

Chromosomal Microarray Prenatal Patient Information

Instructions: The accurate interpretation and reporting of genetic results is contingent upon the reason for referral, clinical information provided, and family history. Supply the information requested below and send paperwork with the specimen or return by fax to the Cytogenetics Laboratory, Attn: Genetic Counselors. Fax: 507-284-0043 Phone: 507-538-2952 / International clients: +1 507 266 5700 or email mmlglobal@mayo.edu

Patient Name <i>(Last, First, Middle Initial)</i>	Birth Date <i>(Month DD, YYYY)</i>	Sex <input type="checkbox"/> Male <input type="checkbox"/> Female
MML Account Number (if known)	Physician Phone	Fax

Clinical Information – check all that apply

<p>Primary Indication for Testing</p> <p><input type="checkbox"/> Abnormal aneuploidy screening</p> <p><input type="checkbox"/> Advanced maternal age</p> <p><input type="checkbox"/> Fetal abnormality</p> <p><input type="checkbox"/> Other _____</p> <p>Perinatal History</p> <p><input type="checkbox"/> 2 vessel cord</p> <p><input type="checkbox"/> Hydrops</p> <p><input type="checkbox"/> Increased NT (includes cystic hygroma)</p> <p><input type="checkbox"/> IUGR</p> <p><input type="checkbox"/> Oligohydramnios</p> <p><input type="checkbox"/> Polyhydramnios</p> <p><input type="checkbox"/> Other _____</p> <p>Neurological</p> <p><input type="checkbox"/> Abnormal gyri (lissencephaly)</p> <p><input type="checkbox"/> Agenesis of the corpus callosum</p> <p><input type="checkbox"/> Cerebellar hypoplasia</p> <p><input type="checkbox"/> Dandy Walker</p> <p><input type="checkbox"/> Decreased fetal movement</p> <p><input type="checkbox"/> Holoprosencephaly</p> <p><input type="checkbox"/> Neural tube defect</p> <p><input type="checkbox"/> Structural brain anomaly</p> <p><input type="checkbox"/> Ventriculomegaly/hydrocephaly</p> <p><input type="checkbox"/> Other _____</p> <p>Craniofacial</p> <p><input type="checkbox"/> Cleft lip + / - palate</p> <p><input type="checkbox"/> Macrocephaly</p> <p><input type="checkbox"/> Microcephaly</p> <p><input type="checkbox"/> Other _____</p>	<p>Cardiac</p> <p><input type="checkbox"/> Aortic atresia</p> <p><input type="checkbox"/> Atrial septal defect</p> <p><input type="checkbox"/> AV canal defect</p> <p><input type="checkbox"/> Coarctation of the aorta</p> <p><input type="checkbox"/> Dextrocardia/situs inversus</p> <p><input type="checkbox"/> Double outlet right ventricle</p> <p><input type="checkbox"/> Ebstein anomaly</p> <p><input type="checkbox"/> Echogenic intracardiac focus</p> <p><input type="checkbox"/> Hypoplastic left heart</p> <p><input type="checkbox"/> Hypoplastic right heart</p> <p><input type="checkbox"/> Tetralogy of Fallot</p> <p><input type="checkbox"/> Transposition of the great vessels</p> <p><input type="checkbox"/> Truncus arteriosus</p> <p><input type="checkbox"/> Ventricular septal defect</p> <p><input type="checkbox"/> Other _____</p> <p>Pulmonary</p> <p><input type="checkbox"/> CCAM/small thoracic cavity</p> <p><input type="checkbox"/> Diaphragmatic hernia</p> <p><input type="checkbox"/> Pleural effusion</p> <p><input type="checkbox"/> Pulmonary sequestration</p> <p><input type="checkbox"/> Other _____</p>	<p>Gastrointestinal</p> <p><input type="checkbox"/> Absent stomach</p> <p><input type="checkbox"/> Echogenic focus</p> <p><input type="checkbox"/> Gastroschisis</p> <p><input type="checkbox"/> Meconium ileus/anal atresia</p> <p><input type="checkbox"/> Omphalocele</p> <p><input type="checkbox"/> Tracheoesophageal fistula</p> <p><input type="checkbox"/> Other _____</p> <p>Musculoskeletal</p> <p><input type="checkbox"/> Acromelia</p> <p><input type="checkbox"/> Clenched hands</p> <p><input type="checkbox"/> Club foot</p> <p><input type="checkbox"/> Contractures (arthrogryposis)</p> <p><input type="checkbox"/> Limb anomaly</p> <p><input type="checkbox"/> Mesomelia/micromelia</p> <p><input type="checkbox"/> Polydactyly</p> <p><input type="checkbox"/> Skeletal dysplasia</p> <p><input type="checkbox"/> Syndactyly</p> <p><input type="checkbox"/> Vertebral anomaly</p> <p><input type="checkbox"/> Other _____</p> <p>Genitourinary</p> <p><input type="checkbox"/> Ambiguous genitalia</p> <p><input type="checkbox"/> Hydronephrosis kidney malformation</p> <p><input type="checkbox"/> Megacystis (including posterior valves)</p> <p><input type="checkbox"/> Polycystic kidneys</p> <p><input type="checkbox"/> Renal agenesis</p> <p><input type="checkbox"/> Urethra/ureter obstruction</p> <p><input type="checkbox"/> Other _____</p> <p>Family History</p> <p><input type="checkbox"/> Parents with greater than or equal to two miscarriages</p> <p><input type="checkbox"/> Other relatives with previous pregnancies with similar clinical history. (explain below)</p> <p><input type="checkbox"/> Other _____</p>
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Clinical Descriptions—include any additional relevant clinical information not provided above (list karyotype if known and provide a report)