

THIS IS NOT A TEST REQUEST FORM.
The information below is required to perform Alport Syndrome testing.
Please fill out this form and submit it with the test request form or electronic packing list.

PATIENT HISTORY FOR ALPORT SYNDROME 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)

- | | | | |
|---|---|--|--------------------------------------|
| <input type="checkbox"/> African American | <input type="checkbox"/> Ashkenazi Jewish | <input type="checkbox"/> Asian | <input type="checkbox"/> Caucasian |
| <input type="checkbox"/> Hispanic | <input type="checkbox"/> Middle Eastern | <input type="checkbox"/> Native American | <input type="checkbox"/> Other _____ |

Does the patient have SYMPTOMS of Alport Syndrome? No Yes (check all that apply)

- | | | |
|--|--|---|
| <input type="checkbox"/> Hematuria | <input type="checkbox"/> Corneal erosion | <input type="checkbox"/> Proteinuria |
| <input type="checkbox"/> Hypertension | <input type="checkbox"/> Renal insufficiency | <input type="checkbox"/> End stage renal disease (Age _____) |
| <input type="checkbox"/> Anterior lenticonus | <input type="checkbox"/> Maculopathy | <input type="checkbox"/> Sensorineural hearing loss (Age _____) |
| <input type="checkbox"/> Other _____ | | |

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

If yes, **please attach PEDIGREE** or specify the RELATIONSHIP of the family member(s) to the patient and detail the symptoms/age of onset in each symptomatic relative.

Please attach a copy of the relative's DNA laboratory result (REQUIRED for familial mutation testing).

Has the patient undergone previous testing for Alport Syndrome? No Yes

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

Circle the test you intend to order.

2002398 Alport Syndrome, X-Linked (COL4A5) Sequencing and Deletion/Duplication Detects >80% of mutations in males and females with X-linked Alport syndrome.

0051786 Alport Syndrome, X-linked (COL4A5) Sequencing Offer to patients who have a family history of X-linked renal failure or to individuals with unexplained hematuria or chronic kidney disease. Clinical sensitivity > 80% in males and >70% in females with X-linked Alport syndrome.

2002394 Alport Syndrome, X-Linked (COL4A5) Deletion/Duplication For patients with negative COL4A5 sequencing result. Clinical sensitivity 10%. **Also order for familial COL4A5 large deletion or duplication.**

2001961 Familial Mutation, Targeted Sequencing Targeted sequencing for a COL4A5 mutation previously identified in a family member; copy of relative's lab result is REQUIRED.

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

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