

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

**PATIENT HISTORY FORM FOR PRENATAL CYTOGENETICS**

Patient Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_ Sex:  Female  Male  
 Ordering Provider: \_\_\_\_\_ Provider's Phone: \_\_\_\_\_  
 Practice Specialty: \_\_\_\_\_ Provider's Fax: \_\_\_\_\_  
 Genetic Counselor: \_\_\_\_\_ Counselor's Phone: \_\_\_\_\_  
 Date of Draw: \_\_\_\_\_ Gestational Age at Draw: \_\_\_\_\_ weeks \_\_\_\_\_ days

|  |   |   |
|--|---|---|
| <p><b>Chorionic Villus (CVS)</b></p> <p><input type="checkbox"/> 0040203 CVS, FISH</p> <p><input type="checkbox"/> 2002291 Chromosome Analysis, CVS</p> <p><input type="checkbox"/> 2002366 Cytogenomic SNP Microarray—Fetal</p> <p><input type="checkbox"/> 2011131 Chromosome FISH, CVS with Reflex to Chromosome Analysis or Genomic Microarray</p> | <p><b>Amniotic Fluid (AF)</b></p> <p><input type="checkbox"/> 2002297 Chromosome FISH, Prenatal</p> <p><input type="checkbox"/> 2002293 Chromosome Analysis, AF</p> <p><input type="checkbox"/> 2002366 Cytogenomic SNP Microarray—Fetal</p> <p><input type="checkbox"/> 2008367 Chromosome Analysis, Amniotic fluid, with Reflex to Genomic Microarray</p> <p><input type="checkbox"/> 2011130 Chromosome FISH, AF with Reflex to Chromosome Analysis or Genomic Microarray</p> <p><input type="checkbox"/> 3000142 Alpha Fetoprotein (AFP) with Reflex Acetylcholinesterase and Fetal Hemoglobin</p> <p><input type="checkbox"/> 2008367 Chromosome Analysis, AF, with Reflex to Genomic Microarray</p> | <p><b>Products of Conception; Fresh/FFPE (POC)</b></p> <p><input type="checkbox"/> 2002288 Chromosome Analysis, POC</p> <p><input type="checkbox"/> 2005633 Genomic SNP Microarray, POC</p> <p><input type="checkbox"/> 2005762 Chromosome Analysis, POC, with Reflex to Genomic Microarray</p> <p><input type="checkbox"/> 2010795 Cytogenomic Molecular Inversion Probe Array, FFPE Tissue—POC</p> <p><b>Maternal Blood</b></p> <p><input type="checkbox"/> 0050608 Maternal Cell Contamination, Maternal Specimen</p> <p><input type="checkbox"/> 2002369 Microarray Genomic, Maternal Confirm</p> |
|--|---|---|

Fetal sex by ultrasound:  Male  Female  Ambiguous  Unknown

For microarray and MCC studies only: Is the patient the biological mother of the fetus? .....  No  Yes  
 Is there consanguinity? .....  No  Yes

**Indication for Testing (check all that apply)**

|   |   |
|---|---|
| <input type="checkbox"/> Advanced maternal age  | <input type="checkbox"/> Familial chromosome abnormality (provide relationship to fetus, specific abnormality, and a copy of the family member's result): _____ |
| <input type="checkbox"/> Abnormal maternal serum screen: <input type="checkbox"/> T21 <input type="checkbox"/> T18 <input type="checkbox"/> High AFP<br><input type="checkbox"/> Other: _____   | <input type="checkbox"/> Fetus with KNOWN chromosome abnormality (describe and provide a copy of the chromosome report): _____                                  |
| <input type="checkbox"/> Abnormal cfDNA (NIPT): <input type="checkbox"/> T21 <input type="checkbox"/> T18 <input type="checkbox"/> T13 <input type="checkbox"/> TS <input type="checkbox"/> SCA<br><input type="checkbox"/> Atypical: CHR <input type="checkbox"/> Other: _____ |   |

**Ultrasound Abnormality (check finding(s) or list under "other")**

|   |   |   |  |   |
|---|---|---|--|---|
| <p><b>Abdominal/Chest</b></p> <p><input type="checkbox"/> Diaphragmatic hernia</p> <p><input type="checkbox"/> Duodenal atresia</p> <p><b>Amniotic Fluid</b></p> <p><input type="checkbox"/> Oligohydramnios</p> <p><input type="checkbox"/> Polyhydramnios</p> <p><b>Cardiac</b></p> <p><input type="checkbox"/> ASD <input type="checkbox"/> VSD</p> <p><input type="checkbox"/> HLH <input type="checkbox"/> TOF</p> <p><input type="checkbox"/> Aortic Stenosis</p> <p><input type="checkbox"/> Other: _____</p> <p><b>Neural Tube</b></p> <p><input type="checkbox"/> Anencephaly</p> <p><input type="checkbox"/> Encephalocele</p> <p><input type="checkbox"/> Spina bifida</p> | <p><b>Cranial Facial</b></p> <p><input type="checkbox"/> Agenesis of the corpus callosum</p> <p><input type="checkbox"/> Absent CSP</p> <p><input type="checkbox"/> Cleft lip</p> <p><input type="checkbox"/> Dandy-Walker</p> <p><input type="checkbox"/> Holopresencephaly</p> <p><input type="checkbox"/> Hydrocephaly</p> <p><input type="checkbox"/> Microcephaly</p> <p><input type="checkbox"/> Micrognathia</p> <p><input type="checkbox"/> Ventriculomegaly</p> <p><b>Fetal Well-Being</b></p> <p><input type="checkbox"/> Fetal demise</p> <p><input type="checkbox"/> IUGR</p> <p><input type="checkbox"/> SGA/size &lt; dates</p> | <p><b>Fluid Collection</b></p> <p><input type="checkbox"/> Ascites</p> <p><input type="checkbox"/> Cystic hygroma</p> <p><input type="checkbox"/> Hydrops</p> <p><input type="checkbox"/> Increased NT</p> <p><input type="checkbox"/> Skin edema</p> <p><input type="checkbox"/> Pericardial effusion</p> <p><input type="checkbox"/> Pleural effusion</p> <p><b>Limb/Joint</b></p> <p><input type="checkbox"/> Arthrogryposis</p> <p><input type="checkbox"/> Clenched hands</p> <p><input type="checkbox"/> Clubfoot</p> <p><input type="checkbox"/> Polydactyly</p> <p><input type="checkbox"/> Rocker bottom foot</p> <p><input type="checkbox"/> Syndactyly</p> | <p><b>Markers/Soft Signs</b></p> <p><input type="checkbox"/> Absent nasal bone</p> <p><input type="checkbox"/> Pyelectasis</p> <p><input type="checkbox"/> Choroid plexus cyst</p> <p><input type="checkbox"/> SUA</p> <p><input type="checkbox"/> Echogenic bowel</p> <p><input type="checkbox"/> Echogenic Cardiac focus</p> <p><input type="checkbox"/> Thickened nuchal fold</p> <p><b>Skeletal</b></p> <p><input type="checkbox"/> "Bent" Bones</p> <p><input type="checkbox"/> Radial ray defect</p> <p><input type="checkbox"/> Short long bones</p> <p><input type="checkbox"/> Short ribs</p> <p><input type="checkbox"/> Vertebral anomalies</p> | <p><b>Urinary Tract</b></p> <p><input type="checkbox"/> Bladder exstrophy</p> <p><input type="checkbox"/> Bladder outlet obstruction</p> <p><input type="checkbox"/> Hydronephrosis</p> <p><input type="checkbox"/> Multicystic kidney</p> <p><input type="checkbox"/> Posterior Urethral Valves</p> <p><input type="checkbox"/> Renal agenesis</p> <p><b>Ventral Wall Defect</b></p> <p><input type="checkbox"/> Gastroschisis</p> <p><input type="checkbox"/> Limb-body wall defect</p> <p><input type="checkbox"/> Omphalocele</p> |
|---|---|---|--|---|

Other: \_\_\_\_\_

**Additional Testing on Sample**

Culture cells for additional testing. Test desired: \_\_\_\_\_

Culture/hold cells for possible additional testing (samples retained for 3 weeks)

Store long-term back-up cultures (two T-25 flasks frozen and retained for 6 months)



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