

BIOCHEMICAL TEST REQUISITION

SHIP TO: MEDICAL GENETICS LABORATORIES

Baylor College of Medicine
2450 Holcombe, Grand Blvd. - Receiving Dock
Houston, TX 77021-2024

!STOP IMPORTANT BILLING INFORMATION!

Indicate Institution Billing Code here: _____, **OR**
complete and fax billing information form (page 2 of requisition)
to 713-798-4187. **Samples received without the Institution
Code or Billing Form cannot be processed.**

FOR PRENATALS, PLEASE USE PRENATAL REQUISITION FORM.

DATE SAMPLE OBTAINED (MM/DD/YY): ____/____/____

PATIENT DATA

NAME (Last, First, Middle Initial): _____

DATE OF BIRTH (MM/DD/YY): ____/____/____

Please check one: MALE FEMALE UNKNOWN

HOSPITAL #: _____

ACCESSION #: _____

REPORTING INFORMATION

PHYSICIAN/INSTITUTION: _____

ADDRESS: _____

CITY, STATE, ZIP: _____

PHONE #: (____) _____ FAX #: (____) _____

Additional Reports to:

1. NAME: _____ FAX #: (____) _____

2. NAME: _____ FAX #: (____) _____

ANALYTE TEST REQUESTED

- | | | |
|--|---|---|
| 4100 <input type="checkbox"/> Amino Acids Analysis - Plasma | 4260 <input type="checkbox"/> Creatine/Guanidinoacetate Analysis - Urine | 4110 <input type="checkbox"/> Phenylalanine/Tyrosine - Plasma |
| 4160 <input type="checkbox"/> Amino Acids Analysis - CSF | 4627 <input type="checkbox"/> Cystine, White Blood Cell - Heparinized Blood | 4340 <input type="checkbox"/> Polyols, Urinary |
| 4240 <input type="checkbox"/> Amino Acids Analysis - Urine | 4140 <input type="checkbox"/> Homocysteine, Total Plasma | 4220 <input type="checkbox"/> Purine Panel - Urine |
| 4300 <input type="checkbox"/> Acylcarnitine/Carnitine Combination - Plasma | 4150 <input type="checkbox"/> Methylmalonic Acid - Plasma | 4250 <input type="checkbox"/> Succinylacetone - Urine |
| 4310 <input type="checkbox"/> Carnitine, Free and Total - Plasma | 4200 <input type="checkbox"/> Organic Acid Screen - Urine | 4330 <input type="checkbox"/> Thymidine - Plasma |
| 4130 <input type="checkbox"/> Creatine/Guanidinoacetate Analysis - Plasma | 4210 <input type="checkbox"/> Orotic/Orotidine Analysis - Urine | |

ENZYME TEST REQUESTED

DISEASE NAME	SAMPLE TYPE				
	RBC	Serum	SFC	WBC	Other
<input type="checkbox"/> Adenosine Deaminase Deficiency	<input type="radio"/> 4509		<input type="checkbox"/> 4510	<input type="checkbox"/> 4511	
<input type="checkbox"/> Argininemia	<input type="radio"/> 4536				<input type="checkbox"/> 4535 (L)
<input type="checkbox"/> Argininosuccinate Lyase Deficiency	<input type="radio"/> 4524		<input type="checkbox"/> 4525		<input type="checkbox"/> 4523 (L)
<input type="checkbox"/> Aspartylglucosaminuria			<input type="checkbox"/> 4514		
<input type="checkbox"/> Biotinidase		<input type="radio"/> 4555			
<input type="checkbox"/> Carbamyl Phosphate Synthetase Deficiency					<input type="radio"/> 4561 (L)
<input type="checkbox"/> Citrullinemia			<input type="radio"/> 4545		<input type="checkbox"/> 4544 (L)
<input type="checkbox"/> Fabry Disease			<input type="checkbox"/> 4516	<input type="radio"/> 4517	
<input type="checkbox"/> Fucosidosis			<input type="checkbox"/> 4512	<input type="radio"/> 4513	
<input type="checkbox"/> Galactosemia (UDPG Transferase)	<input type="radio"/> 4619				
<input type="checkbox"/> Gaucher Disease			<input type="checkbox"/> 4553	<input type="radio"/> 4554	
<input type="checkbox"/> Glucuronidase Deficiency (Sly, MPSVII)		<input type="radio"/> 4550	<input type="checkbox"/> 4551		
<input type="checkbox"/> GM1 Gangliosidosis (Morquio B, MPS IV B)			<input type="checkbox"/> 4548	<input type="radio"/> 4549	
<input type="checkbox"/> Hurler Disease (Hurler-Scheie, MPS I)			<input type="checkbox"/> 4575	<input type="radio"/> 4576	
<input type="checkbox"/> Krabbe Disease			<input type="checkbox"/> 4565	<input type="radio"/> 4566	
<input type="checkbox"/> Lesch-Nyhan Disease	<input type="radio"/> 4572		<input type="checkbox"/> 4573		
<input type="checkbox"/> Lowe Syndrome			<input type="radio"/> 4585		
<input type="checkbox"/> Mannosidosis		<input type="checkbox"/> 4526	<input type="checkbox"/> 4527	<input type="radio"/> 4528	
<input type="checkbox"/> Maroteaux-Lamy (MPS VI)			<input type="checkbox"/> 4540		
<input type="checkbox"/> Metachromatic Leukodystrophy			<input type="checkbox"/> 4537	<input type="radio"/> 4538	
<input type="checkbox"/> Niemann-Pick Disease			<input type="checkbox"/> 4607	<input type="radio"/> 4608	
<input type="checkbox"/> Ornithine Transcarbamylase Deficiency					<input type="radio"/> 4582 (L)
<input type="checkbox"/> Purine Nucleoside Phosphorylase Deficiency	<input type="radio"/> 4592		<input type="checkbox"/> 4593	<input type="checkbox"/> 4594	
<input type="checkbox"/> Pompe Disease			<input type="radio"/> 4520		<input type="radio"/> 4519 (M)
<input type="checkbox"/> Sanfilippo B Mucopolysaccharidosis (MPS III)		<input type="radio"/> 4579	<input type="checkbox"/> 4580		
<input type="checkbox"/> Sialidosis (Neuraminidase Deficiency)			<input type="radio"/> 4603		
<input type="checkbox"/> Tay-Sachs Disease		<input type="radio"/> 4569			
<input type="checkbox"/> Tay-Sachs Disease Carrier Testing		<input type="radio"/> 4617			
<input type="checkbox"/> Wolman Disease (Cholesterol Ester Storage Disease)			<input type="checkbox"/> 4503	<input type="radio"/> 4504	<input type="checkbox"/> 4502 (L)
<input type="checkbox"/> X-Linked Ichthyosis (Steroid Sulfatase Deficiency)			<input type="checkbox"/> 4614	<input type="radio"/> 4615	

* - Preferred specimen type.

Sample types are: Serum; RBC - Red Blood Cells; WBC - White Blood Cells; SFC - Skin Fibroblast Culture; M - Muscle; L - Liver.

PLEASE CONSULT THE LABORATORY PRIOR TO SUBMITTING SKIN FIBROBLAST CULTURES.

BILLING INFORMATION FORM

STOP! ONE OF THE THREE FOLLOWING BILLING OPTIONS MUST BE INDICATED BELOW.
The Self-Pay option must include payment with sample. We require and provide insurance pre-verification service. Please fax the *Patient Insurance Verification Form* (available at www.bcmgeneticlabs.org) to 713-798-4187. If the Billing Information section is incomplete, the referring physician, hospital, or laboratory will automatically be billed, or sample processing suspended. Please forward billing questions to: medgenbilling@bcm.edu

PATIENT INFORMATION:

Name (Last, First, Middle Initial): _____
Address: _____
City, State, Zip: _____
Phone #: (____) _____ Email: _____

PAYMENT OPTIONS:

1. **Institution or referring MD Code (as assigned by BCM):** _____
(or) Institution Name: _____
Billing Address: _____
City, State, Zip: _____
Financial Contact: _____
E-mail (required): _____
Phone #: (____) _____ Fax #: (____) _____

2. **Self-Pay: Check, Money Order, or Credit Card payment must accompany sample.**
Credit Card (Please check one): AMEX Discover MC VISA
Valid Card #: _____ Exp date (mm/yy): ____/____ **CVC Code:** _____
Cardholder printed name: _____
Cardholder signature: _____

3. **Insurance:** Please refer to the Financial Policy at <http://www.bcm.edu/geneticlabs/billing.html> for complete insurance filing information and managed care contract list. Insurance is filed to our contracted carriers as a courtesy. Patients are responsible for non-covered services, deductibles, co-insurance, contract exclusions, non-authorized services, and remaining balances after insurance reimbursement. HMO policies must have required authorizations. We do not file out-of-state Medicaid. Prenatal CMA requires a prepayment amount. Contact medgenbilling@bcm.edu with questions.

ICD9 Diagnosis Code(s) must be provided or insurance cannot be filed: **ICD-9 CODE:** _____
 PPO, Commercial Insurance-provide Patient Insurance Verification form (PIVF) and front/back copy of card
 HMO-provide PIVF, authorization, front/back copy of insurance card
 Texas Medicaid/Texas Medicaid HMO-provide PIVF, authorization, front/back copy of Medicaid card

Insured Policyholder's Information:

Name: _____ Date Of Birth (mm/dd/yy): ____/____/____
Insured SS or ID #: _____ Gender (Please check one): M F
Authorization: _____
Relationship to Patient: _____
Insurance Name: _____
Employer: _____ Group #: _____
Insurance Address: _____
Insurance City, State, Zip: _____
Insurance Phone #: (____) _____

I authorize BCM Medical Genetics Laboratories to furnish any medical information requested on myself, or my covered dependents. In consideration of services rendered, I transfer and assign any benefits of insurance to BCM Medical Genetics Laboratories. I understand I am responsible for any co-pay, deductible, or non-covered service amounts. I understand I am fully responsible for payment of my account if the BCM Medical Genetics Laboratories is not a participant with my health plan, and my health plan does not fully reimburse my medical services due to lack of authorization or medical necessity.

Printed Name: _____

Signature: _____ Date (mm/dd/yy): ____/____/____