



## Patient Information

1 Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_  
 PREFERRED Phone: \_\_\_\_\_ OTHER Phone: \_\_\_\_\_ E-mail: \_\_\_\_\_

## Billing

2  Bill to Patient Insurance  Other (Please Explain) \_\_\_\_\_  
 \* In-network only.

## 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: _____  |

### 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

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

- |  |   |
|--|---|
| <input type="checkbox"/> Clinic note outlining history of disease/suspected diagnosis                        | <input type="checkbox"/> Patient 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. Patient face sheet (Demographics).

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

\*\* We will not provide interpretation.

## Provider Information

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Medical Center/Practice		Practice Contact	
Phone	Fax	E-mail	
Address		City	State Zip
Referring Provider		Fax (required)	
NPI		Referring Provider's Signature	

## Fax completed form to:

6 (760)203-1194

www.InformedDNA.com

For questions, please call  
**800-975-4819**