Complete and print.

Reset Form

MAYO CLINIC LABORATORIES

Biochemical Genetics Test Request

Client Information (required)

Client Name		
Client Account No.		
Client Phone	Client Order No.	
Street Address		
City	State	ZIP Code

Patient Information (required)

Patient ID (Medical Record No.)		
Patient Name (Last, First Middle)		
Sex ☐ Male ☐ Female	Birth Date (mm-dd-yyyy)	
Collection Date (mm-dd-yyyy)	Time	☐ am ☐ pm

Submitting Healthcare Professional Information (required)

Submitting Healthcare Professional Name (Last, First)	
Title/Credentials	
Phone (with area code)	Fax* (with area code)
National Provider Identification (NPI)	
Email**	

**Any communication sent via email will comply with applicable HIPAA regulations.*
**Fax number given must be from a fax machine that complies with applicable HIPAA regulation.*

Note: It is the client's responsibility to maintain documentation of the order.

New York State Patients: Informed Consent for Genetic Testing

"I hereby confirm that informed consent has been signed by an individual legally authorized to do so and is on file with this office or the individual's provider's office."

Signature ▶

Note: Test requests without a signature will not be performed.

Reason for Testing (required)

Has molecular/DNA testing already been performed?
☐ Yes ☐ No If Yes, results:

Gene	Variant	Classification
Gene	Variant	Classification

For molecular testing options, see www.MayoClinicLabs.com

MCL Internal Use Only

Ship specimens to:
Mayo Clinic Laboratories
3050 Superior Drive NW
Rochester, MN 55905

Customer Service: 800-533-1710

Visit www.MayoClinicLabs.com for the most up-to-date test and shipping information.

Billing Information
- An itemized invoice will be sent each month.
- Payment terms are net 30 days.

Call the Business Office with billing-related questions:
800-447-6424 (US and Canada)
507-266-5490 (outside the US)

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AMINO ACID METABOLISM

- ☐ AAQP Amino Acids, Quantitative, Plasma
- ☐ AAPD Amino Acids, Quantitative, Random, Urine
- ☐ AACSF Amino Acids, Quantitative, Spinal Fluid
- ☐ AACYL Aminoacylase-1 Deficiency, Urine
- ☐ TRYPP Tryptophan, Plasma
- ☐ TRYPU Tryptophan, Random, Urine

Cystinuria

- ☐ CYSGP Cystinuria Gene Panel
- ☐ CYSQN Cystinuria Profile, Quantitative, 24 Hour, Urine
- ☐ CYSR Cystinuria Profile, Quantitative, Random, Urine

Homocystinuria

- ☐ CMMPP Cobalamin, Methionine, and Methylmalonic Acid Pathways, Plasma
- ☐ CMMPS Cobalamin, Methionine, and Methylmalonic Acid Pathways, Serum
- ☐ HCYSP Homocysteine, Total, Plasma
- ☐ HCYSS Homocysteine, Total, Serum

Maple Syrup Urine Disease

- ☐ ALLOI Allo-isoleucine, Blood Spot
- ☐ AAMSD Amino Acids, Maple Syrup Urine Disease Panel, Plasma
- ☐ MSUSC Branched-Chain Amino Acids, Self-Collect, Blood Spot
- ☐ MSUDP Maple Syrup Urine Disease Gene Panel

Phenylketonuria

- ☐ PKUBS Phenylalanine and Tyrosine, Blood Spot
- ☐ PHEGP Phenylalanine Disorders Gene Panel
- ☐ PKU Phenylalanine and Tyrosine, Plasma
- ☐ PKUSC Phenylalanine and Tyrosine, Self-Collect, Blood Spot

Tyrosinemia

- ☐ TYRGP Tyrosine Disorders Gene Panel
- ☐ TYRBS Tyrosinemia Follow up Panel, Blood Spot
- ☐ TYRSC Tyrosinemia Follow up panel, Self-Collect, Blood Spot
- ☐ SUAC Succinylacetone, Blood Spot

CARBOHYDRATE METABOLISM

Congenital Disorders of Glycosylation

- ☐ CDG Carbohydrate Deficient Transferrin for Congenital Disorders of Glycosylation, Serum
- ☐ CDGGP Congenital Disorders of Glycosylation Gene Panel
- ☐ CDGN Congenital Disorders of N-Glycosylation, Serum
- ☐ OLIGU Oligosaccharide Screen, Random, Urine
- ☐ PMMIL Phosphomannomutase and Phosphomannose Isomerase, Leukocytes
- ☐ SORBU Sorbitol and Mannitol, Quantitative, Random, Urine

Galactosemia

- ☐ GATOL Galactitol, Quantitative, Urine
- ☐ GALK Galactokinase, Blood
- ☐ GAL1P Galactose-1-Phosphate, Erythrocytes
- ☐ GALT Galactose-1-Phosphate Uridyltransferase, Blood
- ☐ GALTP Galactose-1-Phosphate Uridyltransferase Biochemical Phenotyping, Erythrocytes
- ☐ GALP Galactose, Quantitative, Plasma
- ☐ GALZ Galactosemia, GALT Gene, Full Gene Analysis
- ☐ GCT Galactosemia Reflex, Blood
- ☐ GALE Uridine Diphosphate-Galactose 4' Epimerase, Blood

Transaldolase and Ribose-5-phosphate (RPI) Deficiencies

- ☐ TALDO Polyols, Quantitative, Urine

CHOLESTATIC LIVER DISEASE

- ☐ CHLGP Cholestasis Gene Panel

CHOLESTEROL BIOSYNTHESIS AND TRANSPORT

- ☐ CTXWB Cerebrotendinous Xanthomatosis, Blood
- ☐ CTXBS Cerebrotendinous Xanthomatosis, Blood Spot
- ☐ CTXP Cerebrotendinous Xanthomatosis, Plasma
- ☐ HSMBS Hepatosplenomegaly Panel, Blood Spot
- ☐ HSMWB Hepatosplenomegaly Panel, Blood
- ☐ HSMP Hepatosplenomegaly Panel, Plasma
- ☐ OXYWB Oxysterols, Blood
- ☐ OXYBS Oxysterols, Blood Spots
- ☐ OXNP Oxysterols, Plasma
- ☐ SLO Smith-Lemli-Opitz Screen, Plasma
- ☐ DHCRZ Smith Lemli Opitz Syndrome, DHCR7 Gene, Full Gene Analysis
- ☐ STER Sterols, Plasma

CONGENITAL ADRENAL HYPERPLASIA

- ☐ CAH2T Congenital Adrenal Hyperplasia Newborn Screen, Blood Spot
- ☐ CAH21 Congenital Adrenal Hyperplasia (CAH) Profile for 21-Hydroxylase Deficiency, Serum
- ☐ CYPZ 21-Hydroxylase Gene, CYP21A2, Full Gene Analysis

CONGENITAL LACTIC ACIDOSIS

- ☐ CLADP Congenital Lactic Acidosis Gene Panel

CREATINE DISORDERS

- ☐ CRDPP Creatine Disorders Panel, Plasma
- ☐ CRDPU Creatine Disorders Panel, Random, Urine

CUSTOM GENE PANEL

- ☐ CGPH Custom Gene Panel, Hereditary, Next-Generation Sequencing
  Gene List ID (if known) or Genes Requested for Testing:

FAMILIAL AMYLOIDOSIS

- ☐ TTRX Amyloidosis, Transthyretin-Associated Familial, Reflex, Blood
- ☐ TTRZ *TTR* Gene, Full Gene Analysis

FATTY ACID METABOLISM (BETA-OXIDATION)

- ☐ ACRN Acylcarnitines, Quantitative, Plasma
- ☐ ACRNS Acylcarnitines, Quantitative, Serum
- ☐ AGU20 Acylglycines, Quantitative, Random, Urine
- ☐ C4U C4 Acylcarnitine, Quantitative, Random, Urine
- ☐ CARN Carnitine, Plasma
- ☐ CARNS Carnitine, Serum
- ☐ CARNU Carnitine, Random, Urine
- ☐ HFAOP Fatty Acid Oxidation Gene Panel
- ☐ FAO Fatty Acid Oxidation Probe Assay, Fibroblast Culture
- ☐ PFAPC Fatty Acid Profile, Comprehensive (C8-C26), Plasma
- ☐ FAPCP Fatty Acid Profile, Comprehensive (C8-C26), Serum
- ☐ FAPM Fatty Acid Profile, Mitochondrial (C8-C18), Serum
- ☐ MCADZ Medium-Chain Acyl-CoA Dehydrogenase (MCAD) Deficiency Full Gene Analysis
- ☐ OAU Organic Acids Screen, Random, Urine
- ☐ VLCZ Very Long Chain Acyl-CoA Dehydrogenase Deficiency, Full Gene Analysis

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GLYCOGEN STORAGE DISORDERS

- ☐ GSDGP Glycogen Storage Disease Gene Panel

HYPEROXALURIA

- ☐ HYOX Hyperoxaluria Panel, Random, Urine
- ☐ RSCGP Nephrocalcinosis, Nephrolithiasis, and Renal Electrolyte Imbalance Gene Panel

LYSOSOMAL METABOLISM AND STORAGE DISORDERS

- ☐ CTSU Ceramide Trihexosides and Sulfatides, Random, Urine
- ☐ HSMWB Hepatosplenomegaly Panel, Blood
- ☐ HSMP Hepatosplenomegaly Panel, Plasma
- ☐ PLSD Lysosomal and Peroxisomal Storage Disorders Screen, Blood Spot
- ☐ LSDGP Lysosomal Storage Disease Gene Panel
- ☐ LSDS Lysosomal Storage Disorders Screen, Random, Urine
- ☐ LSD6W Lysosomal Storage Disorders, Six-Enzyme Panel, Leukocytes
- ☐ MPSQU Mucopolysaccharides Quantitative, Random, Urine
- ☐ MP8BS Mucopolysaccharidoses, Eight-Enzyme Panel, Blood Spot
- ☐ OLIGU Oligosaccharide Screen, Random, Urine
- ☐ OXNP Oxysterols, Plasma

Fabry Disease

- ☐ FABRZ Krabbe Disease, Full Gene Analysis
- ☐ AGABS Alpha-Galactosidase, Blood Spot
- ☐ AGAW Alpha-Galactosidase, Leukocytes
- ☐ AGAS Alpha-Galactosidase, Serum
- ☐ CTSU Ceramide Trihexosides and Sulfatides, Random, Urine
- ☐ LGB3S Globotriaosylsphingosine, Serum

Fucosidosis

- ☐ FUCW Alpha-Fucosidase, Leukocytes

Gaucher Disease

- ☐ GBAW Beta-Glucosidase, Leukocytes
- ☐ GBAZ Gaucher Disease, Full Gene Analysis
- ☐ GPSYW Glucopsychosine, Blood
- ☐ GPSY Glucopsychosine, Blood Spot
- ☐ GPSYP Glucopsychosine, Plasma

GM1 Gangliosidosis

- ☐ BGA Beta-Galactosidase, Leukocytes
- ☐ MPS4B Mucopolysaccharidosis IV Enzyme Panel, Blood Spot
- ☐ MPS4W Mucopolysaccharidosis IV Enzyme Panel, Leukocytes

Krabbe Disease

- ☐ GALCW Galactocerebrosidase, Leukocytes
- ☐ KRABZ Krabbe Disease, Full Gene Analysis and Large (30 kb) Deletion
- ☐ PSY Psychosine, Blood Spot
- ☐ PSYCF Psychosine, Spinal Fluid
- ☐ PSYR Psychosine, Whole Blood

Lysosomal Acid Lipase Deficiency

- ☐ LALB Lysosomal Acid Lipase, Blood
- ☐ LALBS Lysosomal Acid Lipase, Blood Spot

Mannosidosis

- ☐ MANN Alpha-Mannosidase, Leukocytes

Metachromatic Leukodystrophy

- ☐ ARSU Arylsulfatase A, 24 Hour, Urine
- ☐ ARSAW Arylsulfatase A, Leukocytes
- ☐ CTSU Ceramide Trihexosides and Sulfatides, Random, Urine

Mucopolysaccharidoses (MPS)

- ☐ MPSQU Mucopolysaccharides Quantitative, Random, Urine
- ☐ MPSER Mucopolysaccharides Quantitative, Serum
- ☐ MPSWB Mucopolysaccharidosis, Blood
- ☐ MPSBS Mucopolysaccharidosis, Blood Spot

MPS Type I (Hurler/Scheie syndrome)

- ☐ IDUAW Alpha-L-Iduronidase, Leukocytes
- ☐ MPS1B Endogenous Mucopolysaccharidosis Type I (IDUA [Alpha-L-Iduronidase]) Biomarker, Blood Spot
- ☐ MPS1R Endogenous Mucopolysaccharidosis Type I (IDUA [Alpha-L-Iduronidase]) Biomarker Reflex, Blood Spot
- ☐ MPS1Z Hurler Syndrome, Full Gene Analysis

MPS Type II (Hunter syndrome)

- ☐ MPS2R Endogenous Mucopolysaccharidosis Type II (I2S [Iduronate-2-Sulfatase]) Biomarker Reflex, Blood Spot
- ☐ MPS2B Endogenous Mucopolysaccharidosis Type II (I2S [Iduronate-2-Sulfatase]) Biomarker, Blood Spot
- ☐ I2SB Iduronate-2-Sulfatase, Blood Spot
- ☐ I2SWB Iduronate-2-Sulfatase, Leukocytes
- ☐ MPS2Z Hunter Syndrome, Full Gene Analysis

MPS Type III (Sanfilippo syndrome)

- ☐ MPS3B Mucopolysaccharidosis III, Three-Enzyme Panel, Blood Spot
- ☐ MPS3W Mucopolysaccharidosis III, Four-Enzyme Panel, Leukocytes

MPS Type IV (Morquio syndrome)

- ☐ BGA Beta-Galactosidase, Leukocytes
- ☐ MPS4B Mucopolysaccharidosis IV Enzyme Panel, Blood Spot
- ☐ MPS4W Mucopolysaccharidosis IV Enzyme Panel, Leukocytes

MPS VI (Maroteaux-Lamy syndrome)

- ☐ ARSBB Arylsulfatase B, Blood Spot
- ☐ ARSBW Arylsulfatase B, Leukocytes

MPS VII (Sly syndrome)

- ☐ GUSBW Beta-Glucuronidase, Leukocytes
- ☐ GUSBB Beta-Glucuronidase, Blood Spot

Multiple Sulfatase Deficiency

- ☐ MSDBS Multiple Sulfatase Deficiency, Blood Spot
- ☐ MSDW Multiple Sulfatase Deficiency, Leukocytes

Niemann-Pick Types A and B

- ☐ ASMW Acid Sphingomyelinase, Leukocytes
- ☐ OXNP Oxysterols, Plasma

Niemann-Pick Type C

- ☐ OXNP Oxysterols, Plasma

Neuronal Ceroid Lipofuscinoses

- ☐ NCLGP Neuronal Ceroid Lipofuscinosis (Batten Disease) Gene Panel
- ☐ NCLBS Neuronal Ceroid Lipofuscinosis, Two-Enzyme Panel, Blood Spot
- ☐ NCLW Neuronal Ceroid Lipofuscinosis, Two-Enzyme Panel, Leukocytes

Pompe Disease

- ☐ GAAW Acid Alpha-Glucosidase, Leukocytes
- ☐ GAAZ Pompe Disease, Full Gene Analysis
- ☐ HEX4 Glucotetrasaccharides, Random, Urine
- ☐ PDBS Pompe Disease, Blood Spot

Tay-Sachs and Sandhoff Diseases

- ☐ NAGW Hexosaminidase A and Total Hexosaminidase, Leukocytes
- ☐ NAGS Hexosaminidase A and Total Hexosaminidase, Serum
- ☐ NAGR Hexosaminidase A and Total, Leukocytes/Molecular Reflex, Whole Blood
- ☐ MUGS Hexosaminidase A, Serum
- ☐ HEXBZ Sandhoff Disease, HEXB Gene, Full Gene Analysis
- ☐ HEXAZ Tay-Sachs Disease, HEXA Gene, Full Gene Analysis

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Birth Date (mm-dd-yyyy)	

MITOCHONDRIAL DISEASES

- ☐ Q10 — Coenzyme Q10, Reduced and Total, Plasma
- ☐ TQ10 — Coenzyme Q10, Total, Plasma
- ☐ FAPM — Fatty Acid Profile, Mitochondrial (C8-C18), Serum
- ☐ CMITO — Combined Mitochondrial Full Genome and Nuclear Gene Panel
- ☐ DMITO — Mitochondrial DNA Deletion Heteroplasmy, ddPCR
- ☐ GDF15 — Growth Differentiation Factor 15, Plasma
- ☐ LAPYP — Lactate Pyruvate Panel, Plasma
- ☐ MITOP — Mitochondrial Full Genome Analysis, Next-Generation Sequencing (NGS)
- ☐ MMPP — Mitochondrial Metabolites, Plasma
- ☐ NMITO — Nuclear Mitochondrial Gene Panel, Next-Generation Sequencing
- ☐ OAU — Organic Acids Screen, Random, Urine
- ☐ PYRC — Pyruvate, Spinal Fluid
- ☐ PYR — Pyruvic Acid, Blood

NEUROLOGIC DISORDERS

- ☐ FFRWB — Friedreich Ataxia, Frataxin, Quantitative, Blood
- ☐ FFRBS — Friedreich Ataxia, Frataxin, Quantitative, Blood Spot
- ☐ AFXN — Friedreich Ataxia, Repeat Expansion Analysis
- ☐ SORD — Sorbitol and Xylitol, Quantitative, Random, Urine

NEWBORN SCREENING

Screening Panels

- ☐ LDALD — Lysosomal and Peroxisomal Disorders Newborn Screen, Blood Spot
- ☐ SNS — Supplemental Newborn Screen, Blood Spot

Second Tier Tests

- ☐ ALLOI — Allo-isoleucine, Blood Spot
- ☐ CAH2T — Congenital Adrenal Hyperplasia Newborn Screen, Blood Spot
- ☐ GPSY — Glucopsychosine, Blood Spot
- ☐ HCMM — Homocysteine (Total), Methylmalonic Acid, and Methylcitric Acid, Blood Spot
- ☐ HGEM — Hydroxyglutaric Acids, Glutaric Acid, Ethylmalonic Acid, and Methylsuccinic Acid, Blood Spot
- ☐ LPCBS — Lysophosphatidylcholines, LC MS/MS, Blood Spot
- ☐ MPSBS — Mucopolysaccharidosis, Blood Spot
- ☐ OXYBS — Oxysterols, Blood Spot
- ☐ PD2T — Pompe Disease Second-Tier Newborn Screening, Blood Spot
- ☐ PSY — Psychosine, Blood Spot
- ☐ SUAC — Succinylacetone, Blood Spot

ORGANIC ACID METABOLISM

- ☐ 3MGAP — 3-Methylglutaconic Aciduria Gene Panel
- ☐ AGU20 — Acylglycines, Quantitative, Random, Urine
- ☐ C5OHU — C5-OH Acylcarnitine, Quantitative, Random, Urine
- ☐ KETGP — Ketone Disorders Gene Panel
- ☐ NAACD — N-Acetylaspartic Acid, Canavan Disease, Random, Urine
- ☐ OAU — Organic Acids Screen, Random, Urine
- ☐ OAUS — Organic Acid Screen, Urine Spot

2-Hydroxyglutaric Aciduria

- ☐ 2OHGP — 2-Hydroxyglutaric Aciduria Gene Panel
- ☐ 2HGA — 2-Hydroxyglutaric Acid Chiral Analysis, Quantitative, Random, Urine

Biotinidase Deficiency

- ☐ BIOTS — Biotinidase, Serum
- ☐ BTDZ — Biotinidase Deficiency, *BTD* Full Gene Analysis

Glutaric Acidemia

- ☐ C5DCU — C5-DC Acylcarnitine, Quantitative, Random, Urine
- ☐ GA2P — Glutaric Aciduria Type II Gene Panel
- ☐ HGEM — Hydroxyglutaric Acids, Glutaric Acid, Ethylmalonic Acid, and Methylsuccinic Acid, Blood Spot
- ☐ HGEMP — Hydroxyglutaric Acids, Glutaric Acid, Ethylmalonic Acid, and Methylsuccinic Acid, Plasma
- ☐ HGEMS — Hydroxyglutaric Acids, Glutaric Acid, Ethylmalonic Acid, and Methylsuccinic Acid, Serum
- ☐ TRYPP — Tryptophan, Plasma
- ☐ TRYPU — Tryptophan, Random, Urine

Methylmalonic Acidemia/Cobalamin/ Propionic Acidemia

- ☐ CMMPP — Cobalamin, Methionine, and Methylmalonic Acid Pathways, Plasma
- ☐ CMMPS — Cobalamin, Methionine, and Methylmalonic Acid Pathways, Serum
- ☐ MMAGP — Methylmalonic Aciduria Gene Panel
- ☐ MPAGP — Methylmalonic Aciduria-Propionic Aciduria Combined Gene Panel
- ☐ MMAP — Methylmalonic Acid, Quantitative, Plasma
- ☐ MMAS — Methylmalonic Acid, Quantitative, Serum
- ☐ MMAU — Methylmalonic Acid, Quantitative, Urine

PEROXISOMAL BIOGENESIS & METABOLISM

- ☐ BAIPD — Bile Acids for Peroxisomal Disorders, Serum
- ☐ POXP — Fatty Acid Profile, Peroxisomal (C22-C26), Plasma
- ☐ POX — Fatty Acid Profile, Peroxisomal (C22-C26), Serum
- ☐ PDGP — Peroxisomal Disorder Gene Panel
- ☐ PIPA — Pipecolic Acid, Serum
- ☐ PIPU — Pipecolic Acid, Random, Urine
- ☐ PGRBC — Plasmalogens, Blood
- ☐ PGDBS — Plasmalogens, Blood Spot
- ☐ XALDZ — X-Linked Adrenoleukodystrophy, Full Gene Analysis

PORPHYRIAS

Urine

- ☐ ALAUR — Aminolevulinic Acid, Urine
- ☐ PBGU — Porphobilinogen, Quantitative, Random, Urine
- ☐ PQNU — Porphyrins, Quantitative, 24 Hour, Urine
- ☐ PQNRU — Porphyrins, Quantitative, Random, Urine

Plasma

- ☐ PBALP — Porphobilinogen and Aminolevulinic Acid, Plasma
- ☐ PTP — Porphyrins, Total, Plasma

Fecal

- ☐ FQPPS — Porphyrins, Feces

Blood

- ☐ PEWE — Porphyrins Evaluation, Washed Erythrocytes
- ☐ PEE — Porphyrins Evaluation, Whole Blood
- ☐ PPFWE — Protoporphyrins, Fractionation, Washed Erythrocytes
- ☐ PPFE — Protoporphyrins, Fractionation, Whole Blood

Enzymes

- ☐ PBGDW — Porphobilinogen Deaminase, Washed Erythrocytes
- ☐ PBGD_ — Porphobilinogen Deaminase, Whole Blood
- ☐ UPGC — Uroporphyrinogen III Synthase (Co-Synthase), Erythrocytes
- ☐ UPGDW — Uroporphyrinogen Decarboxylase, Washed Erythrocytes
- ☐ UPGD — Uroporphyrinogen Decarboxylase, Whole Blood

Molecular

- ☐ APGP — Acute Porphyria Gene Panel
- ☐ PCGP — Porphyria Comprehensive Gene Panel

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POSTMORTEM BIOCHEMICAL TESTING

☐	PMSBB	Postmortem Screening, Bile and Blood Spot

PURINE AND PYRIMIDINE METABOLISM

☐	PUPYP	Purine and Pyrimidine Panel, Plasma
☐	PUPYU	Purine and Pyrimidine Panel, Random, Urine
☐	SSCTU	S-Sulfocysteine Panel, Urine

SIALIC ACID DISORDERS

☐	SAU	Sialic Acid, Free and Total, Random, Urine

UREA CYCLE DISORDERS

☐	AAQP	Amino Acids, Quantitative, Plasma
☐	AAPD	Amino Acids, Quantitative, Random, Urine
☐	AAUCD	Amino Acids, Urea Cycle Disorders Panel, Plasma
☐	OAU	Organic Acids Screen, Random, Urine
☐	OROT	Orotic Acid, Random, Urine
☐	UCDP	Urea Cycle Disorders Gene Panel

WILSON DISEASE

☐	CERS	Ceruloplasmin, Serum
☐	CUU	Copper, 24 Hour, Urine
☐	CUS1	Copper, Serum
☐	WNDZ	Wilson Disease, *ATP7B* Full Gene Sequencing with Deletion/Duplication

WHOLE EXOME

☐	WESMT	Whole Exome and Mitochondrial Genome Sequencing
☐	WESDX	Whole Exome Sequencing for Hereditary Disorders
☐	WESR	Whole Exome Sequencing Reanalysis

WHOLE GENOME

☐	WGSDX	Whole Genome Sequencing for Hereditary Disorders
☐	WGSR	Whole Genome Sequencing Reanalysis

ADDITIONAL TESTS (INDICATE TEST ID AND NAME)

Print