

For PDF Fillable Requisitions, the following applies:

1. The form shall be completed using a Digital Health assigned computer.
2. Absolutely no personal health information shall be electronically saved on a computer.
3. The completed form shall not be shared electronically. If you reasonably believe that e-mailing the information is the only available method of communication or the only way to send the information then you must adhere to the Privacy guideline titled "E-mailing Personal Health Information".
4. All forms must be completed in their entirety, e.g. if a staff member has only completed half of a form they cannot save their work and then come back to complete it at a later date.
5. Once the personal health information has been recorded onto the form, it is to be printed immediately, deleted (not saved) from the computer, and then stored securely inside the client (paper) health record or scanned into an electronic record.
6. Do not print unnecessary duplicate copies of the form.
7. Regular audits of the Digital Health assigned computer shall be undertaken to ensure that no personal health information is being duplicated and saved.

# Patient and Specimen Information Form

5424 Glenridge Drive NE | Atlanta, GA 30342 USA | phone: 844.664.8378 | fax: 678.225.0212 | mnglabs.com

## 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]	
<b>Gender</b> <input type="checkbox"/> Male <input type="checkbox"/> Female	<b>Specimen Type</b> <input type="checkbox"/> Whole Blood <input type="checkbox"/> Buccal Swab	<input type="checkbox"/> CSF <input type="checkbox"/> Urine <input type="checkbox"/> Fibroblasts	<input type="checkbox"/> Plasma/Serum <input type="checkbox"/> DNA Tissue: _____ <input type="checkbox"/> Muscle

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

## Referring Physician Information

Physician Name		NPI # or equivalent (Required)	
Facility / Organization		Signature	
Facility Address City, State, Zip Code		<input type="checkbox"/> Same as billing	
Report Delivery <input type="checkbox"/> Fax	<input type="checkbox"/> Email	Phone	

## Billing Information (REQUIRED)

Self-Pay? <input type="checkbox"/> Yes    If yes, <b>MUST</b> include payer contact name & details below. Payment must be received in full prior to testing.			
Facility		Contact Name	
Billing Address			
City, State, Zip Code			
Phone	Fax	Email	

## Results

Authorized Recipient Name		Authorized Recipient Name	
Facility	Phone	Facility	Phone
<input type="checkbox"/> Fax		<input type="checkbox"/> Fax	
<input type="checkbox"/> Email		<input type="checkbox"/> Email	

## Testing Checklist

All of the following are encouraged to be included with test orders (please check the following):

- ☐ All specimens that will be analyzed must be received - please note if samples will ship separately
- ☐ Clinical Information Form completed
- ☐ Informed Consent for Genetic Testing completed and signed

# Neurochemistry & Metabolic Test Request Form

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Patient Name \_\_\_\_\_

DOB \_\_\_\_\_

## Metabolic

### CSF

- |   |  |  |
|---|--|--|
| <input type="checkbox"/> (MET01) Amino Acids <sup>†</sup>                                   | <input type="checkbox"/> (NC04) Neurotransmitter Metabolites (5HIAA, HVA, 3OMD) <i>[Includes Biomarkers for Pyridoxine Responsive Seizures]</i>    | <input type="checkbox"/> (NC07) Sialic Acid <i>[Disorders with Hypomyelination of Unknown Etiology/ Sialic Acid Storage Disorders]</i> |
| <input type="checkbox"/> (MET07) Lactate  | <input type="checkbox"/> (NC05) Pyridoxal 5'-phosphate <i>[Pyridox[am]ine Phosphateoxidase Deficiency + CNS Pyridoxal 5'-phosphate Deficiency]</i> | <input type="checkbox"/> (NC08) Alpha-Aminoadipic Semialdehyde <i>[Pyridoxine-Responsive Seizures]</i>                                 |
| <input type="checkbox"/> (MET11) Pyruvate*  | <input type="checkbox"/> (NC06) Succinyladenosine <i>[Adenylosuccinate Lyase Deficiency]</i>   | <input type="checkbox"/> (NC10) Glucose <i>[Glucose Transporter Deficiency]</i>  |
| <input type="checkbox"/> (NC01) 5-Methyltetrahydrofolate                                    |  | <input type="checkbox"/> (NC15) Sepiapterin & Dihydrobiopterin   |
| <input type="checkbox"/> (NC02) Neopterin <i>[Marker for CNS Immune System Stimulation]</i> |  |  |
| <input type="checkbox"/> (NC03) Neopterin/Tetrahydrobiopterin                               |  |  |

### Blood & Muscle

- |  |   |  |
|--|---|--|
| <input type="checkbox"/> (MET02) Amino Acids (Plasma) <sup>†</sup> | <input type="checkbox"/> (MET08) Lactate (Plasma)                         | <input type="checkbox"/> (MET23) Creatine & Guanidinoacetate (Plasma)  |
| <input type="checkbox"/> (MET04) Coenzyme Q10 Level (Leukocytes)   | <input type="checkbox"/> (MET10) Pyruvate* (Blood)                        | <input type="checkbox"/> (MET24) Glucose (Plasma)  |
| <input type="checkbox"/> (MET05) Coenzyme Q10 Level (Muscle)       | <input type="checkbox"/> (MET12) Thymidine/Deoxyuridine Analytes (Plasma) | <input type="checkbox"/> (MET29) 3-O-Methyldopa (Plasma) <i>[Specific Marker for Aromatic L-Amino Acid Decarboxylase Deficiency]</i> |

### Urine

- |   |  |   |
|---|--|---|
| <input type="checkbox"/> (MET03) Amino Acids <sup>†</sup> | <input type="checkbox"/> (MET19) Creatine & Guanidinoacetate | <input type="checkbox"/> (MET20) Alpha Amino adipic Semialdehyde <i>[Urine; for Pyridoxine-Responsive Seizures]</i> |
|---|--|---|

## Enzymology

### Blood

- |   |  |
|---|--|
| <input type="checkbox"/> (ENZ01) Aromatic L-amino Acid Decarboxylase Enzyme Analysis (Plasma) - <b>STAT Not Available</b> | <input type="checkbox"/> (ENZ06) Thymidine Phosphorylase Enzyme Analysis (Blood) - <b>STAT Not Available</b> |
|---|--|

<sup>†</sup> Denotes testing performed at LabCorp, Burlington, NC - **STAT Not Available**

\*Denotes testing requires deproteinization



**MNG LABORATORIES**

A LabCorp Company

# Clinical Information Form

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Patient Name \_\_\_\_\_ DOB \_\_\_\_\_

## Clinical (Check All That Apply)

<b>Eye</b> <input type="checkbox"/> Retinitis Pigmentosa <input type="checkbox"/> Optic Atrophy <input type="checkbox"/> Other	<b>Hearing</b> <input type="checkbox"/> Sensorineural <input type="checkbox"/> Stickler <input type="checkbox"/> Usher	<b>Neuronal Migration</b> <input type="checkbox"/> Meckel <input type="checkbox"/> Joubert <input type="checkbox"/> Other	<input type="checkbox"/> Stroke
<b>Cognitive/Neurobehavioral</b> <input type="checkbox"/> Intellectual Disability (ID) <input type="checkbox"/> Syndromic ID <input type="checkbox"/> Nonsyndromic ID <input type="checkbox"/> Autism <input type="checkbox"/> Dementia			
<b>Movement Disorders</b> <input type="checkbox"/> Ataxia <input type="checkbox"/> Episodic Ataxia <input type="checkbox"/> Dystonia <input type="checkbox"/> Chorea/Athetosis <input type="checkbox"/> Parkinson Disease <input type="checkbox"/> L-Dopa Response			
<b>Epilepsy</b> <input type="checkbox"/> Myoclonic <input type="checkbox"/> Other <input type="checkbox"/> Absence <input type="checkbox"/> Tonic Clonic <input type="checkbox"/> Epileptic Encephalopathy	<b>Spasticity</b> <input type="checkbox"/> Spastic Paraplegia <input type="checkbox"/> Other <input type="checkbox"/> Spastic Quadriplegia	<b>Connective Tissue &amp; Bone</b> <input type="checkbox"/> Ehlers Danlos <input type="checkbox"/> Marfan <input type="checkbox"/> Aneurysms <input type="checkbox"/> Other	
<b>Neuromuscular</b> <input type="checkbox"/> Distal <input type="checkbox"/> Proximal <input type="checkbox"/> Muscle Atrophy <input type="checkbox"/> Contractures <input type="checkbox"/> Malignant Hyperthermia <input type="checkbox"/> Arthrogryposis <input type="checkbox"/> Rhabdomyolysis <input type="checkbox"/> Periodic Paralysis <input type="checkbox"/> Statin Use <input type="checkbox"/> Myasthenia		<b>Nerve/Anterior Horn Cell</b> <input type="checkbox"/> Neurofibromas <input type="checkbox"/> Charcot-Marie-Tooth <input type="checkbox"/> Sensory <input type="checkbox"/> Autonomic <input type="checkbox"/> Pain <input type="checkbox"/> Motor <input type="checkbox"/> Nerve Conduction <input type="checkbox"/> Other	
<b>Cardiomyopathy</b> <input type="checkbox"/> Dilated <input type="checkbox"/> Hypertrophic <input type="checkbox"/> Noncompaction	<b>Arrhythmias</b> <input type="checkbox"/> Ventricular Tachycardia <input type="checkbox"/> Brugada <input type="checkbox"/> Long or Short QT <input type="checkbox"/> Conduction Defect	<b>Congenital Heart Defects</b> <input type="checkbox"/> Heterotaxy <input type="checkbox"/> Other	<b>Endocrine</b> <input type="checkbox"/> Hypothyroidism <input type="checkbox"/> Other <input type="checkbox"/> Diabetes Mellitus

## Imaging (Check All That Apply)

<b>Brain MRI</b> <input type="checkbox"/> Leigh Disease <input type="checkbox"/> Basal Ganglia Calcification <input type="checkbox"/> Stroke <input type="checkbox"/> Cerebellar Atrophy <input type="checkbox"/> Abnormal Myelin (describe)	<b>EEG (Describe Findings)</b>	<b>EMG/NVC (Describe Findings)</b>
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## Laboratory

<b>Metabolic (Describe Findings)</b>	<b>Genetic (Describe Findings)</b> <input type="checkbox"/> Chromosomal Microarray <input type="checkbox"/> Deletion/Insertion Testing <input type="checkbox"/> Other (comment)
<b>CPK</b> Maximum _____ Minimum _____	

## Family History

<b>Ethnicity (please check)</b> <input type="checkbox"/> Caucasian <input type="checkbox"/> Sephardic Jewish <input type="checkbox"/> African American (or Black) <input type="checkbox"/> Asian <input type="checkbox"/> Hispanic <input type="checkbox"/> Ashkenazi Jewish <input type="checkbox"/> Native American (or American Indian) <input type="checkbox"/> Other: _____		
<b>Affected Maternal Lineage</b> Relationship to Proband  Symptoms	<b>Affected Paternal Lineage</b> Relationship to Proband  Symptoms	<b>Siblings</b> Number (specify gender)  Healthy/Affected

## Additional Comments

# STAT Test Request Form

5424 Glenridge Drive NE | Atlanta, GA 30342 USA | phone: 844.664.8378 | fax: 678.225.0212 | mnglabs.com

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 an additional fee, the following tests are available for STAT Testing:

<b>Neurochemistry (NC) &amp; Metabolic (MET) Tests</b> 7 day TAT	<b>Molecular (MOL) Tests</b> 2 week TAT	<b>Next-Generation Sequencing (NGS) Panels</b> 2 week TAT
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*NOTE: MNG Laboratories will ensure any STAT orders meet the stated deadline, or the STAT fee will be waived.*

## Patient and Specimen Information

Patient Last Name	Patient First Name
Patient ID #	Date of Birth [MM/DD/YYYY]

## Test Code

*IMPORTANT: Enzymology, familial variants, and RNA tests NOT available 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 (**REQUIRED**)

Self-Pay? <input type="checkbox"/> Yes      If yes, <b>MUST</b> include payer contact name & details below. Payment must be received in full prior to testing.		
Facility	Contact Name	
Billing Address		
City, State, Zip Code		
Phone	Fax	Email

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

- ☐ I acknowledge that the responsible 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):** \_\_\_\_\_