

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