

# DIAGNOSIS

REQUIRES 2 OF 3 FEATURES

- ✓ Elevated Alkaline Phosphatase (ALP)
- ✓ Anti-mitochondrial Antibody (AMA) ⊕
- ✓ Compatible Liver Biopsy

Primary Biliary Cholangitis (PBC) is a global autoimmune liver disease predominantly afflicting women of all races and ethnicities. **Early diagnosis is key to success with current therapies.**

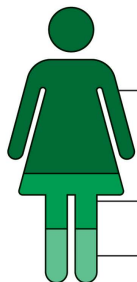


PBC afflicts ~10 females for every 1 male.  
PBC does not normally occur in childhood.

# TREATMENT

**UDCA** Ursodeoxycholic Acid

**OCA** Obeticholic Acid



~60% responders to optimal doses of UDCA

~20% responders to combination of OCA + UDCA <sup>1</sup>

~20% sub-optimal responders <sup>2</sup>

<sup>1</sup> OCA monotherapy is an option for patients unable to tolerate UDCA. <sup>2</sup> New therapies are in clinical trials.

# ABOUT PBCers

We are an organization that supports Primary Biliary Cholangitis patients throughout their journey. One of the main reasons we emphasize education and support is to bolster doctor-patient relationships because a knowledgeable and supported patient is a compliant and empowered patient.

We offer education and support to Primary Biliary Cholangitis patients, family members and friends, and we raise funds to help research the cause(s) and discover a cure for PBC.

# CONTACT US

☎ (346) 302-1620 ✉ [pbcers@pbcers.org](mailto:pbcers@pbcers.org)

# JOIN TODAY

PBCers EXISTS SO THAT NO ONE FEELS ALONE WITH PBC

- ✓ Join our PBC email daily digest
- ✓ Share your story with fellow PBCers
- ✓ Attend a PBC conference or local meeting
- ✓ Follow us on Facebook and Twitter



<http://pbcers.org>

 [facebook.com/PBCersOrg](https://www.facebook.com/PBCersOrg)

 @pbcers

THE PBCers GUIDE TO

# PRIMARY BILIARY CHOLANGITIS (PBC)



