

THIS IS NOT A TEST REQUEST FORM.
The information below is required to perform molecular genetic testing.
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) TESTING

Patient Name _____ Date of Birth ____/____/____ Gender F M

Physician _____ Physician Phone (____) _____ Practice Specialty _____

Genetic Counselor _____ Counselor Phone (____) _____

Patient's Ethnicity (check all that apply)

- African American Ashkenazi Jewish Asian Caucasian
 Hispanic Middle Eastern Native American 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 of ADPKD? No Yes; please check all that apply

- | | | |
|---|--|--|
| <input type="checkbox"/> Renal cysts:
<input type="checkbox"/> bilateral <input type="checkbox"/> unilateral;
total # _____ | <input type="checkbox"/> Renal insufficiency
<input type="checkbox"/> End stage renal disease (ESRD)
Age of onset: _____ | <input type="checkbox"/> Aortic dissection/dilatation
<input type="checkbox"/> Abdominal wall hernia
<input type="checkbox"/> Mitral valve prolapse
<input type="checkbox"/> Intracranial aneurysms
<input type="checkbox"/> Other _____ |
| <input type="checkbox"/> Enlarged kidneys | <input type="checkbox"/> Hypertension | |

Has the patient had renal imaging studies? No Yes Unknown

Imaging Results: Normal Abnormal _____ Uncertain _____

Does the patient have a FAMILY HISTORY of ADPKD? No Yes Unknown

If yes, attach a PEDIGREE or specify the RELATIONSHIP of the family members to the patient and detail the symptoms/age of onset in each symptomatic relative:

Has DNA testing been performed for these family member(s)? No Yes Unknown

If yes, please attach a copy of the relative's DNA laboratory result (REQUIRED for familial mutation testing)

Has the patient undergone previous DNA testing for PKD? No Yes Unknown

If yes, please describe test(s) and results: _____

Circle the test you intend to order

Recommended first tier testing for Autosomal Dominant Polycystic Kidney Disease:	
2012250	Polycystic Kidney Disease, Autosomal Dominant (<i>PKD1</i> and <i>PKD2</i>) Sequencing and Deletion/Duplication: Clinical sensitivity 90% for ADPKD.
2012255	Polycystic Kidney Disease, Autosomal Dominant (<i>PKD1</i> and <i>PKD2</i>) Sequencing: Clinical sensitivity 87% for ADPKD.
2012246	Polycystic Kidney Disease, Autosomal Dominant (<i>PKD1</i> and <i>PKD2</i>) Deletion/Duplication: Clinical sensitivity 3% for ADPKD. Tests for large deletions/duplications when <i>PKD1</i> and <i>PKD2</i> sequencing was negative.
Targeted testing for known mutation (laboratory report from family member REQUIRED)	
2012246	Polycystic Kidney Disease, Autosomal Dominant (<i>PKD1</i> and <i>PKD2</i>) Deletion/Duplication: Tests for a large deletion/duplication previously identified in a family member.
2001961	Familial Mutation, Targeted Sequencing: Targeted testing for a known familial sequence mutation.

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

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