

**THIS IS NOT A TEST REQUEST FORM.**

Please fill out this form and submit it with the test request form or electronic packing list.

**PATIENT HISTORY FOR LAMINOPATHIES (LMNA) GENE TESTING**

**Patient Name:** \_\_\_\_\_ **Date of Birth:** \_\_\_\_\_ **Sex:**  Female  Male

**Physician:** \_\_\_\_\_ **Physician Phone:** \_\_\_\_\_

**Practice Specialty:** \_\_\_\_\_ **Physician Fax:** \_\_\_\_\_

**Genetic Counselor:** \_\_\_\_\_ **Counselor Phone:** \_\_\_\_\_

**Patient's Ethnicity** (check all that apply)

- African American/Black     Asian     Hispanic     Native American  
 Ashkenazi Jewish     Caucasian/White     Middle Eastern     Other: \_\_\_\_\_

**Clinical Diagnosis**

- Atypical Werner syndrome     Emery-Dreifuss muscular dystrophy     Limb girdle muscular dystrophy  
 Charcot-Marie-Tooth     Familial partial lipodystrophy     Mandibulo-acral dysplasia  
 Dilated cardiomyopathy     Hutchinson-Gilford progeria     Restrictive Dermopathy  
 Other: \_\_\_\_\_

**Does the patient have symptoms?**     No     Yes (check all that apply and describe)

- |   |   |  |
|---|---|--|
| <input type="checkbox"/> Absent tendon reflexes                                 | <input type="checkbox"/> Failure to Thrive                            | <input type="checkbox"/> Mottled cutaneous pigmentation      |
| <input type="checkbox"/> Alopecia   | <input type="checkbox"/> Fetal akinesia                               | <input type="checkbox"/> Premature aging                     |
| <input type="checkbox"/> Cardiac conduction defects                             | <input type="checkbox"/> Growth retardation                           | <input type="checkbox"/> Progressive muscle weakness/wasting |
| <input type="checkbox"/> Cataracts  | <input type="checkbox"/> Hypogonadism                                 | <input type="checkbox"/> Progressive ventricular dilation    |
| <input type="checkbox"/> Craniofacial and skeletal anomalies;<br>specify: _____ | <input type="checkbox"/> Impaired systolic function                   |  |
| _____   | <input type="checkbox"/> Joint contractures                           |  |
| <input type="checkbox"/> Distal muscle weakness                                 | <input type="checkbox"/> Joint degeneration                           |  |
| <input type="checkbox"/> Excess fat on face/neck                                | <input type="checkbox"/> Loss of subcutaneous fat<br>from extremities |  |
| <input type="checkbox"/> Other symptom(s): _____                                |   |  |

**Has the patient undergone previous DNA testing for a laminopathy?**     No     Yes     Unknown

If yes, describe the genes, disorder, methodology, and results: \_\_\_\_\_

**Is there any relevant family history?**     No     Yes     Unknown

If yes, attach a pedigree or specify the relative's relationship to the patient. List their symptoms and age of onset: \_\_\_\_\_

**Has DNA testing been performed for the family member(s)?**     No     Yes     Unknown

If yes, attach a copy of the relative's DNA laboratory result (REQUIRED for familial mutation testing).

**Check the test you intend to order.**

- 2004543 LMNA-Related Disorders (LMNA) Sequencing:** Sequence analysis of LMNA coding regions; clinical sensitivity is dependent upon the specific LMNA-related disorder.
- 2001961 Familial Mutation, Targeted Sequencing:** Tests for a mutation previously identified in a family member; a copy of relative's lab result is REQUIRED

<p><b>Master Label</b></p>
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**For questions, contact an ARUP genetic counselor at 800-242-2787 ext. 2141**