

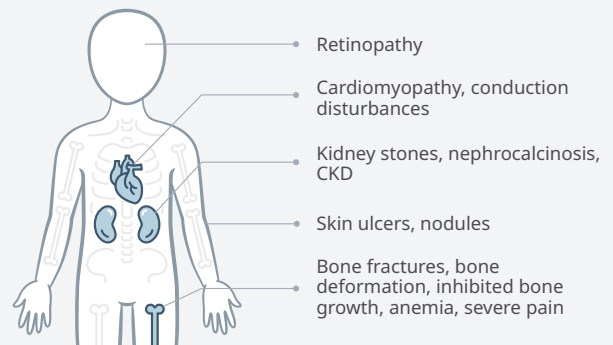
Primary hyperoxaluria (PH), a kidney stone disease that starts in the liver and can lead to systemic damage^{1,2}

PH is a family of ultra-rare genetic disorders in which excess levels of **oxalate** produced by the liver lead to^{1,2}:

- **Recurrent kidney stones (CaOx crystals)**
- **Nephrocalcinosis**
- **Chronic kidney disease**

Progressive kidney damage can result in **end-stage kidney disease** and **systemic oxalate deposition**.¹

Systemic Oxalate Deposition in PH Causes^{1,3,4}



Warning Signs of PH⁵⁻¹¹ (one or a combination of)

The first warning sign may be a single kidney stone in children or recurrent stones in adults.



Family history
of kidney or bladder stones



CKD with no known etiology



Systemic oxalosis



Recurrent stones in adults



Nephrocalcinosis



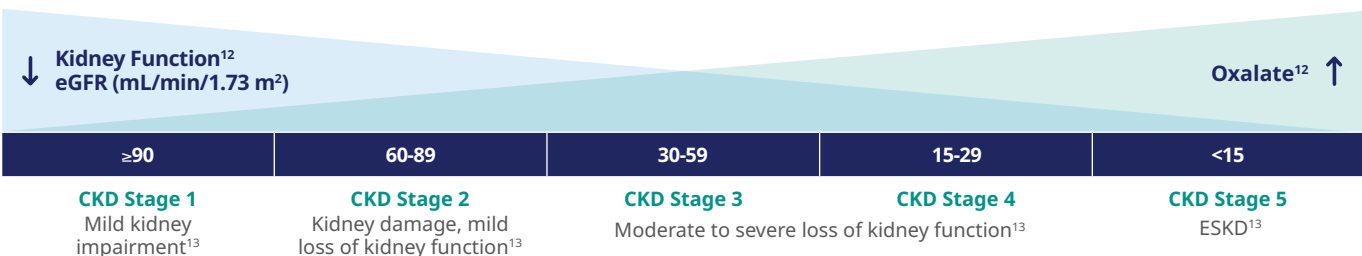
Severe infantile form:
Failure to thrive, ESKD, severe retinal abnormalities



Single kidney stone in a child



ESKD



Oxalate accumulation can occur even in the absence of current symptoms^{12,14}

For more information, visit UNCOVERINGPH.com

Abbreviations: BSA=body surface area; CaOx=calcium oxalate; CKD=chronic kidney disease; COD=calcium oxalate dihydrate; COM=calcium oxalate monohydrate; eGFR=estimated glomerular filtration rate; ESKD=end-stage kidney disease; ESWL=extracorporeal shock wave lithotripsy; HOG=4-hydroxy-2-oxoglutarate; PCNL=percutaneous nephrolithotomy; Pox=plasma oxalate; RNAi=ribonucleic acid interference; siRNA=small interfering ribonucleic acid; Uox=urinary oxalate; URS=ureteroscopy.

For full list of references, please scan the QR code



Please note that this content is not meant to provide diagnosis or treatment recommendations.
The best diagnosis strategy and management options are to be determined by the patient's physician.

PH Diagnosis

Diagnostic Workup

Or

Refer to a Nephrologist for Diagnosis
Visit [OHF.org](https://www.ohf.org) for Resources and for Guidance

Urine Collection (eGFR >30 mL/min/1.73 m²)¹⁵:

24-hour collection (preferred)^{15,16}

Elevated Uox on at least 2 assessments:

- >0.83 mmol/24 h/1.73 m² or >75 mg/24 h

Spot urine¹⁵

Oxalate: creatinine ratio > normal for age

Plasma Collection (eGFR <30 mL/min/1.73 m²)^{1,15}:

Pox >50 μmol/L

Stone Analysis¹⁷⁻¹⁹:

- **PH1:** >95% COM (whewellite)
- **PH2:** Typically >80% COM
- **PH3:** Composed of COM and COD, with 24% to 100% COM

Urinary Metabolites (Nonconfirmatory)^{1,4,15}:

- Elevated glycolate → **PH1**
- Elevated L-glycerate → **PH2**
- Elevated HOG → **PH3**

Rule Out Secondary Causes^{5,15}:

- Enteric causes (eg, chronic pancreatitis, cystic fibrosis, inflammatory bowel syndrome, bariatric surgery)
- Very high-oxalate, low-calcium diet
- Premature infant

Genetic Testing^{5,15}:

- AGXT mutation → **PH1**
- GRHPR mutation → **PH2**
- HOGA1 mutation → **PH3**

Urine and genetic testing can be done simultaneously or sequentially.

Visit [NovoDETECT.com](https://www.novodetect.com) for More Information on Diagnosis

PH Management

Current Available Management Options

Or

Refer to Nephrology for Management
Visit [OHF.org](https://www.ohf.org) for Resources and for Guidance

CaOx Crystal Inhibition¹⁵:

Hyperhydration

- Adults: 3.5 L to 4 L daily
- Children: 2 L to 3 L/m² BSA daily
- Infants: gastrostomy tube

CaOx crystallization inhibitors

- Potassium citrate (preserved renal function patients)

Kidney Stone Management¹⁵:

PCNL or URS instead of ESWL

Oxalate Reduction:

Pyridoxine (vitamin B6) in responsive patients with PH1¹⁵

- Test for responsiveness¹⁵
- B6-responsive AGXT mutations: Gly170Arg, Phe152Ile, Ile244Thr^{11,20,21}

RNAi/siRNA therapy for patients with PH1¹⁵

Advanced CKD (eGFR <30 mL/min/1.73 m²)¹⁵:

Dialysis

For CKD stage 4 to 5, high Pox, systemic oxalosis

Organ transplant

- **PH1**, eGFR <30 mL/min/1.73 m², B6-unresponsive, no access to RNAi → **liver-kidney transplant**
- **PH1**, CKD stage 5, B6-responsive → **kidney transplant**
- **PH2**, eGFR <30 mL/min/1.73 m² → **liver transplant**