

## MUSCULAR DYSTROPHY MOLECULAR GENETICS REQUISITION INTERNATIONAL

UI Diagnostic Laboratories Department of Pathology 200 Hawkins Drive, 5231 RCP Iowa City, Iowa 52242

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Toll Free: 866-844-2522 Local: 319-384-7212

FOR UIDL USE ONLY: MRN# PATH# FOR CLIENT USE ONLY: Requisition Date Completed by: PART A – PATIENT INFORMATION – Required PART B - PROVIDER INFORMATION - Required Patient Last Name: Referring Institution: Patient First Name: Street: Street: City: St: Zip: St: City: Zip: Phone: Fax: Phone: Fax: Referring Physician: Date of Birth: Gender:  $\square M \square F$ Email: Phone: PART C - SPECIMEN INFORMATION (Complete the appropriate information below or include in an accompanying letter.) Ship on Mondays or Tuesdays only, to avoid potential weekend delays. Ship at ambient temperature Priority Overnight. MATERIALS SUBMITTED: ☐ EDTA Whole Blood  $\square$  DNA ☐ Other (Specify): \_\_\_\_\_ \_\_\_\_SPECIMEN COLLECTION TIME: \_\_\_\_ SPECIMEN COLLECTION DATE: \_\_\_\_\_ PERTINENT CLINICAL HISTORY AND FINDINGS (please attach pedigree if available): 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. SELECT TEST: Refer to the UIDL TEST DIRECTORY for specimen requirements and CPT Codes at https://www.healthcare.uiowa.edu/path\_handbook/rindex.html ☐ Congenital Muscular Dystrophy type 1C, MDC1C (*FKRP* sequencing) ☐ Limb Girdle Muscular Dystrophy type 2I (*FKRP* sequencing) ☐ Congenital Muscular Dystrophy type 1D, MDC1D (*LARGE* sequencing) ☐ Limb Girdle Muscular Dystrophy type 2K (*POMT1* sequencing) ☐ Fukuvama Congenital Muscular Dystrophy (FKTN sequencing) ☐ Limb Girdle Muscular Dystrophy type 2M (FKTN sequencing)  $\square$  Fukuyama Congenital Muscular Dystrophy (Japanese Founder Mutation PCR) ☐ Limb Girdle Muscular Dystrophy type 2N (*POMT2* sequencing) ☐ Limb Girdle Muscular Dystrophy type 2O (*POMGnT1* sequencing) ☐ *ISPD* Gene Sequencing ☐ Limb Girdle Muscular Dystrophy, autosomal recessive ☐ Muscle-Eve-Brain Disease (*POMGnT1* sequencing) common mutation panel ☐ Limb Girdle Muscular Dystrophy, autosomal recessive ☐ Myotonic Dystrophy type 1 (DM1) common mutation panel with FKRP sequencing ☐ Limb Girdle Muscular Dystrophy type 2B (*Dysferlin* sequencing) ☐ Walker Warburg Syndrome (*POMT1*, *POMT2*, *POMGnT1*, *FKTN*, FKRP, ISPD and LARGE sequencing) PART D - BILLING - REQUIRED Complete this section for all international samples. UIDL requires international specimens to be accompanied by payment in US dollars. If prepayment is not received, there will be a delay in result reporting. Please call our Billing Department at 00-1-319-353-7958 or email your questions to UIDL@healthcare.uiowa.edu. **Pre-Payment Required** ☐ Bank Check Enclosed (made payable to UI Diagnostic Laboratories) ☐ Credit Card: ☐ Visa ☐ MasterCard Card # \_\_\_\_\_-\_\_\_\_ Expires: \_\_\_\_/\_\_\_ (MM/YY) Security Code: \_\_\_\_\_ Name of Cardholder (print first name, last name): Phone: Cardholder Address: \_\_\_

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.