

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

AUTISM AND INTELLECTUAL DISABILITY PATIENT HISTORY FORM

Patient Name: _____ Date of Birth: _____
 Sex Assigned at Birth: Female Male Intersex Gender Identity (optional): Female Male _____
 Ordering Provider: _____ Provider's Phone: _____
 Practice Specialty: _____ Provider's Fax: _____
 Genetic Counselor: _____ Counselor's Phone: _____

Patient's Ethnicity/Ancestry (check all that apply)
 African American/Black Asian Hispanic White Other: _____
 List country of origin (if known): _____

Clinical Findings: (check all that apply)

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

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, deidentified (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 deidentified 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 aruplab.com/genetics

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

