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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 GENOMIC MICROARRAY TESTING

Patient Name		Date of Birth	Sex □ F □ M
Physician		Physician Phone	
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Constin Councilor		Councelor Phone	
Clinical Information (check all that a			
Perinatal History	Behavioral	Craniofacial	Gastrointestinal
☐ Not evaluated	☐ Not evaluated	☐ Not evaluated	☐ Not evaluated
☐ Within normal limits	☐ Within normal limits	☐ Within normal limits	☐ Within normal limits
☐ Prematurity	☐ Asperger syndrome features	☐ Cleft lip +/- cleft palate	☐ Gastroschisis
, □ IUGR	☐ Autism	☐ Cleft palate alone	☐ Hirschsprung disease
☐ Oligohydramnios	☐ Oppositional defiant disorder	Coloboma	☐ Omphalocele
☐ Polyhydramnios	☐ Obsessive compulsive disorder	☐ Craniosynostosis	☐ Pyloric stenosis
☐ Other:	☐ Pervasive developmental delay	☐ Dysmorphic facial features	
	☐ Other:	•	Other:
Growth		☐ Macrocephaly	
☐ Not evaluated	Neurological	☐ Microcephaly	Genitourinary
☐ Within normal limits	☐ Not evaluated	☐ List HC, if known:	☐ Not evaluated
☐ Failure to thrive	☐ Within normal limits	☐ Other:	☐ Within normal limits
☐ Overgrowth	☐ Ataxia/dystonia/chorea		☐ Ambiguous genitalia
☐ Short stature	☐ Hypotonia	Cutaneous	☐ Hydronephrosis
Other:	☐ Neural tube defect	☐ Not evaluated	☐ Hypospadias
Development	☐ Seizures	☐ Within normal limits	☐ Kidney malformation
☐ Not evaluated	☐ Spasticity	☐ Hyperpigmentation	☐ Undescended testis
☐ Within normal limits	☐ Structural brain anomaly	☐ Hypopigmentation	☐ Urethra/ureter obstruction
☐ Fine motor delay	Other:	Musculoskeletal	Other:
☐ Gross motor delay	Cardiac	☐ Not evaluated	Family History
☐ Speech delay	☐ Not evaluated	☐ Within normal limits	☐ Not evaluated
☐ Other:	☐ Within normal limits	☐ Contractures	☐ No relevant family history
Cognitive	☐ ASD	☐ Club foot	☐ Parents with
☐ Not evaluated	☐ AV canal defect	☐ Diaphragmatic hernia	≥ two miscarriages
☐ Within normal limits	☐ Coarctation of aorta	\square Limb anomaly	☐ Other relatives with
☐ Learning disability	☐ Hypoplastic left heart	☐ Polydactyly	similar clinical history
☐ Intellectual disability	☐ Tetralogy of fallot	☐ Scoliosis	(explain below)
☐ List IQ/DQ, if known:	□ VSD	☐ Syndactyly	
☐ Other:	Other:	☐ Vertebral anomaly	
- other.		☐ Other:	
Clinical Descriptions—Include any a	additional relevant clinical informatic	on not provided above.	
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			Master Label
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.			
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For questions, contact an A			