

NEUROGENETICS LAB REQUISITION FORM

<p>Collection Information</p> <p>Date of Collection: ____ / ____ / ____ MM DD YY</p> <p>Sample Type: <input type="checkbox"/> BLOOD <input type="checkbox"/> SALIVA <input type="checkbox"/> BUCCAL SWAB</p> <hr/> <p>Indication for Study</p> <p>ICD-10 Diagnosis Codes: _____</p>	<p>Patient Information</p> <p style="text-align: center;">(Affix patient label here)</p>
Reporting Information	
<p>Physician: Dr. _____</p> <p>Institution: _____</p> <p>Office: _____</p> <p>Phone: _____ Fax: - _____</p> <p>E-mail: _____@_____</p>	<p>Additional Professional Report Recipients</p> <p>Name: _____</p> <p>Institution: _____</p> <p>Office: _____</p> <p>Phone: - _____ Fax: - _____</p> <p>E-mail: _____@_____</p>
First-line Tests	FISH Tests for Microdeletion Syndromes
<input type="checkbox"/> CBR Chromosome Analysis (Karyotype), Constitutional	<input type="checkbox"/> FDEL01 1p36 (LSIp58)
<input type="checkbox"/> CGH SNP Microarray Analysis, Constitutional	<input type="checkbox"/> FDEL02 Angelman (D15S10), 15q11-q13
<input type="checkbox"/> FXMO Fragile X FMR1 molecular analysis	<input type="checkbox"/> FDEL03 Cri-du-Chat (CDC), 5p15.2
<input type="checkbox"/> DMD DMD Deletion/Duplication ONLY	<input type="checkbox"/> FDEL06 Miller Dieker (LIS1), 17p13.3
Next Generation Sequencing	<input type="checkbox"/> FDEL07 Prader-Willi (SNRPN), 15q11-q13
<input type="checkbox"/> ALS ALS Panel	<input type="checkbox"/> FDEL08 Smith Magenis (SMS), 17p11
<input type="checkbox"/> AIDP Autism/Intellectual Disability Panel	<input type="checkbox"/> FDEL10 Williams (ELN), 7q11.23
<input type="checkbox"/> EPIL Epilepsy Panel	<input type="checkbox"/> FDEL11 Wolf Hirschhorn (WHS), 4p16.1
<input type="checkbox"/> MITOSEQ Mitochondrial DNA Sequencing	
<input type="checkbox"/> NEURO Comprehensive Neurology Panel <i>(includes ALS, Autism/Intellectual Disability, Epilepsy, and Neuromuscular Panels **DOES NOT include Fragile X)</i>	<p>LAB USE ONLY</p> <p>Genetics # _____</p> <p>Lab/Specimen # _____</p> <p>Date/Time received _____</p>
<input type="checkbox"/> IRDP Inherited Recessive Disease Panel	
Disease-Specific Tests	
<input type="checkbox"/> FXMO Fragile X FMR1 molecular analysis	
<input type="checkbox"/> PWS/AS Prader-Willi/Angelman syndrome methylation	
<input type="checkbox"/> SMA SMN1 and SMN2 gene dosage	

DIRECTIONS:

- 1) Complete the above requisition form.
 - 2) Collect one EDTA/lavender top tube 2-3cc blood (NOTE: for FISH tests and Chromosome Karyotype include an extra tube Sodium Heparin 2-3cc blood).
- 2) Attach the following:
- Copy of patient's insurance card
 - Patient's face sheet
 - Patient clinical information (e.g. recent note)