

THIS IS NOT A TEST REQUEST FORM.
Please fill out this form and submit it with the test request form or electronic packing list.

PATIENT HISTORY FOR AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE (ADPKD)

Patient Name _____ Date of Birth _____ Sex F M
 Physician _____ Physician Phone _____
 Practice Specialty _____ Physician Fax _____
 Genetic Counselor _____ Counselor Phone _____

Patient's Ethnicity (check all that apply)
 African-American Asian Hispanic Native American
 Ashkenazi Jewish Caucasian Middle Eastern Other: _____

Does the patient have a clinical diagnosis of ADPKD? No Confirmed Suspected Unknown

Is this patient being evaluated as a potential living related kidney donor? No Yes Unknown

Does the patient have symptoms? No Yes (check all that apply and describe)
 Abdominal wall hernia Enlarged kidneys Renal cysts (total number: _____)
 Aortic dissection/dilatation Hypertension bilateral unilateral
 End stage renal disease (ESRD) Intracranial aneurysms Renal insufficiency
 (age of onset: _____) Mitral valve prolapse
 Other symptom(s): _____

Has the patient had renal imaging studies? No Yes Unknown
 If yes, describe imaging results: Normal Abnormal: _____ Uncertain: _____

Has the patient undergone previous DNA testing? No Yes Unknown
 If yes, describe the genes, disorder, methodology, and results: _____

Is there any relevant family history? No Yes Unknown
 If yes, attach a pedigree or specify the relative's relationship to the patient. List their symptoms and age of onset:

Has DNA testing been performed for the family member(s)? No Yes Unknown
 If yes, attach a copy of the relative's DNA laboratory result (REQUIRED for familial mutation testing).

Check the test you intend to order.

Recommended first tier testing for ADPKD:

- 2012250 Polycystic Kidney Disease, Autosomal Dominant (*PKD1* and *PKD2*) Sequencing and Deletion/Duplication: Clinical sensitivity 90% for ADPKD.
- 2012255 Polycystic Kidney Disease, Autosomal Dominant (*PKD1* and *PKD2*) Sequencing: Clinical sensitivity 87% for ADPKD.

Targeted testing for known mutation (a copy of the laboratory report from a family member is REQUIRED):

- 2012246 Polycystic Kidney Disease, Autosomal Dominant (*PKD1* and *PKD2*) Deletion/Duplication: Tests for a large deletion/duplication previously identified in a family member.
- 2001961 Familial Mutation, Targeted Sequencing: Tests for a mutation previously identified in a family member; copy of relative's lab result is REQUIRED.

Master Label

For questions, contact an ARUP genetic counselor at (800) 242-2787, ext. 2141