

**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 AUTISM AND INTELLECTUAL DISABILITY**

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: \_\_\_\_\_

**Clinical Findings:** (check all that apply)

<u>Growth</u>	<u>Genitourinary</u>	<u>Neurologic</u>	<u>Musculoskeletal</u>
<input type="checkbox"/> Failure to thrive	<input type="checkbox"/> Ambiguous genitalia	<input type="checkbox"/> Ataxia/dystonia/chorea	<input type="checkbox"/> Scoliosis
<input type="checkbox"/> Overgrowth	<input type="checkbox"/> Hydronephrosis	<input type="checkbox"/> Hypotonia	<input type="checkbox"/> Vertebral anomaly
<input type="checkbox"/> Short stature	<input type="checkbox"/> Undescended testes	<input type="checkbox"/> Seizures	<input type="checkbox"/> Limb anomaly
<u>Development</u>	<u>Cognition/behavior</u>	<input type="checkbox"/> Spasticity	<input type="checkbox"/> Organomegaly
<input type="checkbox"/> Fine motor delay	<input type="checkbox"/> Autism spectrum disorder	<input type="checkbox"/> Structural brain abnormality	<u>Craniofacial</u>
<input type="checkbox"/> Gross motor delay	<input type="checkbox"/> Learning disability	<u>Cardiac</u>	<input type="checkbox"/> Dysmorphic facial features
<input type="checkbox"/> Speech delay	<input type="checkbox"/> Intellectual disability; (IQ/DQ: _____)	<input type="checkbox"/> Cardiomyopathy	<input type="checkbox"/> Cleft lip +/- cleft palate
<u>Biochemical</u>	<input type="checkbox"/> Oppositional/defiant disorder	<input type="checkbox"/> Structural heart defect (describe: _____)	<input type="checkbox"/> Craniosynostosis
<input type="checkbox"/> Acidosis	<input type="checkbox"/> Obsessive compulsive disorder		<input type="checkbox"/> Macrocephaly
<input type="checkbox"/> Hypoglycemia	<input type="checkbox"/> Pervasive developmental delay		<input type="checkbox"/> Microcephaly
<input type="checkbox"/> Hyperammonemia			<input type="checkbox"/> Corneal clouding

List patient's medications and, if applicable, describe modified diet. Include formulas, vitamins, supplements, antibiotics, anticonvulsants, or enzyme replacement therapy: \_\_\_\_\_

Has the patient undergone previous DNA testing?  No  Yes  Unknown  
 If yes, describe the test(s) 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 diagnoses and age of onset: \_\_\_\_\_

Has DNA testing been performed for the family member(s)?  No  Yes  Unknown  
 If yes, describe the test(s) and results: \_\_\_\_\_

Are the patient's parents related to one another?  No  Yes (describe: \_\_\_\_\_)  Unknown

In cooperation with the National Institutes of Health's effort to improve understanding of specific genetic variants, ARUP submits HIPAA-compliant, de-identified (cannot be traced back to the patient) genetic test results and health information to public databases. The confidentiality of each sample is maintained. If you prefer that your test result not be shared, call ARUP at (800) 242-2787, ext. 3301. Your de-identified information will not be disclosed to public databases after your request is received, but a separate request is required for each genetic test. Additionally, patients have the opportunity to participate in patient registries and research. To learn more, visit [www.aruplab.com/genetics/resources](http://www.aruplab.com/genetics/resources).

**Check the test you intend to order.**

Primary panel for Autism/Intellectual Disability:

- 2014314 Autism and Intellectual Disability Comprehensive Panel  
 Individual components are available separately:
  - 2003414 Cytogenomic SNP Microarray
  - 2009033 Fragile X (FMR1) with Reflex to Methylation Analysis
  - 2014312 Autism and Intellectual Disability Metabolic Panel



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