



Children's Hospital Colorado

Children's Hospital Colorado
Department of Pathology & Laboratory Medicine
Biochemical Genetics & Mitochondrial Lab Requisition
Phone (720) 777-6711
Fax (720) 777-7118

Specimen Shipping Address:
 Children's Hospital Colorado
 Clinical Laboratory - Room B0200
 13123 E. 16th Ave
 Aurora, CO 80045

FAILURE TO COMPLETE BELOW FIELDS WILL DELAY RESULTS

*****PLEASE PROVIDE COMPLETE BILLING INFORMATION ON THE BACK OF THIS FORM*****

Contact Information

Ordering Institution Name		Ordering Institution Address	
		Street _____	
		City, State, Zip _____	
Ordering Provider (Last, First, and Middle Initial)		Ordering Provider Phone _____	
Result Contact Name	Result Phone	Result Fax	

Patient Information

Last Name	First Name	Middle I	Birthdate (MM/DD/YYYY)	Sex
Client Medical Record Number	Client Specimen Number		Diagnosis/ICD-10 Code	

Specimen Information

Date Collected (MM/DD/YY)	Time Collected (HHMM)	AM / PM
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Biochemical Genetics Test Information

Amino Acids	Other
<input type="checkbox"/> Amino Acids [quantitative] Please select source: <input type="checkbox"/> Serum/Plasma (L6606) <input type="checkbox"/> Urine* (L6574) <input type="checkbox"/> CSF (L6608)	<input type="checkbox"/> Acylcarnitines, serum/plasma [qualitative] (L6615) <input type="checkbox"/> D- & L- isomers of 2-hydroxyglutaric acid, Urine* [qualitative] (L7084) <input type="checkbox"/> Ethylmalonic acid, Urine* [quantitative] (L6759) <input type="checkbox"/> Expanded Newborn Screen, Dried Blood Spot [qualitative] (L6996) <input type="checkbox"/> Glutaric & 3-hydroxyglutaric acids [quantitative]
<input type="checkbox"/> Amino Acid screen [qualitative] Please select source: <input type="checkbox"/> Serum/Plasma (L6751) <input type="checkbox"/> Urine* (L6753)	Please select source: <input type="checkbox"/> Serum/Plasma (L6742) <input type="checkbox"/> Urine* (L6743)
<input type="checkbox"/> Alanine, Serum (L6631)	<input type="checkbox"/> Homovanillic acid [HVA] & Vanillylmandelic acid [VMA], Urine* [quantitative] (L7177)
<input type="checkbox"/> Aspartylglucosamine, Urine* [qualitative] (L6894)	<input type="checkbox"/> 3-Methylglutaconic acid, Urine* [quantitative] (L6860)
<input type="checkbox"/> Benzoic acid, Serum/Plasma [quantitative] (L6760)	<input type="checkbox"/> Methylmalonic acid [quantitative]
<input type="checkbox"/> Branch-chain amino acids (MPS), Serum/Plasma [quantitative] (L6607)	Please select source: <input type="checkbox"/> Serum/Plasma (L6750) <input type="checkbox"/> Urine* (L6762)
<input type="checkbox"/> Cystine, Serum (L6765)	<input type="checkbox"/> Mucopolysaccharide screen, Urine* [qualitative] (L6771)
<input type="checkbox"/> Glycine, Serum (L6630)	<input type="checkbox"/> Organic acid, Urine* [qualitative] (L6619)
<input type="checkbox"/> Glycine, CSF (L6629)	<input type="checkbox"/> Orotic acid, Urine* [quantitative] (L6754)
<input type="checkbox"/> Homocystine, Urine* (L6954)	<input type="checkbox"/> Phytanic acid, Serum/Plasma [quantitative] (L6761)
<input type="checkbox"/> Methionine, Serum (L6767)	<input type="checkbox"/> Succinylacetone, Urine [quantitative] (L6752)
<input type="checkbox"/> Phosphoethanolamine, Urine* [quantitative] (L6893)	<input type="checkbox"/> Succinylpurine [Bratton-Marshall Test] [qualitative]
<input type="checkbox"/> S-sulfo-cysteine, Urine* [quantitative] (L6863)	Please select source: <input type="checkbox"/> Urine (L6862) <input type="checkbox"/> CSF (L6861)
Enzyme Activity Assays	<input type="checkbox"/> Trimethylamine [TMA] & TMA n-oxide [TMAO], Urine* [quantitative] (L6949)
<input type="checkbox"/> GAI; glutaryl-CoA dehydrogenase activity in fibroblasts (L7077)**	<input type="checkbox"/> Pre-choline load collection date/time _____
<input type="checkbox"/> GAI; ETF and ETF:QO activity in fibroblasts (L7089)**	<input type="checkbox"/> Post-choline load collection date/time _____
<input type="checkbox"/> VLCAD; very long chain acyl-CoA dehydrogenase activity in blood (L6895) ¹	

*Due to specimen source, this test includes creatinine **Mycoplasma Tested Y N

¹ For VLCAD genetic sequencing please use our molecular genetics requisition

Mitochondrial Diagnostic Test Information

*****Please complete the clinical information portion on the back of this requisition*****

<input type="checkbox"/> Blue native electrophoresis with in-gel activity staining Please select source: <input type="checkbox"/> Muscle/Liver/Heart (L7086) Weight: _____ mg <input type="checkbox"/> Fibroblasts (L7078) Mycoplasma tested Y <input type="checkbox"/> N <input type="checkbox"/>	<input type="checkbox"/> Mitochondrial respiratory chain enzyme assay Please select source: <input type="checkbox"/> Muscle/Liver/Heart (L7086) Weight: _____ mg <input type="checkbox"/> Fibroblasts (L7079) Mycoplasma tested Y <input type="checkbox"/> N <input type="checkbox"/>
<input type="checkbox"/> Glycine cleavage enzyme assay, Liver	<input type="checkbox"/> Pyruvate dehydrogenase enzyme assay, Fibroblasts (L7085)
Tissue Source: <input type="checkbox"/> Biopsy <input type="checkbox"/> Autopsy (collected _____ Hrs after death)	Specimen Storage Prior to Shipment: <input type="checkbox"/> Liquid Nitrogen <input type="checkbox"/> -20°C <input type="checkbox"/> -70°C
Use of Antibiotics linezolid, macrolides, chloramphenicol in last week? <input type="checkbox"/> Y <input type="checkbox"/> N	

Additional Comments:

Specimen requirements and shipping and handling information can be found on our website at
www.childrenscolorado.org/lab By submitting this document you agree to the terms and conditions listed on our website



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Please do not send patient insurance. We bill clients only, referring provider will be held responsible for payment if no billing information is provided.

FAILURE TO COMPLETE WILL DELAY RESULTS

Bill To: **Billing Facility and Address same as listed on page 1**

Institution Name: _____

Address: _____

City, State, Zip: _____

Phone: _____

Fax: _____

Billing Contact Information:

Name: _____

Email: _____

Phone: _____

Clinical Information (for Mitochondrial testing ONLY)

Please check all that apply

- | | | | |
|---|--|---|--|
| <p>General</p> <p><input type="checkbox"/> failure to thrive</p> <p><input type="checkbox"/> short stature</p> <p>Brain</p> <p><input type="checkbox"/> microcephaly</p> <p><input type="checkbox"/> encephalophaly</p> <p><input type="checkbox"/> seizures:</p> <p><input type="checkbox"/> myclonic seizures</p> <p><input type="checkbox"/> infantile</p> <p><input type="checkbox"/> other seizures</p> <p><input type="checkbox"/> chorea</p> <p><input type="checkbox"/> dystonia</p> <p><input type="checkbox"/> parkinson</p> <p><input type="checkbox"/> ataxia</p> <p><input type="checkbox"/> neurodegeneration</p> <p><input type="checkbox"/> stroke-like episodes</p> <p><input type="checkbox"/> central apnea</p> <p><input type="checkbox"/> leukodystrophy</p> <p><input type="checkbox"/> other: _____</p> <p>Eye</p> <p><input type="checkbox"/> optic atrophy</p> <p><input type="checkbox"/> retinitis pigmentosa/retinal dystrophy</p> <p><input type="checkbox"/> cataracts</p> <p><input type="checkbox"/> ophthalmoplegia</p> <p>Hearing</p> <p><input type="checkbox"/> nerve deafness</p> <p><input type="checkbox"/> hearing loss</p> <p><input type="checkbox"/> other</p> | <p>Endocrine</p> <p><input type="checkbox"/> diabetes mellitus</p> <p><input type="checkbox"/> hypoparathyroidism</p> <p>Kidney</p> <p><input type="checkbox"/> renal Fanconi</p> <p><input type="checkbox"/> glomerulosclerosis</p> <p><input type="checkbox"/> proteinuria – nephritic syndrome</p> <p>Liver</p> <p><input type="checkbox"/> elevated transaminases</p> <p><input type="checkbox"/> fibrosis/steatosis</p> <p><input type="checkbox"/> liver insufficiency</p> <p><input type="checkbox"/> hypoglycemia</p> <p>GI</p> <p><input type="checkbox"/> pancreatitis</p> <p><input type="checkbox"/> pancreatic insufficiency</p> <p><input type="checkbox"/> pseudo-obstruction</p> <p><input type="checkbox"/> malabsorption</p> <p>Heart</p> <p><input type="checkbox"/> Cardiomyopathy:</p> <p><input type="checkbox"/> dilated</p> <p><input type="checkbox"/> hypertrophic</p> <p><input type="checkbox"/> Conduction defect</p> <p><input type="checkbox"/> Arrhythmias</p> <p>Skin</p> <p><input type="checkbox"/> hyperpigmentation</p> | <p>Muscle</p> <p><input type="checkbox"/> myopathy</p> <p><input type="checkbox"/> early fatigue</p> <p><input type="checkbox"/> abnormal exercise test</p> <p><input type="checkbox"/> rhabdomyolysis</p> <p><input type="checkbox"/> elevated CK: _____ U/L</p> <p><input type="checkbox"/> Biopsy:</p> <p><input type="checkbox"/> ragged red fibers</p> <p><input type="checkbox"/> abnormal e.m.</p> <p>Laboratory Studies</p> <p><input type="checkbox"/> Lactate:</p> <p><input type="checkbox"/> Blood: _____ mM</p> <p><input type="checkbox"/> CSF: _____ mM</p> <p><input type="checkbox"/> lactate/pyruvate ratio: _____</p> <p><input type="checkbox"/> 3OHB/AcAc ratio: _____ O</p> <p><input type="checkbox"/> 3-methylglutaconic acid</p> <p><input type="checkbox"/> low total carnitine</p> <p><input type="checkbox"/> lactate on MRS</p> <p><input type="checkbox"/> elevated alanine: _____ μM</p> <p><input type="checkbox"/> incr. Krebs cycle metabolites</p> <p><input type="checkbox"/> other: _____</p> <p>Radiology</p> <p><input type="checkbox"/> abnormal basal ganglia</p> <p><input type="checkbox"/> abnormal brain stem</p> <p><input type="checkbox"/> abnormal dentate nucleus</p> <p><input type="checkbox"/> brain atrophy</p> <p><input type="checkbox"/> cerebellar atrophy</p> <p><input type="checkbox"/> leukodystrophy</p> <p><input type="checkbox"/> other: _____</p> | <p>Recognized Syndromes</p> <p><input type="checkbox"/> Leigh disease</p> <p><input type="checkbox"/> Kearns-Sayre</p> <p><input type="checkbox"/> MELAS</p> <p><input type="checkbox"/> MERRF</p> <p><input type="checkbox"/> NARP</p> <p><input type="checkbox"/> MNGIE</p> <p><input type="checkbox"/> Diabetes-deafness</p> <p><input type="checkbox"/> Diabetes- retinitis pigmentosa</p> <p><input type="checkbox"/> CPEO</p> |
|---|--|---|--|

Previous Mito Investigations: