



NATIONWIDE CHILDREN'S

Laboratory Services

Biochemical Genetics Laboratory
700 Children's Drive, Columbus, Ohio 43205
P: (614) 722-5477 / (800) 934-6575
F: (614) 722-5478 / (877) 722-5478
NationwideChildrens.org/Lab

Ship samples to the CPA Lab , Room C1955

Practice/ Office Name: _____

Address: _____

Address: _____

City, State, Zip: _____

Phone: _____

Fax: _____

Please Mark Billing Option:

Patient Bill: _____ / Client Bill _____

BIOCHEMICAL GENETICS

Patient Information	Parent/Guardian / Billing Information
Legal Last Name:	Guardian Legal Last Name:
First Name: MI:	Guardian First Name: MI:
MRN/ Patient ID:	Patient Relationship:
DOB:	Guardian Contact Phone #: ()
Address:	Subscriber Legal Last Name:
City, State, Zip:	Subscriber First Name and MI:
Phone #: ()	Subscriber DOB: Sex: [] Male [] Female
Race:	Subscriber Social Security #:
Ethnicity:	Subscriber Phone #: ()
Sex: [] Male [] Female	Subscriber Address (if different from patient):
Specimen Information	
Collection Date: Time: [] AM [] PM	Insurance Co. Name:
Collected By (full name):	Policy #: Group #:
Storage Temp: [] Refrigerated [] Frozen [] Room Temperature	Insurance Address:
Patient Fasting?: [] Yes [] No * If yes, how many hours? []	Secondary Insurance Co. Name:

Biochemical Genetic Tests to be Performed:

- ☐ CAH6 Profile (CAH6) includes Cortisol, in addition to the tests below. Any of the tests below can be requested individually.
- ☐ 11-Desoxycortisol (11DESC)
- ☐ 17-OH-Pregnenolone (17PRE)
- ☐ 17-OH-Progesterone (17OP)
- ☐ Androstenedione (ANDRO)
- ☐ Dehydroepiandrosterone (DHEA)
- ☐ Deoxycorticosterone (DOC)
- ☐ Progesterone (PROG)
- ☐ Testosterone, Total (TESTMS)

Lab CPA -
Do not order
these tests if
CAH6 Profile
has been
selected.

Additional Tests:

- ☐ Homovanillic Acid / Vanillylmandelic Acid - HVA (VMHVP)
- ☐ Methylmalonic Acid (MMA)
- ☐ Phenylalanine / Tyrosine (PATY)
- ☐ Psychosine (PSYC)

Additional Biochemical Genetic Tests to be Performed:

****NOTE: If any of the tests below are selected, the Clinical Data Form on page 2 MUST be filled out completely.****

- ☐ CSF Amino Acids by LC-MS/MS (MSAAC) *NOTE: an additional sample - 3 mL heparin green top **NO** gel is required
- ☐ Plasma Amino Acids by LC-MS/MS (MSAAP)
- ☐ Urine Amino Acids by LC-MS/MS (MSAAUP)
- ☐ Organic Acids, Urine (ORGUP)
- ☐ Acylcarnitines, Quantitative, Plasma (preferred) or Serum (SPACP)
- ☐ Acylcarnitines, Quantitative, Blood Spot Filter Paper (FPACP)
- ☐ Carnitine Total & Free, Plasma (preferred) or Serum (FCTC)

Diagnosis/ ICD 10 _____ Physician (Print full name) _____ Physician's Signature (Required) _____

Date _____ Time _____ [] AM [] PM



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Patients Last Name: _____

Patients First Name: _____

Date of Birth: _____

BIOCHEMICAL GENETICS CLINICAL DATA FORM

CNS Symptoms: (check all that apply)		GI Symptoms: (check all that apply)		Physical Features: (check all that apply)	
<input type="checkbox"/> Decreased mental status	<input type="checkbox"/> Ataxia	<input type="checkbox"/> Diarrhea	<input type="checkbox"/> Splenomegaly	<input type="checkbox"/> Alopecia	<input type="checkbox"/> Dysmorphic features
<input type="checkbox"/> Elevated lactate	<input type="checkbox"/> Coma	<input type="checkbox"/> Hepatomegaly	<input type="checkbox"/> Vomiting	<input type="checkbox"/> Skin rash	<input type="checkbox"/> Unusual odor (sweat or urine)
<input type="checkbox"/> Psychomotor retardation	<input type="checkbox"/> Seizures	<input type="checkbox"/> Other: _____		<input type="checkbox"/> Other: _____	

Abnormal Lab Findings (check all that apply):		
<input type="checkbox"/> Failed State Newborn Screen – **Please send copy of state newborn screen results with sample**		
<input type="checkbox"/> Elevated ammonia	<input type="checkbox"/> Elevated lactate	Other Abnormal Tests: _____
<input type="checkbox"/> Hypoglycemia	<input type="checkbox"/> Ketosis	_____

Chief Reason for Requesting Test(s):

Diet for the past seven days prior to sampling (specify formula type or solid food):
Day 1:
Day 2:
Day 3:
Day 4:
Day 5:
Day 6:
Day 7:

Current Medications (list): **This information is critical to accurate analytical interpretation**

Comments and other relevant or unique patient findings: