

BAYLOR MIRACA GENETICS LABORATORIES

PHONE: 800-411-GENE | FAX: 713-798-2787 | www.bmgl.com

SHIP TO: Baylor Miraca Genetics Laboratories
2450 Holcombe, Grand Blvd. -Receiving Dock
Houston, TX 77021-2024
Phone: 713-798-6555

BIOCHEMICAL TESTING REQUISITION - INSTITUTIONAL ONLY

PATIENT INFORMATION	REPORTING INFORMATION
NAME: _____ <small>LAST NAME FIRST NAME MI</small>	PHYSICIAN: _____
ADDRESS: _____ PHONE #: _____	INSTITUTION: _____
DATE OF BIRTH: ____ / ____ / ____ GENDER (Please select one): <input type="checkbox"/> FEMALE <small>MM DD YY</small> <input type="checkbox"/> MALE <input type="checkbox"/> UNKNOWN	PHONE: _____ FAX: _____
HOSPITAL #: _____ ACCESSION #: _____	EMAIL (INTERNATIONAL CLIENT REQUIREMENT): _____

-OR-
PLACE PATIENT STICKER HERE

ADDITIONAL PROFESSIONAL REPORT RECIPIENTS

NAME: _____
PHONE: _____ FAX: _____
NAME: _____
PHONE: _____ FAX: _____

REQUIRED: NEW YORK STATE PHYSICIAN SIGNATURE OF CONSENT

I certify that the patient specified above and/or their legal guardian has been informed of the benefits, risks, and limitations of the laboratory test(s) requested. I have answered this person's questions. I have obtained informed consent from the patient or their legal guardian for this testing.

Physician's Printed Name: _____

Signature: _____

Date (MM/DD/YY): _____

BILLING INFORMATION

INSTITUTIONAL BILLING CODE: _____

SAMPLE INFORMATION

CHECK ALL THAT APPLY

SAMPLE TYPE:	DATE OF COLLECTION (MM/DD/YY):	SAMPLE TYPE:	DATE OF COLLECTION (MM/DD/YY):
<input type="checkbox"/> PLASMA	____ / ____ / ____	<input type="checkbox"/> WHOLE BLOOD (SODIUM HEPARIN - GREEN TOP)	____ / ____ / ____
<input type="checkbox"/> URINE	____ / ____ / ____	<input type="checkbox"/> WHOLE BLOOD (ACD - YELLOW TOP)	____ / ____ / ____
<input type="checkbox"/> CEREBRAL SPINAL FLUID	____ / ____ / ____	<input type="checkbox"/> LIVER	____ / ____ / ____
<input type="checkbox"/> SKIN FIBROBLAST CULTURE	____ / ____ / ____	<input type="checkbox"/> OTHER (please specify): _____	____ / ____ / ____
<input type="checkbox"/> SERUM	____ / ____ / ____		

INDICATIONS ARE REQUIRED. PLEASE FILL OUT CHECKLIST ON PAGE 4

SEE PAGES 2 & 3 FOR TESTING OFFERED

BIOCHEMICAL TESTING REQUISITION - INSTITUTIONAL ONLY

NAME: _____ DATE OF BIRTH: ____ / ____ / ____ GENDER (Please select one): FEMALE
LAST NAME FIRST NAME MI MM DD YY MALE UNKNOWN

SEVERE COMBINED IMMUNODEFICIENCY (SCID) NEWBORN SCREENING FOLLOW-UP PANEL

4001 This panel includes the following tests:
 Adenosine Deaminase Deficiency Enzyme (4509)
 Purine Nucleoside Phosphorylase Deficiency Enzyme (4592)
 Urine Purine Panel (4220)
 CMA-HR (180K High Resolution Copy Number ≥30Kb) (8655)

SPECIMEN REQUIREMENTS:
 Sample types below must be shipped overnight:
 BLOOD: All Blood samples must be shipped at ambient temperature.
 2-3 cc in EDTA (purple-top) tube
 1-2 cc in Sodium Heparin (green-top) tube
 3-5 cc in ACD (yellow-top) tube
 URINE: 2-4 cc. Do not add preservatives. Urine must be shipped frozen with dry ice.

NEONATAL AND INFANTILE METABOLIC SEIZURES PANEL

4400 This panel includes the following tests:
 PLASMA: Acylcarnitine Analysis (4300); Amino Acid Analysis (4100); Creatine/Guanidinoacetate Determination (4130); Pyridoxine-Dependent Seizures Panel (4811)
 CSF: Amino Acid Analysis (4160); Pyridoxine-Dependent Seizures Panel (4812)
 SERUM: Biotinidase Deficiency (4555)
 URINE: Organic Acid Screen (4200); Purine and Pyrimidine Panel (4010); Sulfocysteine Determination (4225)

SPECIMEN REQUIREMENTS:
 All sample types below must be shipped frozen with dry ice and overnight:
 PLASMA: 2-3cc
 CSF: 0.5-1cc
 SERUM: 1-2 cc
 URINE: 2-4 cc. Do not add preservatives.

BIOCHEMISTRY MULTI-PLEX AUTISM PANEL

4000 This panel includes the following tests:
 PLASMA: Acylcarnitine Analysis (4300); Amino Acid Analysis (4100); Creatine/Guanidinoacetate Determination (4130)
 URINE: Carnitine Biosynthesis Panel (4135); Creatine/Guanidinoacetate Determination (4260); Organic Acid Screen (4200); Purine Panel (4220); Pyrimidine Panel (4215)

SPECIMEN REQUIREMENTS:
 All sample types below must be shipped frozen with dry ice and overnight:
 PLASMA: 2cc
 URINE: 3-5cc

BIOCHEMISTRY 5-PLEX AUTISM PANEL (URINE ONLY)

4165 This panel includes the following tests:
 URINE: Carnitine Biosynthesis Panel (4135); Creatine/Guanidinoacetate Determination (4260); Organic Acid Screen (4200); Purine Panel (4220); Pyrimidine Panel (4215)

SPECIMEN REQUIREMENTS:
 All sample types below must be shipped frozen with dry ice and overnight:
 URINE: 3-5cc

BIOCHEMISTRY 3-PLEX AUTISM PANEL (PLASMA ONLY)

4175 This panel includes the following tests:
 PLASMA: Acylcarnitine Analysis (4300); Amino Acid Analysis (4100); Creatine/Guanidinoacetate Determination (4130)

SPECIMEN REQUIREMENTS:
 All sample types below must be shipped frozen with dry ice and overnight:
 PLASMA: 2cc

ANALYTE PANEL TESTS

Carnitine Biosynthesis Panel

<input type="checkbox"/>	4135	URINE
<input type="checkbox"/>	4145	PLASMA
<input type="checkbox"/>	4155	CSF

Creatine Deficiency Syndromes Panel

<input type="checkbox"/>	4015	PLASMA AND URINE
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Polyols

<input type="checkbox"/>	4340	URINE
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Purine Panel

<input type="checkbox"/>	4220	URINE
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Purine and Pyrimidine Panel

<input type="checkbox"/>	4010	URINE
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Pyridoxine-Dependent Seizures Panel

<input type="checkbox"/>	4811	PLASMA
<input type="checkbox"/>	4812	CSF

Pyrimidine Panel

<input type="checkbox"/>	4215	URINE
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INDICATIONS ARE REQUIRED. PLEASE FILL OUT CHECKLIST ON PAGE 4

SEE PAGE 3 FOR INDIVIDUAL ANALYTE AND ENZYME TESTS

BIOCHEMICAL TESTING REQUISITION - INSTITUTIONAL ONLY

NAME: _____ DATE OF BIRTH: ____ / ____ / ____ GENDER (Please select one): FEMALE
LAST NAME FIRST NAME MI MM DD YY MALE UNKNOWN

INDIVIDUAL ANALYTE ANALYSIS

25-Hydroxyvitamin D2 and D3 <input type="checkbox"/> 4360 SERUM Acylcarnitine Analysis <input type="checkbox"/> 4300 PLASMA Acylglycine Determination <input type="checkbox"/> 4350 URINE Amino Acid Analysis <input type="checkbox"/> 4100 PLASMA <input type="checkbox"/> 4160 CSF <input type="checkbox"/> 4240 URINE Blood spot Phe/Tyr <input type="checkbox"/> 4120 Blood spot Carnitine Determination <input type="checkbox"/> 4310 PLASMA	Coenzyme Q10 Determination <input type="checkbox"/> 4800 MUSCLE Creatine/Guanidinoacetate Determination <input type="checkbox"/> 4130 PLASMA <input type="checkbox"/> 4260 URINE Cystine Determination <input type="checkbox"/> 4627 WHITE BLOOD CELLS Homocysteine Determination <input type="checkbox"/> 4140 PLASMA Methylmalonic Acid <input type="checkbox"/> 4150 PLASMA Organic Acid Screen <input type="checkbox"/> 4200 URINE Orotic/Orotidine Determination <input type="checkbox"/> 4210 URINE	Phenylalanine Determination <input type="checkbox"/> 4110 PLASMA Phenylbutyrate Metabolite Analysis <input type="checkbox"/> 4650 PLASMA <input type="checkbox"/> 4651 URINE <input type="checkbox"/> 4652 CSF Sulfocysteine Determination <input type="checkbox"/> 4225 URINE Succinylacetone Determination <input type="checkbox"/> 4250 URINE Succinyladenosine <input type="checkbox"/> 4180 CSF Thymidine Determination <input type="checkbox"/> 4330 PLASMA
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INDIVIDUAL ENZYME ANALYSIS

Adenosine Deaminase Deficiency <input type="checkbox"/> 4509 RED BLOOD CELLS* <input type="checkbox"/> 4510 SKIN FIBROBLAST CULTURE <input type="checkbox"/> 4511 WHITE BLOOD CELLS Argininemia / Arginase Deficiency <input type="checkbox"/> 4535 LIVER <input type="checkbox"/> 4536 RED BLOOD CELLS* Argininosuccinic Aciduria / Argininosuccinate Lyase Deficiency <input type="checkbox"/> 4523 LIVER <input type="checkbox"/> 4524 RED BLOOD CELLS* <input type="checkbox"/> 4525 SKIN FIBROBLAST CULTURE Aspartylglucosaminuria / Aspartylglucosaminidase Deficiency <input type="checkbox"/> 4514 SKIN FIBROBLAST CULTURE Biotinidase Deficiency <input type="checkbox"/> 4555 SERUM Carbamoyl Phosphate Synthetase I Deficiency <input type="checkbox"/> 4561 LIVER Citrullinemia / Argininosuccinate Synthetase Deficiency <input type="checkbox"/> 4544 LIVER <input type="checkbox"/> 4545 SKIN FIBROBLAST CULTURE* Fabry Disease/Alpha-Galactosidase A <input type="checkbox"/> 4516 SKIN FIBROBLAST CULTURE <input type="checkbox"/> 4517 WHITE BLOOD CELLS*	Gaucher Disease / Beta-Glucosidase <input type="checkbox"/> 4553 SKIN FIBROBLAST CULTURE* <input type="checkbox"/> 4554 WHITE BLOOD CELLS GM1 Gangliosidosis (Morquio B, MPS IVB) / Beta-Galactosidase <input type="checkbox"/> 4548 SKIN FIBROBLAST CULTURE <input type="checkbox"/> 4549 WHITE BLOOD CELLS* Krabbe Disease / Galactocerebrosidase <input type="checkbox"/> 4565 SKIN FIBROBLAST CULTURE <input type="checkbox"/> 4566 WHITE BLOOD CELLS* Lowe Syndrome / Phosphatidylinositol Bisphosphate Phosphatase <input type="checkbox"/> 4585 SKIN FIBROBLAST CULTURE Lesch-Nyhan Disease / Hypoxanthine-Guanine Phosphoribosyltransferase <input type="checkbox"/> 4573 SKIN FIBROBLAST CULTURE Metachromatic Leukodystrophy / Arylsulfatase A Deficiency <input type="checkbox"/> 4537 SKIN FIBROBLAST CULTURE <input type="checkbox"/> 4538 WHITE BLOOD CELLS* Mucopolipidosis I (Sialidosis) / Sialidase <input type="checkbox"/> 4603 SKIN FIBROBLAST CULTURE Mucopolysaccharidosis Type I / Alpha-L-iduronidase <input type="checkbox"/> 4575 SKIN FIBROBLAST CULTURE <input type="checkbox"/> 4576 WHITE BLOOD CELLS*	Niemann-Pick Disease Types A & B / Sphingomyelinase <input type="checkbox"/> 4607 SKIN FIBROBLAST CULTURE <input type="checkbox"/> 4608 WHITE BLOOD CELLS* Ornithine Transcarbamylase Deficiency <input type="checkbox"/> 4582 LIVER Purine Nucleoside Phosphorylase Deficiency <input type="checkbox"/> 4592 RED BLOOD CELLS* <input type="checkbox"/> 4593 SKIN FIBROBLAST CULTURE <input type="checkbox"/> 4594 WHITE BLOOD CELLS Tay-Sachs Disease & Sandhoff Disease / Hexosaminidase A and B <input type="checkbox"/> 4569 SERUM <input type="checkbox"/> 4617 SERUM** <input type="checkbox"/> 4620 WHITE BLOOD CELLS** Wolman Disease / Cholesteryl Ester Storage Disease / Acid Lipase Deficiency <input type="checkbox"/> 4502 LIVER <input type="checkbox"/> 4503 SKIN FIBROBLAST CULTURE <input type="checkbox"/> 4504 WHITE BLOOD CELLS* X-Linked Ichthyosis / Steroid Sulfatase <input type="checkbox"/> 4614 SKIN FIBROBLAST CULTURE <input type="checkbox"/> 4615 WHITE BLOOD CELLS
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*PREFERRED SAMPLE TYPE
 **TAY-SACHS DISEASE CARRIER TESTING

INDICATIONS ARE REQUIRED. PLEASE FILL OUT CHECKLIST ON PAGE 4

INDICATION CHECKLIST

PATIENT NAME: _____

LAST NAME

FIRST NAME

MI

 Clinical management of known diagnosis - Please specify: _____

 Diagnostic Testing - Please complete checklist below.

Please indicate whether each feature is PRESENT by checking the box beside the indication below

CENTRAL NERVOUS SYSTEM

<input type="checkbox"/>	101	dd	Developmental Delay/ID
<input type="checkbox"/>	102	ht	Hypotonia
<input type="checkbox"/>	103	au	Autistic Features
<input type="checkbox"/>	104	enc	Dementia/Encephalopathy
<input type="checkbox"/>	105	ha	Headaches/Migraines
<input type="checkbox"/>	106	stk	Stroke, Ischemic Episodes
<input type="checkbox"/>	107	atx	Ataxia
<input type="checkbox"/>	108	sz	Intractable/Refractory/Myoclonus/Myoclonic Seizures
<input type="checkbox"/>	109	pi	Perinatal Insult
<input type="checkbox"/>	110	ps	Pyramidal Signs
<input type="checkbox"/>	111	hp	Hemiparesis
<input type="checkbox"/>	112	spas	Spasticity
<input type="checkbox"/>	113	dyst	Dystonia
<input type="checkbox"/>	114	cho	Chorea
<input type="checkbox"/>	115	sib	Self-Injury
<input type="checkbox"/>	116	sd	Language Problems/Speech Delay
<input type="checkbox"/>	117	fp	Feeding Problems
<input type="checkbox"/>	118	es	Excessive Sleepiness/Sleep Disturbance
<input type="checkbox"/>	119	let	Lethargy
<input type="checkbox"/>	120	cm	Coma

NEUROMUSCULAR

<input type="checkbox"/>	201	pn	Peripheral Neuropathy
<input type="checkbox"/>	202	exi	Exercise Intolerance
<input type="checkbox"/>	203	pmw	Progressive Muscle Weakness
<input type="checkbox"/>	204	smw	Static Muscle Weakness
<input type="checkbox"/>	205	cr	Muscle Cramps after Exercise
<input type="checkbox"/>	206	fat	Easy Fatigability
<input type="checkbox"/>	207	dcmyo	Dilated Cardiomyopathy
<input type="checkbox"/>	208	hcmyo	Hypertrophic Cardiomyopathy
<input type="checkbox"/>	209	hb	Heart Block
<input type="checkbox"/>	210	ar	Arrhythmia
<input type="checkbox"/>	211	op	Ophthalmoparesis, CPEO
<input type="checkbox"/>	212	emg	Abnormal EMG/NCV
<input type="checkbox"/>	213	pto	Ptosis
<input type="checkbox"/>	214	eh	Cardiomegaly/Enlarged heart

VISCERAL

<input type="checkbox"/>	301	gir	Gastrointestinal Reflux
<input type="checkbox"/>	302	dge	Delayed Gastric Emptying
<input type="checkbox"/>	303	pan	Pancreatitis
<input type="checkbox"/>	304	dia	Diarrhea
<input type="checkbox"/>	305	cst	Constipation
<input type="checkbox"/>	306	cv	Cyclic Vomiting
<input type="checkbox"/>	307	pob	Pseudoobstruction
<input type="checkbox"/>	308	hpf	Hepatic Failure

VISCERAL (cont.)

<input type="checkbox"/>	309	eta	Elevated Transaminases
<input type="checkbox"/>	310	rtd	Renal Tubular Disease
<input type="checkbox"/>	311	ap	Apnea/Hypoventilation
<input type="checkbox"/>	312	rsf	Respiratory Deficiency/Failure
<input type="checkbox"/>	313	ren	Renal Dysfunction
<input type="checkbox"/>	314	lc	Liver Carcinoma
<input type="checkbox"/>	315	jau	Jaundice
<input type="checkbox"/>	316	spm	Splenomegaly/Enlarged Spleen
<input type="checkbox"/>	317	hpm	Hepatomegaly/Enlarged Liver
<input type="checkbox"/>	318	hd	Hepatic Dysfunction

METABOLITES/METABOLIC

<input type="checkbox"/>	400	nbs	Abnormal Newborn Screen: _____
<input type="checkbox"/>	401	kto	Ketosis
<input type="checkbox"/>	402	dca	Dicarboxylic Aciduria
<input type="checkbox"/>	403	la	Lactic Acidosis
<input type="checkbox"/>	404	csfl	High CSF Lactate
<input type="checkbox"/>	405	oa	Organic Aciduria: _____
<input type="checkbox"/>	406	lpc	Low Plasma Carnitine
<input type="checkbox"/>	407	cpk	CPK abnormalities
<input type="checkbox"/>	408	pyr	Elevated Pyruvate
<input type="checkbox"/>	409	ala	Elevated Alanine
<input type="checkbox"/>	410	3mg	3-Methylglutaconic Aciduria
<input type="checkbox"/>	411	acid	Acidosis
<input type="checkbox"/>	412	NH3	Hyperammonemia
<input type="checkbox"/>	413	hypo	Hypoglycemia
<input type="checkbox"/>	414	hyper	Hyperglycemia
<input type="checkbox"/>	415	uco	Unusual Color/Odor

SENSORY

<input type="checkbox"/>	501	rp	Retinitis Pigmentosa
<input type="checkbox"/>	502	opa	Optic Atrophy
<input type="checkbox"/>	503	cat	Cataract
<input type="checkbox"/>	504	hl	Sensorineural Hearing Loss
<input type="checkbox"/>	505	trv	Tortuous Retinal Vessels
<input type="checkbox"/>	506	crs	Cherry Red Spot/Eye
<input type="checkbox"/>	507	co	Corneal Opacity
<input type="checkbox"/>	508	el	Ectopia Lentis
<input type="checkbox"/>	509	pp	Photophobia

ENDOCRINE

<input type="checkbox"/>	601	db	Diabetes
<input type="checkbox"/>	602	pd	Exocrine/Pancreatic Deficiency
<input type="checkbox"/>	603	gf	Gonadal Failure
<input type="checkbox"/>	604	hth	Hypothyroidism
<input type="checkbox"/>	605	hpt	Hypoparathyroidism
<input type="checkbox"/>	606	adr	Hypo/Hyper-adrenal Function
<input type="checkbox"/>	607	ss	Short Stature

ENDOCRINE (cont.)

<input type="checkbox"/>	608	adc	Adrenal Calcification
<input type="checkbox"/>	609	hf	Hydrops Fetalis
<input type="checkbox"/>	610	pg	Pregnant

OTHER CLINICAL

<input type="checkbox"/>	701	ftt	Failure to Thrive
<input type="checkbox"/>	702	mce	Microcephaly
<input type="checkbox"/>	703	sids	SIDS/Unexplained Death
<input type="checkbox"/>	704	ca	Congenital Anomalies
<input type="checkbox"/>	705	dys	Dysmorphic Features
<input type="checkbox"/>	706	id	Immunodeficiency
<input type="checkbox"/>	707	ma	Macrocytic Anemia
<input type="checkbox"/>	708	pcbm	Pancytopenia/Bone Marrow Failure
<input type="checkbox"/>	709	np	Neutropenia
<input type="checkbox"/>	710	mc	Macrocephaly
<input type="checkbox"/>	711	cf	Course Features
<input type="checkbox"/>	712	sa	Skeletal Anomalies
<input type="checkbox"/>	713	art	Arthritis

HAIR/SKIN FINDINGS

<input type="checkbox"/>	714	rash	Rashes w/Hypopigmentation
<input type="checkbox"/>	715	htii	Hypertrichosis
<input type="checkbox"/>	716	alp	Alopecia
<input type="checkbox"/>	717	ac	Acrocyanosis
<input type="checkbox"/>	718	ak	Angiokeratoma
<input type="checkbox"/>	719	ic	Ichthyosis

FAMILY HISTORY

<input type="checkbox"/>	001	mut	Mutation*
<input type="checkbox"/>	002	mi	Evidence of Maternal Inheritance

ELECTROPHYSIOLOGY

<input type="checkbox"/>	801	baers	Abnormal BAERS
<input type="checkbox"/>	802	vers	Abnormal VERS
<input type="checkbox"/>	803	eeg	Abnormal EEG

IMAGING/OTHER STUDIES

<input type="checkbox"/>	804	bg	Increased Signal Basal Ganglia
<input type="checkbox"/>	805	dmy	Delayed Myelination
<input type="checkbox"/>	806	cea	Cerebellar Atrophy
<input type="checkbox"/>	807	pstk	Posterior Stroke
<input type="checkbox"/>	808	leuk	Leukodystrophy
<input type="checkbox"/>	809	mrs	MRS/Lactate Peak
<input type="checkbox"/>	810	mri	Abnormal MRI

MUSCLE BIOPSY

<input type="checkbox"/>	901	his	Abnormal Histology
<input type="checkbox"/>	902	em	Abnormal Ultrastructure (EM)
<input type="checkbox"/>	903	enz	Abnormal Respiratory Enzymes
<input type="checkbox"/>	904	prol	Large Mitochondria/Proliferation
<input type="checkbox"/>	905	cox	COX Deficiency
<input type="checkbox"/>	906	rrf	Ragged Red Fibers

* ATTACH DETAILS

If more detailed clinical information is required, please provide the name, e-mail address, and phone number of the contact person below.