MNG LABORATORIES
Neurogenetic Answers™

5424 Glenridge Drive NE
Atlanta, GA 30342 USA
toll-free: 678.225.0222
fax: 678.225.0212
mnglabs.com

Patient and Specimen Information Form

We gladly accept deliveries Monday-Saturday, excluding holidays
CLIA License #11D0703390; CAP License #1441004; State of Georgia License #060-381

Patient and Specimen Information

Patient Last Name	Patient First Name
Patient ID #	Date of Birth [MM/DD/YYYY]
Diagnosis/ICD-10	Collection Date [MM/DD/YYYY]

Gender ☐ Male ☐ Female

Specimen Type
☐ Whole Blood ☐ Buccal Swab ☐ CSF ☐ Urine ☐ Fibroblasts ☐ Skin [For Culture] ☐ Plasma ☐ Muscle ☐ DNA Tissue:_____

Please complete and include our clinical information form, or attach clinical notes

Referring Physician Information

Referring Physician Name	Print	Signature
Referring Physician **NPI # [Required]** or international equivalent		
Facility / Organization		Phone
Select and Provide Email or Fax for Report Delivery	☐ Email	☐ Fax

[REQUIRED] Billing Information

Facility Responsible for Payment		Phone
Facility Contact Person		Fax
Facility Contact Person Email	Email	
Facility Billing Address		
City, State, Zip Code		

Results (sent by secure HIPAA-compliant email or fax)

Authorized Recipient Name	Authorized Recipient Name
Facility	Facility
Phone	Phone
☐ Fax ☐ Email	☐ Fax ☐ Email

Forms Checklist

All of the following are required before we will process your orders (please check the following):

- ☐ All specimens that will be analyzed must be received
- ☐ Clinical Information Form completed
- ☐ Informed Consent for Genetic Testing completed and signed

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Single Gene, mtDNA & Repeat Expansion Requisition Form

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Patient Name _____ **DOB** _____

STAT Testing Now Available

For STAT Testing, please see page 7.

All genes available for sequencing are listed in alphabetical order. Genes which have associated Deletion/Duplication [Del/Dup] analysis are highlighted gray. Note that some genes listed below only have Del/Dup analysis, but sequencing may be orderable by custom next-generation sequencing.

Mitochondrial DNA Genetic Testing

- ☐ (MOL001) Mitochondrial DNA Depletion Testing (Muscle)
- ☐ (MOL002) Mitochondrial DNA Deletion Analysis
- ☐ (MOL021) Mitochondrial Genome Sequencing
- ☐ (MOL189) Mitochondrial Genome Sequencing + Deletion Analysis
- ☐ (MOL232) Mitochondrial Genome Sequencing + Deletion Analysis + Depletion Testing (Leukocyte)
- ☐ (MOL334) Mitochondrial Depletion Testing (Leukocyte)
- ☐ (MOL340) Mitochondrial Genome Sequencing + Deletion Analysis + Depletion Testing (Muscle)

Repeat Expansions

- ☐ (MOL299) Myotonic Dystrophy 1 (DMPK) Genetic Testing (Repeat Expansion)
- ☐ (MOL303) Myotonic Dystrophy 2 (ZNF9/CNBP) Genetic Testing (Repeat Expansion)
- ☐ (MOL364) C9orf72 Genetic Testing (Repeat Expansion)
- ☐ (MOL366) Huntington Disease (HTT) Genetic Testing (Repeat Expansion)
- ☐ (MOL381) Fragile X Repeat Expansion + Methylation Analysis
- ☐ (MOL379) Spinocerebellar Ataxia Repeat Expansion Panel (SCA 8, 10, 12, 17, 36 & DRPLA)
- ☐ (MOL380) Comprehensive Spinocerebellar Ataxia Repeat Expansion Panel (SCA 1, 2, 3, 6, 7, 8, 10, 12, 17, 36 & DRPLA)
- ☐ (MOL391) Comprehensive Ataxia Repeat Expansion Panel (SCA 1, 2, 3, 6, 7, 8, 10, 12, 17, 36, DRPLA & FRDA)
- ☐ (MOL259) Friedreich Ataxia Genetic Testing (Repeat Expansion)
- ☐ SCA1/ATXN1 (MOL368)
- ☐ SCA2/ATXN2 (MOL369)
- ☐ SCA3/ATXN3 (MOL370)
- ☐ SCA6/CACNA1A (MOL371)
- ☐ SCA7/ATXN7 (MOL372)
- ☐ SCA8/ATXN8 (MOL373)
- ☐ SCA10/ATXN10 (MOL374)
- ☐ SCA12/PPP2R2B (MOL375)
- ☐ SCA17/TBP (MOL376)
- ☐ SCA36/NOP56 (MOL377)
- ☐ DRPLA/ATN1 (MOL378)

Single Genes + MLPA

- ☐ ABCB7 (MOL026)
- ☐ ACAD8 MOL242)
- ☐ ACAD9 (MOL243)
- ☐ ACADM (MOL027)
- ☐ ACADM Mutation Screen: c.985A>G, K329E (MOL023)
- ☐ ACADS (MOL028)
- ☐ ACADVL (MOL029)
- ☐ ACADVL + MLPA [Del/Dup] (MOL261)
- ☐ ACADVL MLPA [Del/Dup Only] (MOL260)
- ☐ ADAR (MOL309)
- ☐ ADCK3 (MOL140)
- ☐ ADSL (MOL209)
- ☐ AIFM1 (MOL183)
- ☐ ALDH5A1 (MOL125)
- ☐ ALDH7A1 (MOL030)
- ☐ AMPD1 (MOL185)
- ☐ APTX (MOL110)
- ☐ ARHGEF9 (MOL306)
- ☐ ATL1/SPG3 + MLPA [Del/Dup] (MOL267)
- ☐ ATL1/SPG3 MLPA [Del/Dup Only] (MOL266)
- ☐ ATP1A2 (MOL032)
- ☐ ATP1A2 MLPA [Del/Dup Only] (MOL214)
- ☐ ATP5E (MOL295)
- ☐ ATPAF2 (MOL112)
- ☐ BCS1L (MOL045)
- ☐ BRAF (MOL147)
- ☐ BSND (MOL271)
- ☐ C10ORF2 (MOL073)
- ☐ C10ORF2 MLPA [Del/Dup Only] (MOL313)
- ☐ CACNA1A + [Del/Dup](MOL033)
- ☐ CACNA1A [Del/Dup Only] (MOL208)
- ☐ CACNA1S (MOL307)
- ☐ CACNB4 (MOL227)
- ☐ CAPN3 (MOL034)
- ☐ CAPN3 + MLPA [Del/Dup] (MOL240)
- ☐ CAPN3 MLPA [Del/Dup Only] (MOL239)
- ☐ CAV3 (MOL035)
- ☐ CHKB (MOL254)
- ☐ CLCN1 (MOL355)
- ☐ CLCNKA (MOL268)
- ☐ CLCNKB + MLPA [Del/Dup] (MOL287)
- ☐ CLCNKB MLPA [Del/Dup Only] (MOL286)
- ☐ COQ2 (MOL046)
- ☐ COQ3) MOL128)
- ☐ COQ4 (MOL129)
- ☐ COQ6 (MOL131)
- ☐ COQ7 (MOL132)
- ☐ COQ9 (MOL133)

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Single Genes + MLPA

- ☐ COX10 (MOL047)
- ☐ COX15 (MOL048)
- ☐ COX6B1 (MOL182)
- ☐ CPT1A (MOL049)
- ☐ CPT2 (MOL050)
- ☐ CYP21A2 MLPA [Del/Dup] (MOL388)
- ☐ DARS (MOL339)
- ☐ DARS2 (MOL094)
- ☐ DBH (MOL141)
- ☐ DDC (MOL025)
- ☐ DEPDC5 (MOL354)
- ☐ DGUOK (MOL051)
- ☐ DHFR (MOL237)
- ☐ DLAT (MOL224)
- ☐ DLD (MOL225)
- ☐ DOK7 (MOL281)
- ☐ DPYS (MOL342)
- ☐ DYSF (MOL052)
- ☐ DMD MLPA [Del/Dup] (MOL302)
- ☐ EIF2B1 (MOL054)
- ☐ EIF2B2 (MOL055)
- ☐ EIF2B3 (MOL056)
- ☐ EIF2B4 (MOL057)
- ☐ EIF2B5 (MOL058)
- ☐ EMD (MOL145)
- ☐ ETFA (MOL163)
- ☐ ETFB (MOL164)
- ☐ ETFDH (MOL142)
- ☐ FASTKD2 (MOL165)
- ☐ FKRP (MOL149)
- ☐ FKRP MLPA [Del/Dup Only] (MOL314)
- ☐ FKTN (MOL150)
- ☐ FOLR1 (MOL166)
- ☐ FOXP2 (MOL238)
- ☐ FOXRED1 (MOL210)
- ☐ FXN (MOL059)
- ☐ GAMT (MOL351)
- ☐ GARS (MOL167)
- ☐ GARS MLPA [Del/Dup Only] (MOL315)
- ☐ GBA MLPA [Del/Dup Only] (MOL389)
- ☐ GATM (MOL352)
- ☐ GCDH (MOL213)
- ☐ GCH1 (MOL060)
- ☐ GCH1 + MLPA [Del/Dup] (MOL234)
- ☐ GCH1 MLPA [Del/Dup Only] (MOL215)
- ☐ GFAP (MOL122)
- ☐ GFER (MOL155)
- ☐ GFM1 (MOL113)
- ☐ GJB1 (MOL332)
- ☐ GJB1 + MLPA [Del/Dup] (MOL333)
- ☐ GJB1 MLPA [Del/Dup Only] (MOL277)
- ☐ GLDC (MOL212)
- ☐ GLDC + MLPA [Del/Dup] (MOL236)
- ☐ GLDC MLPA [Del/Dup Only] (MOL219)
- ☐ GLRA1 (MOL116)
- ☐ GLRB (MOL120)
- ☐ GLUL (MOL308)
- ☐ GNE (MOL341)
- ☐ GPHN (MOL251)
- ☐ GYS2 (MOL061)
- ☐ HADHA (MOL062)
- ☐ HADHA Mutation Screen: c.1528G>C, E510Q (MOL022)
- ☐ HADHB (MOL063)
- ☐ HBA1 & HBA2 [Del/Dup Only] (MOL390)
- ☐ HCCS (MOL146)
- ☐ HSPB1 MLPA [Del/Dup Only] (MOL316)
- ☐ HSPD1 (MOL086)
- ☐ KCNA1 (MOL064)
- ☐ KCNJ1 (MOL270)
- ☐ KCNQ2 (MOL229)
- ☐ KCNQ2 + MLPA [Del/Dup] (MOL330)
- ☐ KCNQ2 MLPA [Del/Dup Only] (MOL216)
- ☐ LAMA2 + [Del/Dup] (MOL065)
- ☐ LAMA2 MLPA [Del/Dup Only] (MOL317)
- ☐ LITAF (MOL335)
- ☐ LMNA (MOL066)
- ☐ LRPPRC (MOL115)
- ☐ MECP2 (MOL067)
- ☐ MECP2 + MLPA [Del/Dup] (MOL257)
- ☐ MECP2 MLPA [Del/Dup Only] (MOL187)
- ☐ MFN2 (MOL114)
- ☐ MFN2 MLPA [Del/Dup Only] (MOL293)
- ☐ MPV17 (MOL068)
- ☐ MPV17 MLPA [Del/Dup Only] (MOL318)
- ☐ MPZ (MOL250)
- ☐ MPZ MLPA [Del/Dup Only] (MOL278)
- ☐ MRPS16 (MOL099)
- ☐ MTHFR (MOL171)
- ☐ MTM1 (MOL139)
- ☐ MTM1 + MLPA [Del/Dup] (MOL290)
- ☐ MTM1 MLPA [Del/Dup Only] (MOL289)
- ☐ MYOT (MOL144)
- ☐ NDUFA1 (MOL102)
- ☐ NDUFA11 (MOL161)
- ☐ NDUFA2 (MOL160)
- ☐ NDUFAF1 (MOL106)
- ☐ NDUFAF2 (MOL101)
- ☐ NDUFAF3 (MOL158)
- ☐ NDUFAF4 (MOL162)
- ☐ NDUFAF5 (MOL151)
- ☐ NDUFS1 (MOL036)
- ☐ NDUFS2 (MOL103)
- ☐ NDUFS3 (MOL104)
- ☐ NDUFS4 (MOL037)
- ☐ NDUFS6 (MOL157)
- ☐ NDUFS7 (MOL038)
- ☐ NDUFS8 (MOL039)
- ☐ NDUFV1 (MOL040)
- ☐ NDUFV2 (MOL105)
- ☐ NEFL MLPA [Del/Dup Only] (MOL319)
- ☐ NUBPL (MOL211)
- ☐ OPA1 (MOL069)
- ☐ OPA1 + MLPA [Del/Dup] (MOL292)
- ☐ OPA1 MLPA [Del/Dup Only] (MOL291)
- ☐ PAH (MOL349)

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- ☐ PC (MOL226)
- ☐ PDHA1 (MOL070)
- ☐ PDHA1 + MLPA [Del/Dup] (MOL235)
- ☐ PDHA1 MLPA [Del/Dup Only] (MOL217)
- ☐ PDHB (MOL222)
- ☐ PDP1 (MOL223)
- ☐ PDSS1 (MOL071)
- ☐ PDSS2 (MOL072)
- ☐ PDXK (MOL274)
- ☐ PEX6 (MOL360)
- ☐ PFKM (MOL272)
- ☐ PGAM2 (MOL253)
- ☐ PHOX2B (MOL184)
- ☐ PKD1/PKD2 MLPA [Del/Dup Only] (MOL350)
- ☐ PMP22 (MOL344)
- ☐ PMP22 + MLPA [Del/Dup](MOL345)
- ☐ PMP22 MLPA [Del/Dup Only] (MOL276)
- ☐ PNPO (MOL074)
- ☐ POLG + [Del/Dup] (MOL075)
- ☐ POLG MLPA [Del/Dup Only] (MOL294)
- ☐ POLG2 (MOL109)
- ☐ POMT1 (MOL153)
- ☐ POMT2 (MOL143)
- ☐ PROSC (MOL367)
- ☐ PRX MLPA [Del/Dup Only] (MOL320)
- ☐ PTS (MOL096)
- ☐ PUS1 (MOL107)
- ☐ PYGM (MOL076)
- ☐ PYGM Mutation Screen: c.148C>T, R50X; c.613G>A, G205S; c.1628A>C, K543T (MOL024)
- ☐ QDPR (MOL117)
- ☐ RAB7A MLPA [Del/Dup Only] (MOL321)
- ☐ RARS2 (MOL152)
- ☐ RNASEH2A (MOL124)
- ☐ RNASEH2B (MOL118)
- ☐ RNASEH2C (MOL168)
- ☐ RRM2B (MOL077)
- ☐ RRM2B MLPA [Del/Dup Only] (MOL322)
- ☐ RYR1 + [Del/Dup] (MOL154)
- ☐ SAMHD1 (MOL172)
- ☐ SCN10A (MOL346)
- ☐ SCN11A (MOL347)
- ☐ SCN1A + [Del/Dup] (MOL248)
- ☐ SCN1A MLPA [Del/Dup Only] (MOL218)
- ☐ SCN1B (MOL255)
- ☐ SCN4A (MOL356)
- ☐ SCN5A (MOL348)
- ☐ SCN9A + [Del/Dup] (MOL258)
- ☐ SCO1 (MOL082)
- ☐ SCO2 (MOL083)
- ☐ SDHA (MOL041)
- ☐ SDHB (MOL042)
- ☐ SDHC (MOL043)
- ☐ SDHD (MOL044)
- ☐ SEPT9 + [Del/Dup] (MOL336)
- ☐ SGCA (MOL078)
- ☐ SGCA MLPA [Del/Dup Only] (MOL324)
- ☐ SGCB (MOL079)
- ☐ SGBC MLPA [Del/Dup Only] (MOL325)
- ☐ SGCD (MOL080)
- ☐ SGCD MLPA [Del/Dup Only] (MOL326)
- ☐ SGCE (MOL247)
- ☐ SGCE + MLPA [Del/Dup] (MOL283)
- ☐ SGCE MLPA [Del/Dup Only] (MOL282)
- ☐ SGCG (MOL081)
- ☐ SGCG MLPA [Del/Dup Only] (MOL327)
- ☐ SLC12A3 (MOL233)
- ☐ SLC12A3 + MLPA [Del/Dup] (MOL245)
- ☐ SLC12A3 MLPA [Del/Dup Only] (MOL244)
- ☐ SLC17A5 (MOL170)
- ☐ SLC18A2 (MOL095)
- ☐ SLC1A3 (MOL228)
- ☐ SLC22A5 MLPA [Del/Dup Only](MOL262)
- ☐ SLC25A19 (MOL169)
- ☐ SLC25A20 (MOL085)
- ☐ SLC25A4 (MOL031)
- ☐ SLC25A4 MLPA [Del/Dup Only] (MOL328)
- ☐ SLC2A1 (MOL121)
- ☐ SLC2A1 + MLPA [Del/Dup] (MOL231)
- ☐ SLC2A1 MLPA [Del/Dup Only] (MOL186)
- ☐ SLC6A3 (MOL097)
- ☐ SLC6A4 (MOL252)
- ☐ SLC6A5 (MOL127)
- ☐ SLC6A8 (MOL246)
- ☐ SMN1/SMN2 Spinal Muscular Atrophy (MOL301) MLPA [Del/Dup Only]
- ☐ SPAST (MOL256)
- ☐ SPAST + MLPA [Del/Dup] (MOL265)
- ☐ SPAST MLPA [Del/Dup Only] (MOL264)
- ☐ SPG11 MLPA [Del/Dup Only] (MOL343)
- ☐ SPG7 (MOL087)
- ☐ SPR (MOL126)
- ☐ SPTLC1 (MOL331)
- ☐ SUCLA2 (MOL088)
- ☐ SUCLG1 (MOL089)
- ☐ SURF1 (MOL090)
- ☐ TACO1 (MOL241)
- ☐ TAZ (MOL111)
- ☐ TCAP (MOL148)
- ☐ TH (MOL091)
- ☐ TIMM8A (MOL098)
- ☐ TK2 (MOL305)
- ☐ TK2 MLPA [Del/Dup Only] (MOL329)
- ☐ TMEM126A (MOL156)
- ☐ TMEM70 (MOL173)
- ☐ TNNI2 (MOL159)
- ☐ TPH2 (MOL092)
- ☐ TPM3 (MOL137)
- ☐ TREX1 (MOL119)
- ☐ TUBB4A (MOL362)
- ☐ TUFM (MOL108)
- ☐ TYMP (MOL053)
- ☐ UBE3A (MOL093)
- ☐ UQCRB (MOL100)
- ☐ UQCRQ (MOL174)
- ☐ ZMPSTE24 (MOL138)

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Patient Name _____ **DOB** _____ **Gender** ☐ Male ☐ Female

Clinical (Check All That Apply)

Eye ☐ Retinitis Pigmentosa ☐ Optic Atrophy ☐ Other

Hearing ☐ Sensorineural ☐ Stickler ☐ Usher

Neuronal Migration ☐ Meckel ☐ Joubert ☐ Other

☐ Stroke

Cognitive/Neurobehavioral ☐ Intellectual Disability (ID) ☐ Syndromic ID ☐ Nonsyndromic ID ☐ Autism ☐ Dementia

Movement Disorders ☐ Ataxia ☐ Episodic Ataxia ☐ Dystonia ☐ Chorea/Athetosis ☐ Parkinson Disease ☐ L-Dopa Response

Epilepsy ☐ Myoclonic ☐ Other ☐ Absence ☐ Tonic Clonic ☐ Epileptic Encephalopathy

Spasticity ☐ Spastic Paraplegia ☐ Other ☐ Spastic Quadriplegia

Connective Tissue & Bone ☐ Ehlers Danlos ☐ Marfan ☐ Aneurysms ☐ Other

Neuromuscular ☐ Distal ☐ Proximal ☐ Muscle Atrophy ☐ Contractures ☐ Malignant Hyperthermia ☐ Arthrogryposis ☐ Rhabdomyolysis ☐ Periodic Paralysis ☐ Statin Use ☐ Myasthenia

Nerve/Anterior Horn Cell ☐ Neurofibromas ☐ Charcot-Marie-Tooth ☐ Sensory ☐ Autonomic ☐ Pain ☐ Motor ☐ Nerve Conduction ☐ Other

Cardiomyopathy ☐ Dilated ☐ Hypertrophic ☐ Noncompaction

Arrhythmias ☐ Ventricular Tachycardia ☐ Brugada ☐ Long or Short QT ☐ Conduction Defect

Congenital Heart Defects ☐ Heterotaxy ☐ Other

Endocrine ☐ Hypothyroidism ☐ Other ☐ Diabetes Mellitus

Additional Comments

Imaging (Check All That Apply)

Brain MRI

☐ Leigh Disease ☐ Basal Ganglia Calcification ☐ Stroke ☐ Cerebellar Atrophy ☐ Abnormal Myelin (describe)

EEG (Describe Findings)

EMG/NVC (Describe Findings)

Laboratory

Metabolic (Describe Findings)

CPK Maximum _____
Minimum _____

Genetic (Describe Findings)

☐ Chromosome Microarray
☐ Deletion/Insertion Testing
☐ Other (comment)

Family History

Affected Maternal Lineage	Affected Paternal Lineage	Siblings
Relationship to Proband	Relationship to Proband	Number (specify gender)
Symptoms	Symptoms	Healthy/Affected

Ethnicity (please check)

☐ South Asian ☐ East Asian ☐ European (Non-Finnish) ☐ European (Finnish) ☐ Latino ☐ African ☐ Other (comment)

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Informed Consent for Genetic Testing

In compliance with New York State Civil Law: Section 79-L

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Please provide a copy of completed consent with sample and requisition. Failure to do so may delay testing.

When signed and dated, this written consent is written authorization to participate in genetic testing.

1. **Purpose of the Test:** My physician has explained the recommended testing: _____ (name of test or MNG test code), which is performed to help diagnose _____ _____(insert disease description). I am aware that all documentation regarding this testing, including the description of the purpose, methodology, and disorders is freely available at www.mnglabs.com/tests and has either been reviewed with me by my physician or I have read the documentation on my own. **Patient (or parent/guardian) initials: _____**

2. **Statement Regarding Test Result:** A positive test result is an indication that the individual has a genetic cause for the specific disease tested for. A negative result may/may not rule out a genetic disorder depending on clinical history and quality/type of specimen tested. The individual may wish to consider further independent testing, consult a personal physician or pursue genetic counseling.

3. **Level of Certainty:** Is test-specific and determined by the methods employed, patient's clinical history and sometimes by the nature of the patient's condition at time of sampling. There is always a small possibility of error or failure in sample analysis; this is true with complex testing in any laboratory. Inclusion of clinical data, such as medical history, family history, images as they relate to the disease or disorder, will decrease the level of uncertainty in an interpretation and are encouraged to be included when submitting samples for analysis. MNG Laboratories will keep personal information private in accordance with HIPAA laws.
   I consent to the retention of these documents by MNG Laboratories in their database.
   Patient (or parent/guardian) Initials: _____

4. **Disclosing Test Results:** The following categories of persons or organizations that test results may be released to include, but are not limited to: hospitals or laboratories involved in the patient's care, referring physician(s) and primary care providers, other physician groups (consultants, surgeons), insurance companies (as provided by the patient or referring physician for payment purposes), and other professionals involved in patient care that assist MNG Laboratories in carrying out treatment, payment, and operational activities. Results are kept confidential. Medical Neurogenetics complies with security and privacy statutes of the federal Health Information Portability and Accountability Act (HIPAA). If a patient chooses to specifically declare where results may be released (other than the referring institution and ordering physician), please provide these *in writing* to the Compliance Officer, MNG Laboratories (quickresponse@mnglabs.com).

5. **Consent to Retain Specimen:** The laboratory does not return any remaining sample to individuals or physicians unless requested. No clinical tests other than those authorized shall be performed on the sample. A request for additional testing must be made by my referring physician or other authorized healthcare professional and there will be an additional charge. If agreed by the patient, MNG Laboratories will retain the samples for longer periods for use in an anonymous fashion for research/development or for quality assurance processes.
   I consent to have my specimens retained after completion of initial testing (this consent may be withdrawn at any time and the laboratory will destroy any remaining sample). Patient (or parent/guardian) Initials: _____

6. **Testing for Genetic Conditions can be Complex:** If warranted, obtain professional genetic counseling prior to giving consent to fully understand what the risks and benefits are to having the testing completed. I hereby consent to participate in testing described above. I understand that a biologic specimen will be obtained from me and/or members of my family. I understand that this biologic specimen will be used for the purpose of attempting to determine if I, or members of my family, are affected or are carriers of a particular disease or are at increased risk to someday be affected with this genetic disease.

_____ _____
Signature of Patient Date

_____ _____
Authorized Signature (Parent/Guardian) Relationship

_____ _____
Name of Patient (*please print clearly*) Name of Ordering MD (*please print clearly*)

_____ _____
Referring Facility (*please print clearly*) **Signature of Ordering MD**

Important: One signature from patient (or parent/guardian), authorized person, or physician is required to complete this form. New York requires completion of all signatures on this form.

MNG LABORATORIES
Neurogenetic Answers™

5424 Glenridge Drive NE
Atlanta, GA 30342 USA
toll-free: 844.TESTMNG
fax: 678.225.0212
mnglabs.com

STAT Testing Request Form

We gladly accept deliveries Monday-Saturday, excluding holidays
CLIA License #11D0703390; CAP License #1441004; State of Georgia License #060-381

Patient Name _____ **DOB** _____

STAT Testing - Expedite Your Results

IMPORTANT: To request STAT Testing, STAT Testing Form must be ***completed, signed and submitted*** with test request form. Failure to do so will delay your order.

For a nominal fee, the following tests are available for STAT Testing:

Neurochemistry (NC) & Metabolic (MET) Tests	Molecular (MOL) Tests	Next-Generation Sequencing (NGS) Panels
$100 per test - 7 day TAT	**$200 per test - 2 week TAT**	**$500 per panel - 2 week TAT**

NOTE: All MNG tests rely heavily on our proprietary Genome MaNaGer™ variant calling process coupled with our Neurogenetic Answers™ first-in-class reporting platform that delivers the actionable results you expect. MNG Laboratories will ensure any STAT orders meet the stated deadline, or the STAT fee will be waived.

Test Code

IMPORTANT: Enzymology tests NOT offered as STAT

Test Code: _____	**Test Code:** _____	**Test Code:** _____
Test Code: _____	**Test Code:** _____	**Test Code:** _____
Test Code: _____	**Test Code:** _____	**Test Code:** _____
Test Code: _____	**Test Code:** _____	**Test Code:** _____

Billing Information For STAT Testing [REQUIRED]

Facility Responsible for Payment	Phone
Facility Contact Person	Fax
Facility Contact Person Email	
Facility Billing Address	
City, State, Zip Code	

I HEREBY ACKNOWLEDGE *(check all & sign below)*:

☐ I acknowledge that the responsibile billing party listed above will pay for the additional costs associated with ordering a STAT Test. I understand that failure to submit payment for STAT Testing will delay my order.

☐ I consent that all requested STAT Tests listed above are either Neurochemistry tests, Metabolic tests, Molecular Tests or Next-Generation Sequencing Panels. I understand that all other tests are not available for STAT Testing and will not be ran as a STAT Test if requested.

Signature of Responsible Billing Party (*required*): _____