

MUSCULAR DYSTROPHY MOLECULAR GENETICS REQUISITION

FOR UIDL USE ONLY: MRN# _____		PATH# _____																					
FOR CLIENT USE ONLY: Requisition Date _____		Completed by: _____																					
PART A – PATIENT INFORMATION – <i>Required</i>		PART B – PROVIDER INFORMATION – <i>Required</i>																					
Patient Last Name: _____		Referring Institution: _____																					
Patient First Name: _____		Street: _____																					
Street: _____		City: _____ St: _____ Zip: _____																					
City: _____ St: _____ Zip: _____		Phone: _____ Fax: _____																					
Phone: _____ Fax: _____		Referring Physician: _____																					
Date of Birth: _____ Gender: <input type="checkbox"/> M <input type="checkbox"/> F		Referring Physician Phone: _____																					
PART C - SPECIMEN INFORMATION (Complete the appropriate information below or include in an accompanying letter.) MATERIALS SUBMITTED: <input type="checkbox"/> EDTA Whole Blood <input type="checkbox"/> DNA <input type="checkbox"/> Other (Specify): _____ Required ICD-10 codes: 1. _____ 2. _____ 3. _____ 4. _____ 5. _____ TISSUE SOURCE/SITE: _____ DATE OF COLLECTION: _____ PERTINENT CLINICAL HISTORY AND FINDINGS (please attach pedigree if available): _____																							
<p>This request to order molecular diagnostic tests from University of Iowa Diagnostic Laboratories (UIDL) certifies to UIDL that the ordering physician has obtained informed consent from the patient as required by applicable state or federal laws for each test ordered and that the ordering physician has authorization from the patient permitting UIDL to report results for each test ordered to the ordering physician. Genetic Counseling and Information: By requesting testing, the ordering physician assumes responsibility for providing the patient with all associated guidance and counseling regarding the test results. Alternatively, patients can be referred to qualified counseling services by contacting client services at ph. (866)844-2522.</p> <p>SELECT TEST: Refer to the UIDL TEST DIRECTORY for specimen requirements and CPT Codes at https://www.healthcare.uiowa.edu/path_handbook/rindex.html</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td><input type="checkbox"/> Congenital Muscular Dystrophy type 1C, MDC1C (<i>FKRP</i> sequencing)</td> <td><input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2I (<i>FKRP</i> sequencing)</td> </tr> <tr> <td><input type="checkbox"/> Congenital Muscular Dystrophy type 1D, MDC1D (<i>LARGE</i> sequencing)</td> <td><input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2K (<i>POMTI</i> sequencing)</td> </tr> <tr> <td><input type="checkbox"/> Fukuyama Congenital Muscular Dystrophy (<i>FKTN</i> sequencing)</td> <td><input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2M (<i>FKTN</i> sequencing)</td> </tr> <tr> <td><input type="checkbox"/> Fukuyama Congenital Muscular Dystrophy (Japanese Founder Mutation PCR)</td> <td><input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2N (<i>POMT2</i> sequencing)</td> </tr> <tr> <td><input type="checkbox"/> <i>ISPD</i> Gene Sequencing</td> <td><input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2O (<i>POMGnTI</i> sequencing)</td> </tr> <tr> <td><input type="checkbox"/> Limb Girdle Muscular Dystrophy, autosomal recessive common mutation panel</td> <td><input type="checkbox"/> Muscle-Eye-Brain Disease (<i>POMGnTI</i> sequencing)</td> </tr> <tr> <td><input type="checkbox"/> Limb Girdle Muscular Dystrophy, autosomal recessive common mutation panel with <i>FKRP</i> sequencing</td> <td><input type="checkbox"/> Myotonic Dystrophy type 1 (DM1)</td> </tr> <tr> <td><input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2B (<i>Dysferlin</i> sequencing)</td> <td><input type="checkbox"/> Walker Warburg Syndrome (<i>POMTI</i>, <i>POMT2</i>, <i>POMGnTI</i>, <i>FKTN</i>, <i>FKRP</i>, <i>ISPD</i> and <i>LARGE</i> sequencing)</td> </tr> <tr> <td><input type="checkbox"/></td> <td><input type="checkbox"/></td> </tr> <tr> <td><input type="checkbox"/></td> <td><input type="checkbox"/></td> </tr> </table>				<input type="checkbox"/> Congenital Muscular Dystrophy type 1C, MDC1C (<i>FKRP</i> sequencing)	<input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2I (<i>FKRP</i> sequencing)	<input type="checkbox"/> Congenital Muscular Dystrophy type 1D, MDC1D (<i>LARGE</i> sequencing)	<input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2K (<i>POMTI</i> sequencing)	<input type="checkbox"/> Fukuyama Congenital Muscular Dystrophy (<i>FKTN</i> sequencing)	<input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2M (<i>FKTN</i> sequencing)	<input type="checkbox"/> Fukuyama Congenital Muscular Dystrophy (Japanese Founder Mutation PCR)	<input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2N (<i>POMT2</i> sequencing)	<input type="checkbox"/> <i>ISPD</i> Gene Sequencing	<input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2O (<i>POMGnTI</i> sequencing)	<input type="checkbox"/> Limb Girdle Muscular Dystrophy, autosomal recessive common mutation panel	<input type="checkbox"/> Muscle-Eye-Brain Disease (<i>POMGnTI</i> sequencing)	<input type="checkbox"/> Limb Girdle Muscular Dystrophy, autosomal recessive common mutation panel with <i>FKRP</i> sequencing	<input type="checkbox"/> Myotonic Dystrophy type 1 (DM1)	<input type="checkbox"/> Limb Girdle Muscular Dystrophy type 2B (<i>Dysferlin</i> sequencing)	<input type="checkbox"/> Walker Warburg Syndrome (<i>POMTI</i> , <i>POMT2</i> , <i>POMGnTI</i> , <i>FKTN</i> , <i>FKRP</i> , <i>ISPD</i> and <i>LARGE</i> sequencing)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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<input type="checkbox"/>	<input type="checkbox"/>																						
<input type="checkbox"/>	<input type="checkbox"/>																						
PART D – BILLING – REQUIRED		NOTE: Claims can't be submitted to a Medicaid Program Outside of Iowa																					
On date of collection, was patient: _____		Admission Date: _____ Discharge Date: _____																					
<input type="checkbox"/> Inpatient <input type="checkbox"/> Outpatient <input type="checkbox"/> Non-Hospital Patient - Facility name where specimen collected: _____																							
<input type="checkbox"/> Bill Client Email Recipient for Invoicing _____ Name: _____ Phone: _____																							
<input type="checkbox"/> Bill Insurance PRE-AUTHORIZATION #: _____ Required for all cases organized outside of Iowa. See website for a comprehensive guide to UIDL Billing - https://www.healthcare.uiowa.edu/uidl/site/billing_services/index.html																							
Primary Insurance Coverage Information		Secondary Insurance Coverage Information																					
Insured by: _____		Insured by: _____																					
Claims Address: _____		Claims Address: _____																					
City: _____ St: _____ Zip: _____		City: _____ St: _____ Zip: _____																					
Policy/ID #: _____ Group #: _____		Policy/ID #: _____ Group #: _____																					
Name of Subscriber: _____ DOB: _____		Name of Subscriber: _____ DOB: _____																					
Relationship to Patient: _____		Relationship to Patient: _____																					
<small>Medicare will only pay for the services that it determines to be "reasonable and necessary" under section 1862(a)(1) of the Medicare Law. If Medicare determines that a particular service, although it would otherwise be covered, is not "reasonable and necessary" under the Medicare standards, Medicare will deny payment for that service or test.</small>																							

The purpose of this form is to obtain information necessary for UIDL Pathology Department to perform consultations and/or testing.
 Failure to properly complete the form may cause delay in the processing of specimens.