

Date _____

Patient Information

1

Patient Name: _____ Date of Birth: ____ / ____ / ____

Phone: _____ Cell: _____ E-mail: _____

Insurance - a copy of the patient's insurance card is required* Please expedite genetic counseling for immediate management decisions (2-4 business days)

Reason for Referral

Personal and/or Family History (known diagnosis):

- | PATIENT | FAMILY MEMBER | |
|--------------------------|--------------------------|---|
| <input type="checkbox"/> | <input type="checkbox"/> | Duchenne or Becker muscular dystrophy |
| <input type="checkbox"/> | <input type="checkbox"/> | Myotonic dystrophy (type 1 or 2) |
| <input type="checkbox"/> | <input type="checkbox"/> | Other muscular dystrophy (i.e. Limb-girdle, Emery Dreifuss) |
| <input type="checkbox"/> | <input type="checkbox"/> | Charcot-Marie-Tooth |
| <input type="checkbox"/> | <input type="checkbox"/> | Alzheimer's disease (suspected or known family history, and/or onset <60 years) |
| <input type="checkbox"/> | <input type="checkbox"/> | Parkinson disease (suspected or known family history, and/or onset <50 years) |
| <input type="checkbox"/> | <input type="checkbox"/> | Frontotemporal dementia (suspected or known family history) |
| <input type="checkbox"/> | <input type="checkbox"/> | Amyotrophic lateral sclerosis (suspected or known family history) |
| <input type="checkbox"/> | <input type="checkbox"/> | Hereditary ataxia |
| <input type="checkbox"/> | <input type="checkbox"/> | Other neuromuscular, neurodegenerative, or neurometabolic disease |
| <input type="checkbox"/> | <input type="checkbox"/> | Known gene mutation/neurogenetic condition. Specify: _____ |
| <input type="checkbox"/> | <input type="checkbox"/> | Other: _____ |

2

Personal and/or Family History (symptoms):

- | PATIENT | FAMILY MEMBER | |
|--------------------------|--------------------------|----------------------------------|
| <input type="checkbox"/> | <input type="checkbox"/> | Young-onset dementia (<60 years) |
| <input type="checkbox"/> | <input type="checkbox"/> | Ataxia, non-acquired |
| <input type="checkbox"/> | <input type="checkbox"/> | Cerebellar atrophy |
| <input type="checkbox"/> | <input type="checkbox"/> | Non-acquired neuropathy |
| <input type="checkbox"/> | <input type="checkbox"/> | Other: _____ |

Genetic Test Status

- Test not yet ordered
- Test ordered
- Results received, please provide results interpretation
- Unknown
- Other: _____

Patient Documentation - Fax with Referral

3

a. Clinical. Please include the following (if performed)

- | | |
|--|---|
| <input type="checkbox"/> Clinic note outlining history of disease/suspected diagnosis | <input type="checkbox"/> Genetic test results |
| <input type="checkbox"/> Lab/imaging results (i.e. creatine kinase, brain MRI, EMG, and/or muscle biopsy).** | <input type="checkbox"/> Copy of mutation in family (if requesting carrier testing) |

b. Insurance documentation. A copy of front and back of the patient's insurance card.

* Used to verify the patient's benefits. | ** We will not provide interpretation

Physician Information

Medical Center/Practice

Primary Provider

NPI

Practice Contact

Phone

Fax

4

E-mail

Address

City

State

Zip

Secondary Provider*

NPI

Referring Physician's Signature

* Will receive copy of Genetic Counseling Summary Report.

Fax completed form to:

(760) 203-1194

To refer by phone
800.975.4819

To refer by email
referral@informedDNA.com

www.informedDNA.com

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5