



**Instructions:** The accurate interpretation and reporting of genetic test results is contingent upon the reason for referral, clinical information provided, and family history. To help provide the best possible service, please check the applicable clinical information below.

**Patient Identification**

**Patient Name:** \_\_\_\_\_ (Last) \_\_\_\_\_ (First)      **Gender:**  Male  Female  
**Date of Birth:** \_\_\_\_\_ (mm/dd/yyyy)

**Clinical Information – Check all that apply. Use additional space at the bottom of the form if needed.**

<p><b>Perinatal History</b></p> <p><input type="checkbox"/> Prematurity  <input type="checkbox"/> Intrauterine growth restriction  <input type="checkbox"/> Oligohydramnios  <input type="checkbox"/> Polyhydramnios  <input type="checkbox"/> Non-immune hydrops fetalis  <input type="checkbox"/> Other: _____</p> <p><b>Growth</b></p> <p><input type="checkbox"/> Failure to thrive  <input type="checkbox"/> Overgrowth  <input type="checkbox"/> Short stature  <input type="checkbox"/> Other: _____</p> <p><b>Cognitive/Developmental</b></p> <p><input type="checkbox"/> Learning disability  <input type="checkbox"/> Developmental delay  <input type="checkbox"/> Gross motor delay  <input type="checkbox"/> Fine motor delay  <input type="checkbox"/> Speech delay  <input type="checkbox"/> Intellectual disability/MR  <input type="checkbox"/> Other: _____</p> <p><b>Behavioral/Psychiatric</b></p> <p><input type="checkbox"/> Autism  <input type="checkbox"/> Pervasive developmental delay  <input type="checkbox"/> Attention deficit hyperactivity disorder  <input type="checkbox"/> Anxiety  <input type="checkbox"/> Behavioral/psychiatric abnormality  Specify: _____  <input type="checkbox"/> Other: _____</p> <p><b>Cutaneous</b></p> <p><input type="checkbox"/> Hyperpigmentation  <input type="checkbox"/> Hypopigmentation  <input type="checkbox"/> Other: _____</p>	<p><b>Neurological</b></p> <p><input type="checkbox"/> Seizures  <input type="checkbox"/> Hypotonia  <input type="checkbox"/> Hypertonia  <input type="checkbox"/> Cerebral palsy  <input type="checkbox"/> Encephalopathy  <input type="checkbox"/> Structural brain anomaly  Specify: _____  <input type="checkbox"/> Other: _____</p> <p><b>Cardiac</b></p> <p><input type="checkbox"/> Atrial septal defect  <input type="checkbox"/> Ventricular septal defect  <input type="checkbox"/> Coarctation of the aorta  <input type="checkbox"/> Tetralogy of Fallot  <input type="checkbox"/> Other structural heart defect  Specify: _____  <input type="checkbox"/> Other cardiac abnormality  Specify: _____</p> <p><b>Craniofacial</b></p> <p><input type="checkbox"/> Dysmorphic facial features  Specify: _____  <input type="checkbox"/> Ear malformation  Specify: _____  <input type="checkbox"/> Cleft lip  <input type="checkbox"/> Cleft palate  <input type="checkbox"/> Macrocephaly  <input type="checkbox"/> Microcephaly  <input type="checkbox"/> Other: _____</p> <p><b>Hearing/Vision</b></p> <p><input type="checkbox"/> Hearing loss  Specify: _____  <input type="checkbox"/> Abnormality of Vision  Specify: _____  <input type="checkbox"/> Abnormality of Eye Movement  Specify: _____  <input type="checkbox"/> Other: _____</p>	<p><b>Musculoskeletal</b></p> <p><input type="checkbox"/> Contractures  <input type="checkbox"/> Club foot  <input type="checkbox"/> Diaphragmatic hernia  <input type="checkbox"/> Limb anomaly  Specify: _____  <input type="checkbox"/> Polydactyly  Specify: _____  <input type="checkbox"/> Syndactyly  Specify: _____  <input type="checkbox"/> Vertebral anomaly  Specify: _____  <input type="checkbox"/> Other: _____</p> <p><b>Gastrointestinal</b></p> <p><input type="checkbox"/> Gastroschisis  <input type="checkbox"/> Omphalocele  <input type="checkbox"/> Anal atresia  <input type="checkbox"/> Tracheoesophageal fistula  <input type="checkbox"/> Pyloric stenosis  <input type="checkbox"/> Other: _____</p> <p><b>Genitourinary</b></p> <p><input type="checkbox"/> Ambiguous genitalia  <input type="checkbox"/> Hydronephrosis  <input type="checkbox"/> Kidney malformation  Specify: _____  <input type="checkbox"/> Cryptorchidism  <input type="checkbox"/> Hypospadias  <input type="checkbox"/> Other: _____</p> <p><b>Family History</b></p> <p><input type="checkbox"/> Parents with ≥ 2 miscarriages  <input type="checkbox"/> Other relatives with similar clinical history  Explain: _____  _____  _____</p>
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Please include any additional relevant information not provided above.

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As a participant in the Clinical Genome Resource (ClinGen), this clinical laboratory contributes submitted clinical information and test results to a HIPAA compliant, de-identified public database as part of the NIH's effort to improve understanding of the relationship between genetic changes and clinical symptoms. Confidentiality is maintained. Patients may request to opt-out of this scientific effort by: 1) checking the box below, 2) calling the laboratory at **xxx-xxx-xxxx** and asking to speak with a laboratory genetic counselor. Please call with any questions.

**Indicate refusal for inclusion in these efforts by checking this box. If the box is not marked, data will be anonymized and used.**