





American Kidney Fund's APOL1 KD page