



## Genetic Tests and Laboratory Matrix

The matrix below contains all the genetic tests which Evolent (formerly National Imaging Associates, Inc.) manages. This matrix is designed to assist in determining if a genetic test requires authorization through Evolent. Genetic tests are grouped by category and include the test name, the laboratory that performs the test, and the genetic testing unit (GTU). The GTU is a unique identifier created by Concert Genetics for each genetic test. To learn more about the GTU, please visit [Concert Genetics' website](#).

As Evolent systems are continually updated, a new matrix will be posted to RadMD the first of every month. To find the most current CPT codes please visit [Concert Genetics' Portal](#).

Prior authorization is not a guarantee of payment. Authorizations are based on medical necessity and are contingent upon member eligibility at the time services are rendered. Experimental and/or investigational (E/I) tests or categories are not considered medically necessary. Payment for these services will be excluded for all lines of business with the exception of Medicare, which is subject to state and federal regulations and guidelines.

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## Genetic Tests and Laboratory Matrix

### Arrhythmia Panel Tests

GTU	Test Name	Laboratory Name
<b>6S34G</b>	Arrhythmia / cardiac conduction defect	Fairview Diagnostic Laboratories
<b>7VV6G</b>	Arrhythmia Comprehensive Panel	The University of Chicago Genetic Services
<b>7VA6G</b>	Arrhythmia NGS Panel	AiLife Diagnostics
<b>7VA7G</b>	Arrhythmia NGS Panel Rapid	AiLife Diagnostics
<b>2YLUG</b>	Arrhythmia Panel	Blueprint Genetics
<b>6L93G</b>	Arrhythmia Panel	GeneDx
<b>7DMRG</b>	Arrhythmia Panel	Washington University in St. Louis Genomics and Pathology Services
<b>2EPVG</b>	Arrhythmia Panel NGS	All Children's Hospital
<b>76V3G</b>	Atrial Fibrillation (A Fib) Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>782SG</b>	Atrial fibrillation NGS Panel	Connective Tissue Gene Tests
<b>4JR5G</b>	Atrial Fibrillation NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>77VHG</b>	Atrial Fibrillation NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>76WLG</b>	Atrial Fibrillation NGS Panel (Sequencing Only)	Fulgent Genetics
<b>782TG</b>	Atrial fibrillation NGS Panel Next Generation Sequencing	Connective Tissue Gene Tests
<b>77ANG</b>	Atrial Fibrillation Panel	Blueprint Genetics
<b>77WVG</b>	Atrial fibrillation, familial	Fairview Diagnostic Laboratories
<b>6MEFG</b>	Atrioventricular block NGS Panel	Connective Tissue Gene Tests
<b>6KDGG</b>	Atrioventricular block NGS Panel Deletion / Duplication	Connective Tissue Gene Tests
<b>6ME7G</b>	Atrioventricular block NGS Panel Next Generation Sequencing	Connective Tissue Gene Tests
<b>5TZFG</b>	Cardiac Arrhythmia Panel	Northwest Clinical Genomics Lab
<b>6MEDG</b>	Cardiac channelopathy NGS Panel	Connective Tissue Gene Tests
<b>6MALG</b>	Cardiac channelopathy NGS Panel Deletion / Duplication	Connective Tissue Gene Tests
<b>6MECG</b>	Cardiac channelopathy NGS Panel Next Generation Sequencing	Connective Tissue Gene Tests
<b>53VJG</b>	Comp Arrhythmia Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>7SCDG</b>	Comprehensive Arrhythmia Gene Panel, Varies	Mayo Clinic Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2QALG</b>	Comprehensive Arrhythmia NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>25DJG</b>	Comprehensive Arrhythmia NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>3EQZG</b>	Comprehensive Arrhythmia NGS Panel (Sequencing Only)	Fulgent Genetics
<b>33XQG</b>	Comprehensive Arrhythmia Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>2Y7CG</b>	Comprehensive Arrhythmia Panel	PerkinElmer Genomics
<b>2LNNG</b>	Comprehensive Arrhythmias Panel	Knight Molecular Diagnostic
<b>5C8DG</b>	Comprehensive Cardiac Arrhythmia Panel	PreventionGenetics, part of Exact Sciences
<b>52PAG</b>	Familial Atrial Fibrillation – FAF (14 genes) Deletion/Duplication Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>2FPEG</b>	Familial Atrial Fibrillation – FAF (14 genes) NGS + Sanger fill-in Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>3BQFG</b>	Familial Atrial Fibrillation – FAF (14 genes) NGS Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>3LMAG</b>	Familial Atrial Fibrillation (FAF) Panel Sequencing	Indiana University Molecular Genetics Diagnostic Laboratory - Indiana University Medical Center, School of Medicine
<b>78FYG</b>	Familial Atrial Fibrillation Syndrome Panel	PreventionGenetics, part of Exact Sciences
<b>4ZNAG</b>	GeneSeq Cardio Familial Arrhythmia Panel	LabCorp
<b>4ZP3G</b>	GeneSeq : Cardio-Familial Arrhythmia Profile	Integrated Genetics
<b>2BXYG</b>	Hereditary Cardiac Arrhythmia (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>76Q4G</b>	Hereditary Ventricular Tachycardia Syndromes (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>25JTG</b>	Invitae Arrhythmia Comprehensive Panel	Invitae Corporation
<b>44YPG</b>	Invitae Arrhythmia Comprehensive Panel-Add-on Preliminary-evidence Genes for Arrhythmia	Invitae Corporation
<b>76S9G</b>	Invitae Arrhythmia Comprehensive Panel-Add-on Sudden Unexpected Death in Epilepsy (SUDEP) Genes	Invitae Corporation
<b>3E7RG</b>	Pan Arrhythmia (54 genes) Deletion/Duplication Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>2FQGG</b>	Pan Arrhythmia (54 genes) NGS + Sanger fill-in Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory

GTU	Test Name	Laboratory Name
<b>54N6G</b>	Pan Arrhythmia (54 genes) NGS Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>7UY5G</b>	Postmortem Arrhythmia Gene Panel, Tissue	Mayo Clinic Laboratories
<b>783JG</b>	Rest of Arrhythmia after Brugada Syndrome Panel	GeneDx
<b>775WG</b>	Rest of Arrhythmia after LQTS Panel	GeneDx
<b>6DDUG</b>	Rest of Arrhythmia after SCA Panel	GeneDx
<b>2Z2MG</b>	RhythmNext	Ambry Genetics
<b>58GWG</b>	SCA Arrhythmia Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>7B7BG</b>	STAT Comprehensive Arrhythmia Panel	PerkinElmer Genomics
<b>6DTGG</b>	Sudden Cardiac Arrest Arrhythmia Panel	GeneDx
<b>77GKG</b>	Sudden Cardiac Arrest Panel	PreventionGenetics, part of Exact Sciences
<b>4AAWG</b>	Ventricular fibrillation, familial	Fairview Diagnostic Laboratories

#### Autism Spectrum Disorder/Intellectual Disability Panel Tests <sup>1</sup>

GTU	Test Name	Laboratory Name
<b>4J42G</b>	Acetyl-CoA carboxylase deficiency	Fairview Diagnostic Laboratories
<b>77SSG</b>	Angelman Syndrome Gene Analysis	University of Minnesota Physicians Outreach Laboratory
<b>78EAG</b>	Angelman-like syndrome panel	Fairview Diagnostic Laboratories
<b>2FHGG</b>	Autism NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>76VUG</b>	Autism NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>78AYG</b>	Autism NGS Panel (Sequencing Only)	Fulgent Genetics
<b>76Z5G</b>	Autism panel	Fairview Diagnostic Laboratories
<b>78FSG</b>	Autism Spectrum Disorders (ASD) Panel	PreventionGenetics, part of Exact Sciences
<b>76U3G</b>	Autism Spectrum Disorders Panel	Blueprint Genetics
<b>7BM2G</b>	AUTISM, INTELLECTUAL DISABILITY, and DEVELOPMENTAL DELAY GENE SEQUENCING PANEL	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>76QWG</b>	AUTISM/AUTISM SPECTRUM DISORDER (53 gene panel)	Center for Human Genetics, Inc.
<b>77KNG</b>	Autism/ID Panel	GeneDx
<b>78K3G</b>	Autism/ID Xpanded Panel	GeneDx
<b>77YKG</b>	AutismNext ®	Ambry Genetics

GTU	Test Name	Laboratory Name
<b>78FHG</b>	Autosomal Recessive Non-Specific Intellectual Disability Panel	The University of Chicago Genetic Services
<b>7RW9G</b>	CHILD Neurodevelopmental Panel	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>77SAG</b>	Comprehensive Intellectual Disability / Autism (NGS Panel and Copy Number Analysis + mtDNA + Fragile X Repeat Expansion and Methylation)	MNG Laboratories
<b>77AUG</b>	Comprehensive Intellectual Disability / Autism (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
<b>5QHHG</b>	Feingold syndrome	Fairview Diagnostic Laboratories
<b>787KG</b>	Focused Autism and Intellectual Disability Panel	PerkinElmer Genomics
<b>77WLG</b>	Glycosylphosphatidylinositol biosynthesis defect	Fairview Diagnostic Laboratories
<b>3BXSG</b>	Hyperphosphatasia with mental retardation syndrome	Fairview Diagnostic Laboratories
<b>5HM6G</b>	Hypomagnesemia, seizures, and mental retardation	Fairview Diagnostic Laboratories
<b>76YDG</b>	Inherited Glycosylphosphatidylinositol Biosynthesis Defects (IGDs) Panel	PreventionGenetics, part of Exact Sciences
<b>76YRG</b>	Intellectual disability	Fairview Diagnostic Laboratories
<b>5FW2G</b>	Intellectual Disability Exome	The University of Chicago Genetic Services
<b>5DPWG</b>	Intellectual Disability NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>78F2G</b>	Intellectual Disability NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77GSG</b>	Intellectual Disability NGS Panel (Sequencing Only)	Fulgent Genetics
<b>79Q6G</b>	Intellectual disability panel - NGS Panel (CNV included)	Centogene
<b>775YG</b>	Intellectual Disability, Epilepsy, and Autism (IDEA) Panel - Family - Duo (IDEA panel of patient + 1 additional family members)	PreventionGenetics, part of Exact Sciences
<b>783YG</b>	Intellectual Disability, Epilepsy, and Autism (IDEA) Panel - Family - Trio (IDEA panel of patient + 2 additional family members)	PreventionGenetics, part of Exact Sciences
<b>77KFG</b>	Intellectual Disability, Epilepsy, and Autism (IDEA) Panel - Patient Only	PreventionGenetics, part of Exact Sciences
<b>76Z2G</b>	Intellectual disability, autosomal dominant	Fairview Diagnostic Laboratories
<b>78E7G</b>	Intellectual disability, autosomal recessive	Fairview Diagnostic Laboratories
<b>7CFRG</b>	Invitae Neurodevelopmental Disorders (NDD) Panel	Invitae Corporation
<b>4AADG</b>	Kleefstra syndrome	Fairview Diagnostic Laboratories
<b>4J4AG</b>	Martsolf syndrome	Fairview Diagnostic Laboratories
<b>77HZG</b>	NeurodevelopmentNext™	Ambry Genetics

GTU	Test Name	Laboratory Name
77H2G	Non-Specific Intellectual Disability Panel	The University of Chicago Genetic Services
76UQG	Nonsyndromic Intellectual Disability (NGS Panel and Copy Number Analysis)	MNG Laboratories
4E6VG	Opitz G/BBB Syndrome Panel	PreventionGenetics, part of Exact Sciences
7BYUG	PGmaxTM - Intellectual Disability, Epilepsy, and Autism (IDEA) Panel	PreventionGenetics, part of Exact Sciences
7768G	PGXome Custom - Intellectual Disability, Autosomal Dominant	PreventionGenetics, part of Exact Sciences
77XDG	PGXome Custom - Intellectual Disability, Autosomal Recessive	PreventionGenetics, part of Exact Sciences
7B62G	STAT Focused Autism and Intellectual Disability Panel	PerkinElmer Genomics
77EPG	Syndromic Autism NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
77SFG	Syndromic Intellectual Disability (NGS Panel and Copy Number Analysis)	MNG Laboratories
78FXG	Top 99 Genetic Causes of Developmental Delay Panel	PreventionGenetics, part of Exact Sciences

#### Bladder Cancer Diagnostic and Recurrence Algorithmic Tests <sup>1</sup>

GTU	Test Name	Laboratory Name
73PCG	Cxbladder Detect	Pacific Edge
7VLUG	Cxbladder Detect	Pacific Edge
6W4EG	Cxbladder Monitor	Pacific Edge
72KHG	Cxbladder Triage	Pacific Edge
7BN4G	Decipher® Bladder Genomic Test	Veracyte

#### Blood-based Post Heart Transplant Gene Expression Panels for Rejection Risk

GTU	Test Name	Laboratory Name
74CFG	AlloMap Heart	Allina Health Laboratory
2TW7G	AlloMap Molecular Expression, Heart	University of Minnesota Physicians Outreach Laboratory
525DG	AlloMapHeart	CareDx, Inc.

## BRCA1/2 Sequencing & Deletion/Duplication Tests

GTU	Test Name	Laboratory Name
<b>7SYQG</b>	BRAC ANALYSIS,COMPREHENSIVE	University of California Davis Health System - UCDMC - Molecular and Cytogenetic Laboratory
<b>6RPGG</b>	BRACAnalysis CDx	Myriad Genetics
<b>2FFUG</b>	BRCA AVANTAGE,COMPREHENSIVE	Empire City Laboratories, Inc.
<b>3GPZG</b>	BRCA Comprehensive (BRCA 1 and BRCA 2 Seq & Del/Dup Analysis)	Genetics Center - Molecular and Cytogenetic Diagnostic Laboratories
<b>26REG</b>	BRCA Panel (BRCA1, BRCA2)	Palo Verde Laboratory - division of Sonora Quest Lab
<b>2E63G</b>	BRCA Panel (BRCA1, BRCA2)	Quest Diagnostics
<b>7V2JG</b>	BRCA1 & BRCA2 Panel	Baylor Genetics, LLC
<b>55MGG</b>	BRCA1 AND BRCA2 ANALYSIS (SEQUENCING & MLPA FOR BOTH GENES)	Center for Human Genetics, Inc.
<b>5BRJG</b>	BRCA1 and BRCA2 Focus (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>6U9AG</b>	BRCA1 and BRCA2 Focus Panel (Germline)	NeoGenomics Laboratories
<b>2TT6G</b>	BRCA1 and BRCA2 Hereditary Gene Analysis Deletion and Duplication	University of Minnesota Physicians Outreach Laboratory
<b>7U27G</b>	BRCA1 and BRCA2 Sequencing and Del/Dup (NGS)	University of Michigan - Michigan Medical Genetics Laboratories
<b>2ELCG</b>	BRCA1 and BRCA2 Sequencing and Deletion/Duplication	Knight Molecular Diagnostic
<b>7QGJG</b>	BRCA1 AND BRCA2, COMPREHENSIVE	Clinical Pathology Laboratories
<b>6UQAG</b>	BRCA1 and BRCA2-Associated HBOC Syndrome Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>6LTEG</b>	BRCA1&2 Analysis	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>79AHG</b>	BRCA1, BRCA2 panel Combi (with MLPA) - NGS Panel (MLPA included)	Centogene
<b>79AJG</b>	BRCA1, BRCA2 panel Plus - NGS Panel (CNV included)	Centogene
<b>2YYZG</b>	BRCA1/2 seq and del/dup	Ambry Genetics
<b>4RLXG</b>	BRCA1/2 Sequencing and Del/Dup Analysis	BioReference Laboratories
<b>6DTQG</b>	BRCA1/2 Sequencing and Del/Dup Analysis	GeneDx
<b>3D8PG</b>	BRCA1/2 Sequencing and Deletion Duplication Analysis	ACL Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>5FUUG</b>	BRCA1/2: Comprehensive BRCA Analysis by Gene Sequencing with Deletion/Duplication Analysis	Medical Diagnostic Laboratories, LLC
<b>7DLBG</b>	BRCA1/BRCA2 Genes, Full Gene Analysis, Varies	Mayo Clinic Laboratories
<b>6LS6G</b>	BRCAssure : BRCA1 and BRCA2 Comprehensive Analysis	LabCorp
<b>546VG</b>	BRCAssure: BRCA1 / 2 Comprehensive Analysis	Labcorp   Oncology
<b>58G6G</b>	BRCAssure: Comprehensive BRCA1 / 2 Analysis	Integrated Genetics
<b>2C88G</b>	Empower - BRCA1 & BRCA2 Hereditary Cancer Test	Natera
<b>57DKG</b>	Hereditary Breast and Ovarian Cancer BRCA1/2 Panel	PreventionGenetics, part of Exact Sciences
<b>32SZG</b>	Hereditary Breast and Ovarian Cancer Syndrome Panel	PerkinElmer Genomics
<b>2V8RG</b>	Integrated BRAC Analysis, EDTA Whole Blood (BRACA 1 & 2 Testing)	Marshfield Labs
<b>6RMHG</b>	Integrated BRACAnalysis	Myriad Genetics
<b>7QDVG</b>	Integrated Reflex BRACAnalysis	Myriad Genetics
<b>442GG</b>	Invitae BRCA1 and BRCA2 Panel	Invitae Corporation
<b>44N2G</b>	Invitae BRCA1 and BRCA2 STAT Panel	Invitae Corporation
<b>7SLMG</b>	PrevenTest, Custom (BRCA1, BRCA2)	Advanced Molecular Diagnostics, LLC
<b>6RN9G</b>	Reflex BRACAnalysis	Myriad Genetics
<b>7B7QG</b>	STAT Hereditary Breast and Ovarian Cancer Syndrome Panel	PerkinElmer Genomics

### BRCA1/2 Sequencing Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7TMWG</b>	BRCA HRR Deficiency NGS Panel with Fusion	University of Minnesota Physicians Outreach Laboratory
<b>7QGKG</b>	BRCA1 and BRCA2 Sequencing	Clinical Pathology Laboratories
<b>7U26G</b>	BRCA1 and BRCA2 Sequencing (NGS)	University of Michigan - Michigan Medical Genetics Laboratories
<b>79AFG</b>	BRCA1, BRCA2 panel - NGS Panel	Centogene
<b>3SEYG</b>	BRCA1/2 Sequencing Test	Gene by Gene
<b>6UQTG</b>	BRCA1-2 with RNAinsight	Ambry Genetics
<b>2CHKG</b>	Hereditary cancer BRCA1 BRCA2	Fairview Diagnostic Laboratories

## Breast Cancer Prognostic Algorithmic Tests

GTU	Test Name	Laboratory Name
<b>4K7JG</b>	Agendia Breast Cancer Test Suite	Agendia, Inc.
<b>4ZMDG</b>	Breast Cancer Prognostic Gene Signature Assay (Prosigna ), IVD	LabCorp
<b>6UVJG</b>	EndoPredict	Myriad Genetics
<b>49Z2G</b>	MammaPrint	Agendia, Inc.
<b>326SG</b>	PROSIGNA BREAST CANCER PROGNOSTIC GENE SIGNATURE	Empire City Laboratories, Inc.
<b>54HTG</b>	Prosigna Breast Cancer Prognostic Gene Signature Assay	Labcorp   Oncology
<b>6UUCG</b>	Prosigna Breast Cancer Prognostic Gene Signature Assay	Quest Diagnostics
<b>2KTBG</b>	Prosigna Breast Cancer Prognostic Gene Signature Assay	Veracyte

## Breast Cancer Treatment and Prognostic Algorithmic Tests

GTU	Test Name	Laboratory Name
<b>2VDEG</b>	Oncotype Dx Breast Cancer Assay	2VDEG
<b>7BNGG</b>	Oncotype DX Breast Recurrence Score Test	7BNGG

## Cell-Free Circulating Tumor DNA Colorectal Cancer Panel Tests

GTU	Test Name	Laboratory Name
<b>2AHTG</b>	OncoBEAM™ CRC1: KRAS, NRAS, BRAF, HRAS	Sysmex Inostics, INC

## Cell-Free Circulating Tumor DNA Lung Cancer Panel Tests

GTU	Test Name	Laboratory Name
<b>7SGYG</b>	CNSide Biomarker Profile, Lung	Biocept
<b>7ACFG</b>	GeneStrat	Biodesix
<b>7ACGG</b>	GeneStrat NGS	Biodesix
<b>4764G</b>	Genestrat Test	ACL Laboratories
<b>7TRFG</b>	InVisionFirst -Lung Liquid Biopsy	Inivata Ltd
<b>4FGPG</b>	InVisionFirst -Lung Liquid Biopsy	NeoGenomics Laboratories
<b>7RVXG</b>	IQlung Treatment Guidance Testing	Biodesix

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>28XAG</b>	OncoBEAM™ Lung2: EGFR, KRAS, BRAF	Sysmex Inostics, INC
<b>6UU4G</b>	Resolution ctDx Lung assay (Liquid Biopsy)	Labcorp   Oncology

#### Cell-Free Circulating Tumor DNA Cancer Profiling Panel Tests (51 or more genes)

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7VVWG</b>	Caris Assure	Caris Life Sciences
<b>6R3DG</b>	FoundationOne Liquid CDx	Foundation Medicine
<b>7BZFG</b>	FoundationOne Liquid CDx (For Allina outpatient use only)	Allina Health Laboratory
<b>27X9G</b>	Guardant 360 Tumor Sequencing Assay	University of Minnesota Physicians Outreach Laboratory
<b>7ABJG</b>	Guardant360	Guardant Health
<b>6UW6G</b>	Guardant360 CDx	Guardant Health
<b>7TMMG</b>	LiquidHALLMARK	Lucence Health
<b>7ACKG</b>	MSK-ACCESS	Memorial Sloan Kettering Cancer Center
<b>7VPQG</b>	Northstar Select	Billion to One
<b>7TXKG</b>	Tempus xF Liquid Biopsy Test, blood	University of Michigan - Michigan Medical Genetics Laboratories
<b>2B64G</b>	Tempus  xF: Liquid Biopsy Panel of 105 Genes	Tempus AI, Inc.
<b>7S8WG</b>	Tempus  xF+: Liquid Biopsy Panel of 523 Genes	Tempus AI, Inc.

#### Cell-Free Circulating Tumor DNA Cancer Profiling Panel Tests (5-50 genes)

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7RWNG</b>	MayoComplete Liquid Biopsy Panel, Next-Generation Sequencing, Cell-Free DNA	Mayo Clinic Laboratories
<b>3FYGG</b>	NeoLAB Solid Tumor Liquid Biopsy	NeoGenomics Laboratories
<b>7VQXG</b>	Neuroblastoma Liquid Biopsy Panel (Circulating Tumor DNA)	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>537NG</b>	PGDx elio plasma resolve*	Personal Genome Diagnostics
<b>7UQXG</b>	Plasma Focus	LabCorp

## Chromosomal Microarray (SNP and CGH) for Invasive Prenatal Tests

GTU	Test Name	Laboratory Name
<b>2KSUG</b>	511810 Follow-Up Prenatal CMA qPCR	LabCorp
<b>79AMG</b>	CentoArrayCyto 750K - Array CGH (prenatal)	Centogene
<b>79APG</b>	CentoArrayCyto HD - Array CGH (prenatal)	Centogene
<b>2EKCG</b>	Chromosomal Microarray - Prenatal Diagnosis	Knight Molecular Diagnostic
<b>2LBVG</b>	Chromosomal Microarray Analysis - Prenatal CMA Amniotic Fluid (Affymetrix CytoScan HD array)	Washington University in St. Louis Pathology Services
<b>2QYYG</b>	Chromosomal Microarray Analysis - Prenatal CMA Chorionic Villi Sampling (Affymetrix CytoScan HD array)	Washington University in St. Louis Pathology Services
<b>33X4G</b>	Chromosomal Microarray Prenatal (CMAP)	Marshfield Labs
<b>2ETFG</b>	Chromosomal Microarray, Preatal, ClariSure	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>6LEBG</b>	Chromosomal Microarray, Prenatal	Hennepin County Medical Center
<b>6R47G</b>	Chromosomal Microarray, Prenatal, Amniotic Fluid/Chorionic Villus Sampling	Mayo Clinic Laboratories
<b>7U36G</b>	Chromosomal Microarray, Prenatal, Amniotic Fluid/Chorionic Villus Sampling	University of Michigan - Michigan Medical Genetics Laboratories
<b>2EQ3G</b>	Chromosomal Microarray, Prenatal, ClariSure Oligo-SNP	Quest Diagnostics
<b>2WTCG</b>	CHROMOSOMAL MICROARRAY, PRENATAL CLARISURE	Empire City Laboratories, Inc.
<b>7U5HG</b>	Cytogenomic SNP Microarray&nbsp;-&nbsp;Fetal	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>25DWG</b>	Cytogenomic SNP Microarray, Fetal	ARUP Laboratories
<b>4XSNG</b>	Cytogenomic SNP Microarray, Fetal	TriCore Reference Laboratories
<b>3DQXG</b>	Cytogenomic SNP Microarray, Fetal	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<b>78KYG</b>	Cytogenomic SNP Microarray, Fetal	University of California Davis Health System - UCDMC - Molecular and Cytogenetic Laboratory
<b>2Z2KG</b>	Cytogenomic SNP Microarray, Fetal	University of Minnesota Physicians Outreach Laboratory
<b>2P6UG</b>	Expanded Chromosomal Microarray Analysis - Prenatal - Amniotic Fluid	Baylor Genetics, LLC
<b>4JAVG</b>	Expanded Chromosomal Microarray Analysis - Prenatal - CVS	Baylor Genetics, LLC
<b>5BR7G</b>	Genomic Microarray, Prenatal (Amniotic Fluid or CVS)	ACL Laboratories
<b>7V33G</b>	IriSight™? CNV AnalysisAccepted Specimens	Variantyx

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>328BG</b>	Prenatal Cytogenomic Microarray	University Hospitals
<b>2LEXG</b>	Prenatal Microarray	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>2EE3G</b>	Prenatal Microarray Analysis with Parental Testing	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>5BNTG</b>	Prenatal Microarray with 5-Cell Chromosome Analysis (No Parental Testing)	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>2E8KG</b>	Prenatal Microarray without Parental Testing	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>2EMWG</b>	Prenatal SNP Microarray	Colorado Genetics Laboratory
<b>7C4VG</b>	Prenatal Whole Genome Chromosomal Microarray	BioReference Laboratories
<b>6WNXG</b>	Prenatal Whole Genome Chromosomal Microarray	GeneDx
<b>6LUQG</b>	Rapid Prenatal Chromosomal Microarray via aCGH and SNP - Prenatal Test	PreventionGenetics, part of Exact Sciences
<b>7C4XG</b>	Reflex to Prenatal Whole Genome Chromosomal Microarray from Chromosomes	BioReference Laboratories
<b>5BNSG</b>	Reveal SNP Microarray – Prenatal	Integrated Genetics
<b>2EPWG</b>	SNP Array for Prenatal Analysis (aka Microarray)	UCSF Molecular Diagnostics Laboratory
<b>3U6CG</b>	SNP Microarray (Direct)-Prenatal (Reveal)	LabCorp
<b>3N8VG</b>	SNP Microarray-Prenatal (Reveal)	LabCorp

### Chromosomal Microarray (SNP and CGH) for Pregnancy Loss Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6S7ZG</b>	Anora Miscarriage Test (Fresh)	Natera
<b>2EXPG</b>	Anora Miscarriage Test (Paraffin)	Natera
<b>2EW5G</b>	CGH-POC: Comparative Genomic Hybridization	New Jersey Medical School - Institute of Genomic Medicine
<b>2EDZG</b>	Chromosomal Microarray Analysis - CMA Products of Conception (POC) (Affymetrix CytoScan HD array)	Washington University in St. Louis Pathology Services
<b>6LRAG</b>	Chromosomal Microarray Analysis (CMA) - Products of Conception	Allina Health Laboratory
<b>6R2VG</b>	Chromosomal Microarray, Autopsy, Products of Conception, or Stillbirth	Mayo Clinic Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7U37G</b>	Chromosomal Microarray, Autopsy, Products of Conception, or Stillbirth	University of Michigan - Michigan Medical Genetics Laboratories
<b>6R3NG</b>	Chromosomal Microarray, Autopsy/Products of Conception/Stillbirth, Tissue	Mayo Clinic Laboratories
<b>7U35G</b>	Chromosomal Microarray, Autopsy/Products of Conception/Stillbirth, Tissue	University of Michigan - Michigan Medical Genetics Laboratories
<b>2E8MG</b>	Chromosomal Microarray, POC, ClariSure Oligo-SNP	Quest Diagnostics
<b>3PJJG</b>	CHROMOSOMAL MICROARRAY, POC, CLARISURE OLIGO-SNP,POC	Empire City Laboratories, Inc.
<b>3KVVG</b>	Comparative Genomic Hybridization (CGH): Products of Conception (POC)	New Jersey Medical School - Institute of Genomic Medicine
<b>7DK8G</b>	Cytogenomic Molecular Inversion Probe Array FFPE Tissue - Products of Conception	ARUP Laboratories
<b>2EKBG</b>	Genomic Microarray, POC	ACL Laboratories
<b>2EXDG</b>	Genomic SNP Microarray, Products of Conception	ARUP Laboratories
<b>7PTBG</b>	Microarray Analysis - PREGNANCY LOSS	Nebraska Medical Center - Molecular Diagnostic Laboratory
<b>2ENDG</b>	Microarray-Products of Conception (POC) Reveal FFPE	LabCorp
<b>2E65G</b>	Microarray-Products of Conception (POC) Reveal FFPE, Data Transfer	LabCorp
<b>5BMSG</b>	POC (Products of Conception) / Tissue Microarray Analysis	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>5BN2G</b>	POC Microarray with 5-Cell Chromosome Analysis	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>2M7XG</b>	Pregnancy Loss Microarray	Colorado Genetics Laboratory
<b>3JXFG</b>	Products of Conception Microarray + SNP	Cleveland Clinic Laboratories
<b>4ZPRG</b>	Reveal SNP Microarray – POC	Integrated Genetics
<b>5FU7G</b>	SNP Array for Tissue and POC (aka Microarray)	UCSF Molecular Diagnostics Laboratory
<b>34WFG</b>	SNP Microarray – Products of Conception	Knight Molecular Diagnostic
<b>3U6BG</b>	SNP Microarray Products of Conception (POC) / Tissue (Reveal )	LabCorp

## Chromosomal Microarray (SNP and CGH) Tests

GTU	Test Name	Laboratory Name
<b>25SKG</b>	2.7 M SNP Chromosomal Microarray	Nicklaus Children's Hospital - Miami Genetic Laboratories
<b>4LCPG</b>	Array CGH	UAB Cytogenetics Laboratory
<b>4ZQCG</b>	Array CGH Constitutional blood (Microarray)	PathGroup
<b>3R4MG</b>	Array Comparative Genomic Hybridization	Children's Hospital and Research Center Oakland - Molecular Genetics
<b>2KYRG</b>	Array-based Comparative Genome Hybridization, Genetic Dx.	Stanford Clinical Laboratories - Biochemical Genetics Laboratory
<b>3GPJG</b>	Assure SNP Microarray Analysis (FDA cleared)	CytoGenX
<b>2E8LG</b>	Assure SNP Microarray Analysis (FDA cleared) – Prenatal & Postnatal	CytoGenX
<b>3AG7G</b>	CGH: Comparative Genomic Hybridization	New Jersey Medical School - Institute of Genomic Medicine
<b>3F5GG</b>	CHROMOSOMAL MICROARRAY	Detroit Medical Center University Laboratories - Molecular Genetics Diagnostic Laboratory
<b>4CURG</b>	CHROMOSOMAL MICROARRAY	Fullerton Genetics Center - Mission Health
<b>5JPRG</b>	Chromosomal Microarray - Postnatal	Knight Molecular Diagnostic
<b>6XNCG</b>	Chromosomal Microarray (MicroarrayDx)	GeneDx
<b>687XG</b>	Chromosomal Microarray Analysis	Center for Genetic Testing at Saint Francis
<b>38VKG</b>	Chromosomal Microarray Analysis	UCLA Diagnostic Molecular Pathology Laboratory
<b>3UXBG</b>	Chromosomal Microarray Analysis - CMA Fibroblasts (Affymetrix CytoScan HD array)	Washington University in St. Louis Pathology Services
<b>3HN3G</b>	Chromosomal Microarray Analysis - CMA Peripheral Blood (Affymetrix CytoScan HD array)	Washington University in St. Louis Pathology Services
<b>6NL2G</b>	Chromosomal Microarray Analysis - HR	Baylor Genetics, LLC
<b>2ZAVG</b>	Chromosomal Microarray Analysis - HR + SNP Screen (Comprehensive)	Baylor Genetics, LLC
<b>6LRBG</b>	Chromosomal microarray analysis (CMA) - Blood	Allina Health Laboratory
<b>4DNFG</b>	Chromosomal Microarray Analysis for Constitutional Abnormalities	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>2EW7G</b>	Chromosomal Microarray only	Washington University in St. Louis Pathology Services
<b>53JEG</b>	Chromosomal Microarray SNP, Constitutional	Cleveland Clinic Laboratories
<b>3A53G</b>	Chromosomal Microarray with abbreviated karyotype	Washington University in St. Louis Pathology Services
<b>32EDG</b>	Chromosomal Microarray with concurrent karyotype	Washington University in St. Louis Pathology Services

GTU	Test Name	Laboratory Name
<b>333GG</b>	Chromosomal Microarray with reflex karyotype	Washington University in St. Louis Pathology Services
<b>4XSMG</b>	Chromosomal Microarray, Congenital, Bld (CMACB)	Marshfield Labs
<b>6R2TG</b>	Chromosomal Microarray, Congenital, Blood	Mayo Clinic Laboratories
<b>7R9LG</b>	Chromosomal Microarray, Hematologic Malignancy, ClariSure Oligo-SNP (90961)	Rady Children's Institute for Genomic Medicine
<b>2Z55G</b>	Chromosomal Microarray, POC FFPE, ClariSure Oligo-SNP	Quest Diagnostics
<b>8EJVG</b>	Chromosomal Microarray, POC, ClariSure Oligo-SNP	Medfusion
<b>8EPBG</b>	Chromosomal Microarray, Postnatal Familial Follow-up, ClariSure Oligo-SNP	Quest Diagnostics
<b>2ETDG</b>	Chromosomal Microarray, Postnatal, ClariSure Oligo-SNP	Quest Diagnostics
<b>2E8NG</b>	Chromosomal Microarray, Postnatal, ClariSure Oligo-SNP (16478)	Rady Children's Institute for Genomic Medicine
<b>2EP9G</b>	Chromosomal SNP Microarray	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>2BG9G</b>	Chromosomal SNP Microarray	Seattle Children's Hospital
<b>68Y6G</b>	Chromosome Array (aCGH)	Center for Genetic Testing at Saint Francis
<b>2DXNG</b>	Chromosome Microarray	Palo Verde Laboratory - division of Sonora Quest Lab
<b>53TQG</b>	Chromosome Microarray	University of Virginia Health System
<b>54CPG</b>	Chromosome Microarray Analysis	Molecular Diagnosis - University of Rochester Medical Center - Strong Memorial Hospital
<b>2ELZG</b>	Chromosome Microarray Analysis - Constitutional	All Children's Hospital
<b>7S97G</b>	Chromosome Microarray, Congenital	Sanford USD Medical Center - Sanford Clinic USD Genetics Laboratory
<b>2DYKG</b>	CHROMOSOME MICROARRAY, POSTNATAL	Empire City Laboratories, Inc.
<b>7QHSG</b>	CHROMOSOME SNP MICROARRAY	University of Texas Medical Branch - UTMB - Porphyria Laboratory
<b>5FQBG</b>	Chromosome Specific Interphase FISH	Center for Genetic Testing at Saint Francis
<b>2LLCG</b>	Combitmatrix Combi SNP Array Tissue	Norton CPA Lab
<b>2FNJG</b>	Compar Genomic Hybrid-SNP	Norton CPA Lab
<b>2EPKG</b>	Comparative Genomic Hybridization (CGH):Mental Retardation/ Developmental Delay	New Jersey Medical School - Institute of Genomic Medicine
<b>5DPRG</b>	Constitutional Chromosomal Microarray (Copy Number)	University of Minnesota Physicians Outreach Laboratory
<b>6T6QG</b>	Constitutional Chromosomal Microarray Analysis – Peripheral Blood	Indiana University Molecular Genetics Diagnostic Laboratory - Indiana University Medical Center, School of Medicine
<b>2E7LG</b>	Constitutional Chromosomal Microarray Analysis (CMA)	Indiana University School of Medicine - Cytogenetics Laboratory

GTU	Test Name	Laboratory Name
7SUJG	Constitutional Limited Chromosomal Microarray (Copy number only) (Charged)	University of Minnesota Physicians Outreach Laboratory
2Z48G	Constitutional or Products of Conception (POC) Chromosomal Microarray (Copy Number/SNP)	University of Minnesota Physicians Outreach Laboratory
2ETHG	Constitutional SNP Array Karyotyping, Constitutional Chromosome Microarray Analysis (CMA)	Beaumont Laboratories - Molecular Pathology Lab
2M4WG	Cytogenomic Microarray Analysis	University of Washington Medicine - Pathology - Cytogenetics and Genomics Laboratory
52XTG	Cytogenomic Microarray Analysis	Veripath Laboratories
49JCG	Cytogenomic Microarray Analysis of Postnatal Blood	Akron Children's Hospital
49W6G	Cytogenomic SNP array (postnatal)	The University of Chicago Genetic Services
2E3BG	Cytogenomic SNP Microarray	ARUP Laboratories
8EBHG	Cytogenomic SNP Microarray	Medfusion
2MTFG	Cytogenomic SNP Microarray	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
2X3MG	Cytogenomic SNP Microarray	University of California Davis Health System - UCDMC - Molecular and Cytogenetic Laboratory
7U9JG	Cytogenomic SNP Microarray	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
2542G	Cytogenomic SNP Microarray Buccal Swab	ARUP Laboratories
4PPTG	Exon-Centric Deletion/Duplication Anaylsis	Knight Molecular Diagnostic
2D5QG	FirstStepDx PLUS	Bionano Laboratories
7UHHG	Genomic Microarray Analysis - Parental study, Genome Wide Array	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
3MA8G	Genomic Microarray, Blood	ACL Laboratories
2ENHG	Genomic SNP Microarray, Products of Conception	University of California Davis Health System - UCDMC - Molecular and Cytogenetic Laboratory
7UHKG	Genomic SNP Microarray, Products of Conception	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
2ETGG	Illumina Global Screening Array Sequencing Test	Gene by Gene
2EQNG	Illumina Microarray Analysis	University of Wisconsin - Madison WSLH - UW Cytogenetic Services
2EMZG	Illumina Microarray Analysis- Targeted Family	University of Wisconsin - Madison WSLH - UW Cytogenetic Services
7TG6G	Invitae Chromosomal Microarray Analysis (CMA)	Invitae Corporation

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2EMYG</b>	Invitae Chromosomal Microarray Analysis (CMA) with follow-up FISH when required	Invitae Corporation
<b>7QHGG</b>	LV CYTOGENOMIC SNP MICROARRAY	Clinical Pathology Laboratories
<b>3UUJG</b>	Micro Array: Comparative Genomic Hybridization (aCGH) with SNP	University of Florida - Health Pathology Laboratories
<b>346KG</b>	MICROARRAY : FAMILY STUDY	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>33ZTG</b>	Microarray : SNP	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>3CAGG</b>	Microarray Analysis	University of Illinois at Chicago - Biochemical Genetics Laboratory
<b>3LQLG</b>	Microarray Analysis - CytoScan SNP	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<b>7PTDG</b>	Microarray Analysis - MALIGNANCY TESTING	Nebraska Medical Center - Molecular Diagnostic Laboratory
<b>7PTAG</b>	Microarray Analysis - POSTNATAL	Nebraska Medical Center - Molecular Diagnostic Laboratory
<b>6LCFG</b>	Microarray Analysis With Interpretation	Hennepin County Medical Center
<b>32YYG</b>	MICROARRAY CGH, WHOLE GENOME(OLIGO)	Ann and Robert Lurie Children's Hospital of Chicago
<b>54P3G</b>	Microarray Single Nucleotide Polymorphism	Molecular Diagnosis - University of Rochester Medical Center - Strong Memorial Hospital
<b>4QK2G</b>	Microarray/aCGH	Genetic Associates
<b>2DYJG</b>	Pediatric and Adult (Postnatal) SNP Microarray	Colorado Genetics Laboratory
<b>7A2PG</b>	POC Microarray Analysis	ProPath
<b>6NNJG</b>	Rapid Chromosomal Microarray via aCGH and SNP Test	PreventionGenetics, part of Exact Sciences
<b>59WPG</b>	Rapid microarray (CGH and SNP)	Allele Diagnostics
<b>3PL6G</b>	Reflex microarray if chromosomes are normal	Genetic Associates
<b>2EKFG</b>	Reflex to SNP Array	UCSF Molecular Diagnostics Laboratory
<b>2DYMIG</b>	Reveal SNP Microarray Pediatric	Integrated Genetics
<b>2EN4G</b>	SNP Array	Ambry Genetics
<b>5FU8G</b>	SNP Array for Blood Analysis (aka Microarray)	UCSF Molecular Diagnostics Laboratory
<b>2DYGG</b>	SNP Array, Family Follow-Up	UCSF Molecular Diagnostics Laboratory
<b>54CTG</b>	SNP CHIP (6.0) WHOLE GENOME COPY NUMBER ANALYSIS	Center for Human Genetics, Inc.
<b>3GSDG</b>	SNP LOH STUDIES	Center for Human Genetics, Inc.
<b>2L3GG</b>	SNP Microarray	Shodair Children's Hospital - Genetics Laboratory

GTU	Test Name	Laboratory Name
2ZALG	SNP Microarray Analysis (Chromosomal Microarray)	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
2Z5JG	SNP Microarray Pediatric	Integrated Regional Laboratories
2EKEG	SNP microarray Processing, Extraction and Storage	UCSF Molecular Diagnostics Laboratory
49AAG	SNP Microarray when Routine Chrom or High Res and Fragile X have ALREADY been billed	Center for Human Genetics, Inc.
2EQQG	SNP Microarray-Pediatric (Reveal)	LabCorp
52MEG	SNP Oligonucleotide Microarray Analysis (SOMA)	Columbia University - Personalized Genomic Medicine
6NNKG	Whole-Genome Chromosomal Microarray (CMA-ISCA) via the aCGH and SNP Test	PreventionGenetics, part of Exact Sciences

### Colorectal Cancer Prognostic Algorithmic Tests <sup>1</sup>

GTU	Test Name	Laboratory Name
7S9GG	miR-31now	GoPath Laboratories
7BNJG	Oncotype DX® Colon Recurrence Score Test	Exact Sciences Laboratories, LLC

### Comprehensive Arrhythmia & Cardiomyopathy (Sudden Cardiac or Unexplained Death) Panel Tests

GTU	Test Name	Laboratory Name
7VV4G	Arrhythmia and Cardiomyopathy Comprehensive Panel	The University of Chicago Genetic Services
23DQG	Cardiomyopathy and Arrhythmia Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
2DTLG	Cardiomyopathy and Arrhythmia Sequencing Panel	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
2LRSG	Combined Cardiac Panel	GeneDx
7SCFG	Comprehensive Arrhythmia and Cardiomyopathy Gene Panel, Varies	Mayo Clinic Laboratories
3FQNG	Comprehensive Cardiac Arrhythmia/Cardiomyopathy Panel	Northwest Clinical Genomics Lab
438XG	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel	Invitae Corporation
43H3G	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Arrhythmia and Cardiomyopathy	Invitae Corporation
43FSG	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Sudden Unexpected Death in Epilepsy (SUDEP) Genes	Invitae Corporation

GTU	Test Name	Laboratory Name
<b>7UY6G</b>	Postmortem Cardiomyopathy and Arrhythmia Gene Panel, Tissue	Mayo Clinic Laboratories
<b>775JG</b>	Rest of Combined Cardiac after Arrhythmia Panel	GeneDx
<b>775TG</b>	Rest of Combined Cardiac after ARVC Panel	GeneDx
<b>78HUG</b>	Rest of Combined Cardiac after Cardiomyopathy Panel	GeneDx
<b>4XFMG</b>	Sudden Death Syndrome NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>77CZG</b>	Sudden Death Syndrome NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77U8G</b>	Sudden Death Syndrome NGS Panel (Sequencing Only)	Fulgent Genetics

### Comprehensive Cardiomyopathy Panel Tests

GTU	Test Name	Laboratory Name
<b>6S35G</b>	Cardiomyopathy	Fairview Diagnostic Laboratories
<b>5DPTG</b>	Cardiomyopathy Gene Analysis	University of Minnesota Physicians Outreach Laboratory
<b>7VAEG</b>	Cardiomyopathy NGS Panel	AiLife Diagnostics
<b>7VAFG</b>	Cardiomyopathy NGS Panel Rapid	AiLife Diagnostics
<b>2JJGG</b>	Cardiomyopathy Panel	Blueprint Genetics
<b>6DDTG</b>	Cardiomyopathy Panel	GeneDx
<b>4MUVG</b>	Cardiomyopathy Panel	Northwest Clinical Genomics Lab
<b>7DMVG</b>	Cardiomyopathy Panel	Washington University in St. Louis Genomics and Pathology Services
<b>2EMVG</b>	Cardiomyopathy panel NGS	All Children's Hospital
<b>45LWG</b>	Cardiomyopathy Pediatric Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>79PLG</b>	CentoCardio - NGS Panel (CNV included)	Centogene
<b>2YZFG</b>	CMNext	Ambry Genetics
<b>28S8G</b>	Comp Cardiomyopathy Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>2BXWG</b>	Comprehensive Cardiomyopathy (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
<b>7SCCG</b>	Comprehensive Cardiomyopathy Gene Panel, Varies	Mayo Clinic Laboratories
<b>3XZHg</b>	Comprehensive Cardiomyopathy NGS Panel (Deletion/Duplication Only)	Fulgent Genetics

GTU	Test Name	Laboratory Name
<b>25DLG</b>	Comprehensive Cardiomyopathy NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>33E5G</b>	Comprehensive Cardiomyopathy NGS Panel (Sequencing Only)	Fulgent Genetics
<b>345YG</b>	Comprehensive Cardiomyopathy Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>332EG</b>	Comprehensive Cardiomyopathy Panel	Knight Molecular Diagnostic
<b>7DQHG</b>	Comprehensive Cardiomyopathy Panel	PerkinElmer Genomics
<b>56ZPG</b>	GeneSeq Cardio Familial Cardiomyopathy Panel	LabCorp
<b>3GBZG</b>	GeneSeq : Cardio-Familial Cardiomyopathy Profile	Integrated Genetics
<b>5FWFG</b>	Invitae Cardiomyopathy Comprehensive Panel	Invitae Corporation
<b>5U3FG</b>	Pan Cardiomyopathy (92 genes) Deletion/Duplication Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>2G3NG</b>	Pan Cardiomyopathy (92 genes) NGS + Sanger fill-in Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>5638G</b>	Pan Cardiomyopathy (92 genes) NGS Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>7BLQG</b>	Pan Cardiomyopathy Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>57C9G</b>	Pan Cardiomyopathy Panel	PreventionGenetics, part of Exact Sciences
<b>7UHGG</b>	Pan Cardiomyopathy Panel (62 Genes)	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>7UY7G</b>	Postmortem Cardiomyopathy Gene Panel, Tissue	Mayo Clinic Laboratories
<b>6DTCG</b>	Rest of Cardiomyopathy after DCM Panel	GeneDx
<b>6DXQG</b>	Rest of Cardiomyopathy after HCM Panel	GeneDx
<b>7B77G</b>	STAT Comprehensive Cardiomyopathy Panel	PerkinElmer Genomics
<b>7AP3G</b>	Suggested Custom Slice - Lymphedema	GeneDx

### Dilated Cardiomyopathy (DCM) Panel Tests

GTU	Test Name	Laboratory Name
<b>6S48G</b>	Cardiomyopathy, dilated	Fairview Diagnostic Laboratories
<b>2YW9G</b>	DCMNext	Ambry Genetics
<b>5NAGG</b>	Dilated & Arrhythmogenic Cardiomyopathy NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory

GTU	Test Name	Laboratory Name
<b>3JZVG</b>	Dilated Cardiomyopathy – DCM (52 genes) Deletion/Duplication Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>2FP7G</b>	Dilated Cardiomyopathy – DCM (52 genes) NGS + Sanger fill-in Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>3CBJG</b>	Dilated Cardiomyopathy – DCM (52 genes) NGS Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>2JLQG</b>	Dilated Cardiomyopathy (DCM) Panel	Blueprint Genetics
<b>3XP4G</b>	Dilated Cardiomyopathy (DCM) Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>7DMXG</b>	Dilated Cardiomyopathy (DCM) Panel	Washington University in St. Louis Genomics and Pathology Services
<b>7SC4G</b>	Dilated Cardiomyopathy and Left Ventricular Noncompaction Cardiomyopathy Gene Panel, Varies	Mayo Clinic Laboratories
<b>7VUXG</b>	Dilated Cardiomyopathy and Left Ventricular Noncompaction Panel	The University of Chicago Genetic Services
<b>44D6G</b>	Dilated Cardiomyopathy NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>25DFG</b>	Dilated Cardiomyopathy NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>3CJ3G</b>	Dilated Cardiomyopathy NGS Panel (Sequencing Only)	Fulgent Genetics
<b>6DD9G</b>	Dilated Cardiomyopathy Panel	GeneDx
<b>3CBUG</b>	Dilated Cardiomyopathy Panel	Knight Molecular Diagnostic
<b>3XNVG</b>	Dilated Cardiomyopathy Panel	PerkinElmer Genomics
<b>5CZ6G</b>	Dilated Cardiomyopathy Panel	PreventionGenetics, part of Exact Sciences
<b>2KTAG</b>	Dilated Cardiomyopathy Panel, Sequencing	ARUP Laboratories
<b>3XPBG</b>	Dilated Cardiomyopathy Sequencing Panel with CNV Detection: Institution-Specific	PreventionGenetics, part of Exact Sciences
<b>24WTG</b>	Invitae Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Cardiomyopathy	Invitae Corporation
<b>2KYQG</b>	Invitae Dilated Cardiomyopathy and Left Ventricular Noncompaction Panel	Invitae Corporation
<b>5FWEG</b>	Invitae Dilated Cardiomyopathy and Left Ventricular Noncompaction Panel-Add-on Preliminary-evidence Genes for Dilated Cardiomyopathy and Left Ventricular Noncompaction	Invitae Corporation
<b>7B78G</b>	STAT Dilated Cardiomyopathy Panel	PerkinElmer Genomics

## Dystonia Panel Tests

GTU	Test Name	Laboratory Name
<b>77W5G</b>	Basal Ganglia Calcification Dystonia (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>5X7HG</b>	Complete Dopa-Responsive Dystonia (DYT5) Evaluation	Athena Diagnostics Inc
<b>78DLG</b>	Comprehensive Dystonia (NGS Panel and Copy Number Analysis + mtDNA + HTT Repeat Expansion Analysis)	MNG Laboratories
<b>77FMG</b>	Comprehensive Dystonia (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
<b>249UG</b>	Dopa-Responsive Dystonia (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>6FW7G</b>	Dopa-Responsive Dystonia NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>6CT3G</b>	Dopa-Responsive Dystonia NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>3PSFG</b>	Dopa-Responsive Dystonia NGS Panel (Sequencing Only)	Fulgent Genetics
<b>77G9G</b>	Dystonia	Fairview Diagnostic Laboratories
<b>6USCG</b>	Dystonia Chorea Parkinson Exome	The University of Chicago Genetic Services
<b>4WB4G</b>	Dystonia Dyskinesia NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>76VXG</b>	Dystonia Dyskinesia NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>76WNG</b>	Dystonia Dyskinesia NGS Panel (Sequencing Only)	Fulgent Genetics
<b>5JVSG</b>	Dystonia Exome	The University of Chicago Genetic Services
<b>77AHG</b>	Dystonia Panel	Blueprint Genetics
<b>775XG</b>	Dystonia Panel	GeneDx
<b>775BG</b>	Dystonia Panel	Knight Molecular Diagnostic
<b>77JMG</b>	Dystonia Panel	PerkinElmer Genomics
<b>78FUG</b>	Dystonia Panel	PreventionGenetics, part of Exact Sciences
<b>778BG</b>	Invitae Dystonia Comprehensive Panel	Invitae Corporation
<b>76S3G</b>	Invitae Dystonia Comprehensive Panel-Add-on Preliminary-evidence Genes for Dystonia	Invitae Corporation
<b>4SRKG</b>	Isolated Dystonia Evaluation	Athena Diagnostics Inc
<b>5X6QG</b>	Myoclonic Dystonia Panel	PerkinElmer Genomics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>4AANG</b>	Myoclonus dystonia	Fairview Diagnostic Laboratories
<b>77MVG</b>	OXPHOS Defect Dystonia (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
<b>76Q3G</b>	Primary Dystonia (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>7AYEG</b>	STAT Dystonia Panel	PerkinElmer Genomics
<b>7AYCG</b>	STAT Myoclonic Dystonia Panel	PerkinElmer Genomics

### Epilepsy and Seizure Disorder Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>65CUG</b>	Actionable Epilepsy NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>78D5G</b>	Actionable Epilepsy NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>76Y6G</b>	Actionable Epilepsy NGS Panel (Sequencing Only)	Fulgent Genetics
<b>3UT6G</b>	Adolescent/Adult Epilepsy NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>78BJG</b>	Adolescent/Adult Epilepsy NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77UVG</b>	Adolescent/Adult Epilepsy NGS Panel (Sequencing Only)	Fulgent Genetics
<b>77RAG</b>	Beyond Paediatric Epilepsy Panel – for Europe and Middle East	Blueprint Genetics
<b>3PZSG</b>	Childhood Epilepsy NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>77UAG</b>	Childhood Epilepsy NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>76XSG</b>	Childhood Epilepsy NGS Panel (Sequencing Only)	Fulgent Genetics
<b>784QG</b>	Childhood Epilepsy Panel	Knight Molecular Diagnostic
<b>87ASG</b>	Clinical Epilepsy NGS Panel	LabCorp
<b>78L9G</b>	Clinical Epilepsy NGS Panel	MNG Laboratories
<b>77ASG</b>	Comprehensive Epilepsy (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
<b>73X4G</b>	Comprehensive Epilepsy NGS Panel	LabCorp
<b>77RBG</b>	Comprehensive Epilepsy Panel	Blueprint Genetics
<b>78KDG</b>	Comprehensive Epilepsy Panel	GeneDx
<b>778XG</b>	Comprehensive Epilepsy Panel	PerkinElmer Genomics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7UUBG</b>	Comprehensive Epilepsy Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>7BNUG</b>	Comprehensive Epilepsy With or Without Encephalopathy Gene Panel, Varies	Mayo Clinic Laboratories
<b>77GAG</b>	Early infantile epilepsy with encephalopathy	Fairview Diagnostic Laboratories
<b>78FGG</b>	Early Infantile Epileptic Encephalopathy (EIEE) Panel	The University of Chicago Genetic Services
<b>77V3G</b>	Early Infantile Epileptic Encephalopathy NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>77FPG</b>	Early Infantile Epileptic Encephalopathy Panel	PreventionGenetics, part of Exact Sciences
<b>5DRKG</b>	Early Onset Epileptic Encephalopathy NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>78CPG</b>	Early Onset Epileptic Encephalopathy NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77TTG</b>	Early Onset Epileptic Encephalopathy NGS Panel (Sequencing Only)	Fulgent Genetics
<b>776MG</b>	Epilepsy	Fairview Diagnostic Laboratories
<b>77GTG</b>	Epilepsy Advanced Sequencing and CNV Evaluation	Athena Diagnostics Inc
<b>78FNG</b>	Epilepsy Advanced Sequencing and CNV Evaluation - Epileptic Encephalopathy	Athena Diagnostics Inc
<b>77J4G</b>	Epilepsy Advanced Sequencing and CNV Evaluation - Generalized, Absence, Focal, Febrile and Myoclonic Epilepsies	Athena Diagnostics Inc
<b>77HNG</b>	Epilepsy Advanced Sequencing and CNV Evaluation - Infantile Spasms	Athena Diagnostics Inc
<b>78FPG</b>	Epilepsy Advanced Sequencing and CNV Evaluation - Intellectual Disability	Athena Diagnostics Inc
<b>77HQG</b>	Epilepsy Advanced Sequencing and CNV Evaluation - Syndromic Disorders	Athena Diagnostics Inc
<b>7DP9G</b>	Epilepsy and Seizure Panel	PreventionGenetics, part of Exact Sciences
<b>5DSFG</b>	Epilepsy Comprehensive NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>7724G</b>	Epilepsy Comprehensive NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77D4G</b>	Epilepsy Comprehensive NGS Panel (Sequencing Only)	Fulgent Genetics
<b>6DDAG</b>	Epilepsy Deletion/Duplication Panel	GeneDx
<b>77GZG</b>	Epilepsy Exome	The University of Chicago Genetic Services

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>772EG</b>	EPILEPSY GENE PANEL COMPREHENSIVE	Ann and Robert Lurie Children's Hospital of Chicago
<b>77HKG</b>	EPILEPSY GENE PANEL, COMPREHENSIVE WHEN ADFLE GENES NEGATIVE	Ann and Robert Lurie Children's Hospital of Chicago
<b>772DG</b>	EPILEPSY GENE PANEL, COMPREHENSIVE WHEN INFANTILE PANEL NEGATIVE	Ann and Robert Lurie Children's Hospital of Chicago
<b>78FKG</b>	EPILEPSY GENE PANEL, COMPREHENSIVE WHEN THERAPEUTIC PANEL NEGATIVE	Ann and Robert Lurie Children's Hospital of Chicago
<b>77HJG</b>	EPILEPSY GENE PANEL, INFANTILE	Ann and Robert Lurie Children's Hospital of Chicago
<b>34NLG</b>	EPILEPSY GENE PANEL, NOCTURNAL FRONTAL LOBE	Ann and Robert Lurie Children's Hospital of Chicago
<b>77YAG</b>	EPILEPSY GENE PANEL, THERAPEUTIC	Ann and Robert Lurie Children's Hospital of Chicago
<b>7V2RG</b>	Epilepsy Panel	Baylor Genetics, LLC
<b>78FEG</b>	Epilepsy Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>79Q4G</b>	Epilepsy panel - NGS Panel (CNV included)	Centogene
<b>24DMG</b>	Epilepsy, familial focal	Fairview Diagnostic Laboratories
<b>6MJQG</b>	Epilepsy, familial temporal lobe	Fairview Diagnostic Laboratories
<b>34NJG</b>	Epilepsy, nocturnal frontal lobe	Fairview Diagnostic Laboratories
<b>77Y5G</b>	Epilepsy/Seizure NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>77LQG</b>	Epilepsy/Seizures panel	Knight Molecular Diagnostic
<b>78G2G</b>	EpilepsyNext	Ambry Genetics
<b>78GSG</b>	EpilepsyNext-Expanded	Ambry Genetics
<b>776GG</b>	Epileptic Encephalopathy (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>779KG</b>	Epileptic Encephalopathy Panel	Blueprint Genetics
<b>86PAG</b>	EpiPanelDxPLUS	Bionano Laboratories
<b>772UG</b>	EpiRapid	Ambry Genetics
<b>775EG</b>	EpiXpanded Panel	GeneDx
<b>6YMKG</b>	Genomic Unity Epilepsy Analysis	Variantyx
<b>779WG</b>	Idiopathic Generalized and Focal Epilepsy Panel	Blueprint Genetics
<b>77P9G</b>	Infantile Epilepsy Panel	Knight Molecular Diagnostic
<b>787MG</b>	Infantile Epilepsy Panel	PerkinElmer Genomics
<b>3Q7DG</b>	Infantile Spasms NGS Panel (Deletion/Duplication Only)	Fulgent Genetics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>76XZG</b>	Infantile Spasms NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>76WJG</b>	Infantile Spasms NGS Panel (Sequencing Only)	Fulgent Genetics
<b>76TEG</b>	Invitae Epilepsy Panel	Invitae Corporation
<b>7834G</b>	Invitae Epilepsy Panel-Add-on Preliminary-evidence Genes for Epilepsy	Invitae Corporation
<b>779ZG</b>	Metabolic Epilepsy Panel	Blueprint Genetics
<b>5DQXG</b>	Neonatal Epilepsy NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>77VEG</b>	Neonatal Epilepsy NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77VNG</b>	Neonatal Epilepsy NGS Panel (Sequencing Only)	Fulgent Genetics
<b>78HSG</b>	PGmaxTM - Comprehensive Epilepsy and Seizure Panel	PreventionGenetics, part of Exact Sciences
<b>77XKG</b>	PGXome Custom - Early Epileptic Encephalopathy, Dominant and X-linked	PreventionGenetics, part of Exact Sciences
<b>7764G</b>	PGXome Custom - Early Infantile Epileptic Encephalopathy, Dominant and X-linked	PreventionGenetics, part of Exact Sciences
<b>78EUG</b>	PGXome Custom - Early Infantile Epileptic Encephalopathy, Recessive	PreventionGenetics, part of Exact Sciences
<b>5HKQG</b>	Seizures, benign familial infantile	Fairview Diagnostic Laboratories
<b>34QFG</b>	Seizures, benign neonatal	Fairview Diagnostic Laboratories
<b>7B4AG</b>	STAT Comprehensive Epilepsy Panel	PerkinElmer Genomics
<b>7V2QG</b>	STAT Epilepsy Panel	Baylor Genetics, LLC
<b>7AY4G</b>	STAT Infantile Epilepsy Panel	PerkinElmer Genomics

### Exome Sequencing Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7AY2G</b>	Add Familial Report to Previous Whole Exome Sequencing TRIO Test	PerkinElmer Genomics
<b>2FV5G</b>	Any Panel Expand to Exome Plus	Blueprint Genetics
<b>4BXYG</b>	Any Panel Expand to Exome Plus Family	Blueprint Genetics
<b>79AQG</b>	CentoDx - NGS Panel	Centogene
<b>79ARG</b>	CentoLCV - NGS Panel (CNV included)	Centogene

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>46YFG</b>	CHOP Medical Exome	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>46WXG</b>	CHOP Medical Exome + MitoGenome Combined Test	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>6RG2G</b>	Chromosomal Sequencing Analysis (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77JAG</b>	Clinical Exome (Sequencing & Del/Dup) Trio	Fulgent Genetics
<b>77YSG</b>	Clinical Exome (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>78GBG</b>	Clinical Exome (Sequencing Only)	Fulgent Genetics
<b>78GCG</b>	Clinical Exome (Sequencing) Trio	Fulgent Genetics
<b>537AG</b>	Clinical Exome Sequencing	DNA Diagnostic Laboratory - Johns Hopkins Hospital
<b>24C9G</b>	Clinical Exome Sequencing - Duo (Proband and 1 family member)	UCLA Diagnostic Molecular Pathology Laboratory
<b>2WCZG</b>	Clinical Exome Sequencing - Exome re-analysis	UCLA Diagnostic Molecular Pathology Laboratory
<b>33B2G</b>	Clinical Exome Sequencing - Family Trio (Proband and two family members preferred)	UCLA Diagnostic Molecular Pathology Laboratory
<b>2XH2G</b>	Clinical Exome Sequencing - Individual/Proband only	UCLA Diagnostic Molecular Pathology Laboratory
<b>2ZN9G</b>	Clinical Exome Sequencing - Quad (Proband and 3 family members)	UCLA Diagnostic Molecular Pathology Laboratory
<b>7TYRG</b>	Critical Trio Whole Exome Sequencing	University of Michigan - Michigan Medical Genetics Laboratories
<b>7VBYG</b>	Duo Exome	AiLife Diagnostics
<b>86SUG</b>	Duo Whole Exome Sequencing	Baylor Genetics, LLC
<b>2TQRG</b>	Exome Family Member	University of Minnesota Physicians Outreach Laboratory
<b>3V27G</b>	Exome Select	The University of Chicago Genetic Services
<b>7UNTG</b>	Exome Sequencing	ARUP Laboratories
<b>7VSBG</b>	Exome Sequencing	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>24Q2G</b>	Exome Sequencing - Proband	Knight Molecular Diagnostic
<b>33W5G</b>	Exome Sequencing - Trio	Knight Molecular Diagnostic
<b>2KSYG</b>	Exome Sequencing (Proband Only)	The University of Chicago Genetic Services
<b>2G2KG</b>	Exome Sequencing (Proband)	Northwest Clinical Genomics Lab
<b>4GMQG</b>	Exome Sequencing (Trio)	The University of Chicago Genetic Services
<b>4M9ZG</b>	Exome Sequencing Comparator	Northwest Clinical Genomics Lab

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2WWZG</b>	Exome Sequencing Re-analysis	Northwest Clinical Genomics Lab
<b>7VSAG</b>	Exome Sequencing Result	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>53RNG</b>	Exome Sequencing Symptom-Guided Analysis	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<b>4N3SG</b>	Exome Sequencing Trio	Northwest Clinical Genomics Lab
<b>7UNSG</b>	Exome Sequencing, Familial Control	ARUP Laboratories
<b>46SRG</b>	Exome, Family Member	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>2YZMG</b>	ExomeNext-Duo	Ambry Genetics
<b>28FHG</b>	ExomeNext-Duo plus mtDNA	Ambry Genetics
<b>2YZRG</b>	ExomeNext-Proband	Ambry Genetics
<b>2YZQG</b>	ExomeNext-Proband plus mtDNA	Ambry Genetics
<b>2YZVG</b>	ExomeNext-Trio	Ambry Genetics
<b>5U8XG</b>	ExomeNext-Trio plus mtDNA	Ambry Genetics
<b>33Z3G</b>	ExomeSeq - Whole Exome Sequencing	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>4SUTG</b>	Expand to Exome-Each Additional Family Member	Blueprint Genetics
<b>7TLGG</b>	Expanded Exome Upgrade per Sample	Praxis Genomics, LLC
<b>7TLMG</b>	Expanded Exome, 4 Sample	Praxis Genomics, LLC
<b>7TAKG</b>	Expanded Exome, Duo	Praxis Genomics, LLC
<b>7TLJG</b>	Expanded Exome, Proband	Praxis Genomics, LLC
<b>7TLLG</b>	Expanded Exome, Trio	Praxis Genomics, LLC
<b>7A7AG</b>	Family Member Comparator Specimen for Exome Sequencing, Varies	Mayo Clinic Laboratories
<b>6ZSHG</b>	Genomic Unity Exome Analysis	Variantyx
<b>7ABTG</b>	Genomic Unity Exome Plus Analysis	Variantyx
<b>2BFBG</b>	GML Exome Family Member Peripheral blood draw	UCSF Molecular Diagnostics Laboratory
<b>78GXG</b>	IMMUNOLOGY EXOME	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>6S7CG</b>	Invitae Exome, Duo	Invitae Corporation
<b>6S7DG</b>	Invitae Exome, Proband-Only	Invitae Corporation

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6RLHG</b>	MNG Exome Additional Comparator (only available when trio is ordered)	MNG Laboratories
<b>6RLRG</b>	MNG Exome DUO Sequencing	MNG Laboratories
<b>78DUG</b>	MNG Exome DUO Sequencing + mtDNA	MNG Laboratories
<b>6RLPG</b>	MNG Exome Proband Only Sequencing	MNG Laboratories
<b>6RLJG</b>	MNG Exome Proband Only Sequencing + mtDNA	MNG Laboratories
<b>6RGKG</b>	MNG Exome TRIO Sequencing	MNG Laboratories
<b>77FXG</b>	MNG Exome TRIO Sequencing + mtDNA	MNG Laboratories
<b>4AVGG</b>	NextStepDx PLUS	Bionano Laboratories
<b>5E4FG</b>	PGxome Diagnostic Exome Test - Duo	PreventionGenetics, part of Exact Sciences
<b>5E5HG</b>	PGxome Diagnostic Exome Test - Trio	PreventionGenetics, part of Exact Sciences
<b>5E4SG</b>	PGxome Diagnostic Exome Test Patient Plus	PreventionGenetics, part of Exact Sciences
<b>6LUFG</b>	PGxome Prenatal Exome Test - Duo	PreventionGenetics, part of Exact Sciences
<b>5NNGY</b>	PGxome Prenatal Exome Test - Trio	PreventionGenetics, part of Exact Sciences
<b>6LUGG</b>	PGxome Prenatal Exome Test Patient Only	PreventionGenetics, part of Exact Sciences
<b>5E3RG</b>	PGxome Diagnostic	PreventionGenetics, part of Exact Sciences
<b>6QZH</b>	Prenatal Exome Sequencing	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>7BM3G</b>	PRENATAL EXOMESEQ	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>55S7G</b>	Prenatal Trio Whole Exome Sequencing	Baylor Genetics, LLC
<b>7VBUG</b>	Proband Exome	AiLife Diagnostics
<b>2ZBFG</b>	Proband Whole Exome Sequencing	Baylor Genetics, LLC
<b>27YXG</b>	Proband Whole Exome Sequencing + Chromosomal Microarray Analysis (CMA) (Comprehensive)	Baylor Genetics, LLC
<b>76VCG</b>	Proband Whole Exome Sequencing + Comprehensive mtDNA Analysis	Baylor Genetics, LLC
<b>2BJCG</b>	Reanalysis and Interpretation for WES (Internal)	PerkinElmer Genomics
<b>2LFVG</b>	REFLEX to Exome Sequencing	Northwest Clinical Genomics Lab
<b>6DM5G</b>	Reflex to Whole Exome after Slice (Proband Only)	GeneDx
<b>365BG</b>	Reflex to Whole Exome after Slice (Trio)	GeneDx

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7B6CG</b>	Reflex to Whole Exome Sequencing (from panel)	PerkinElmer Genomics
<b>2YZUG</b>	Sequencing plus raw data and filtered variant list (no analysis)	Ambry Genetics
<b>2YWFG</b>	Sequencing plus raw data only	Ambry Genetics
<b>2G7GG</b>	Sequential Trio Whole Exome Sequencing	Baylor Genetics, LLC
<b>4PFEG</b>	Single Gene Expand to Exome Plus Family (trios)	Blueprint Genetics
<b>2WP2G</b>	Single Gene Expanded to Exome Plus	Blueprint Genetics
<b>348XG</b>	Total BluePrint Panel	Baylor Genetics, LLC
<b>7VBWG</b>	Trio Exome	AiLife Diagnostics
<b>6QQMG</b>	Trio Whole Exome Sequencing	Baylor Genetics, LLC
<b>776NG</b>	Trio Whole Exome Sequencing + Comprehensive mtDNA Analysis	Baylor Genetics, LLC
<b>2BFEG</b>	UCSF Genomics Blood Draw	UCSF Molecular Diagnostics Laboratory
<b>2LNHG</b>	WES - Additional Affected Sibling	Baylor Genetics, LLC
<b>22DJG</b>	WES Single or Multi-Sample (Duo, Trio, Quad, etc.)	Medical College of Wisconsin - Human and Molecular Genetics Center
<b>5HGMG</b>	Whole Exome	Blueprint Genetics
<b>2TFEG</b>	Whole Exome (Sequencing & Del/Dup) Trio	Fulgent Genetics
<b>3JD6G</b>	Whole Exome (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>3FAPG</b>	Whole Exome (Sequencing Only)	Fulgent Genetics
<b>3AALG</b>	Whole Exome (Sequencing) Trio	Fulgent Genetics
<b>53N9G</b>	Whole Exome Additional Family Members	Blueprint Genetics
<b>5JSMG</b>	Whole Exome Family	Blueprint Genetics
<b>2Z7LG</b>	Whole Exome Sequencing	Gene by Gene
<b>6QZFG</b>	Whole Exome Sequencing	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>7RSXG</b>	Whole Exome Sequencing	Rady Children's Institute for Genomic Medicine
<b>5BY3G</b>	Whole Exome Sequencing	University of Minnesota Physicians Outreach Laboratory
<b>7R8CG</b>	Whole Exome Sequencing	Washington University in St. Louis Genomics and Pathology Services
<b>5JQPG</b>	Whole Exome Sequencing - DUO (Proband)	LabCorp
<b>7PQEG</b>	Whole Exome Sequencing – DUO (Proband), Products of Conception (POC)	LabCorp
<b>5JR8G</b>	Whole Exome Sequencing - Proband Only	LabCorp

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7PQFG</b>	Whole Exome Sequencing – Proband Only, Products of Conception (POC)	LabCorp
<b>2Z5DG</b>	Whole Exome Sequencing - Reanalysis	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>5XKLG</b>	Whole Exome Sequencing - TRIO (Proband)	LabCorp
<b>7PQDG</b>	Whole Exome Sequencing – TRIO (Proband), Products of Conception (POC)	LabCorp
<b>6LSZG</b>	Whole Exome Sequencing Comparator - Additional FM	LabCorp
<b>6LT2G</b>	Whole Exome Sequencing Comparator - Father	LabCorp
<b>6LSYG</b>	Whole Exome Sequencing Comparator - Mother	LabCorp
<b>7A7EG</b>	Whole Exome Sequencing for Hereditary Disorders , Varies	Mayo Clinic Laboratories
<b>3SMCG</b>	Whole Exome Sequencing Proband Only	PerkinElmer Genomics
<b>7AYYG</b>	Whole Exome Sequencing QUAD	PerkinElmer Genomics
<b>2DR8G</b>	Whole Exome Sequencing TRIO	PerkinElmer Genomics
<b>7AYLG</b>	Whole Exome Sequencing, DUO	PerkinElmer Genomics
<b>2D8LG</b>	Whole Exome Sequencing: Duo Analysis	University of Wisconsin - Madison WSLH - UW Cytogenetic Services
<b>2DR9G</b>	Whole Exome Sequencing: Proband Analysis Only	University of Wisconsin - Madison WSLH - UW Cytogenetic Services
<b>33NTG</b>	Whole Exome Sequencing: Trio Analysis	University of Wisconsin - Madison WSLH - UW Cytogenetic Services
<b>5FMPG</b>	Whole exome via NGS (Duo with full parental report)	Allele Diagnostics
<b>5FMNG</b>	Whole exome via NGS (Proband only)	Allele Diagnostics
<b>5FMQG</b>	Whole exome via NGS (Trio with full parental reports)	Allele Diagnostics
<b>24C4G</b>	Whole exome via NGS (Trio with proband report only)	Allele Diagnostics
<b>7TLFG</b>	Whole Exome, 4 Sample	Praxis Genomics, LLC
<b>7TLDG</b>	Whole Exome, Duo	Praxis Genomics, LLC
<b>7TLCG</b>	Whole Exome, Proband	Praxis Genomics, LLC
<b>7TLEG</b>	Whole Exome, Trio	Praxis Genomics, LLC
<b>6L9SG</b>	XomeDx - Proband	GeneDx
<b>6RVYG</b>	XomeDx - Trio	GeneDx
<b>6L9QG</b>	XomeDx - Duo	GeneDx
<b>7TVKG</b>	XomeDx Express (GeneDx), Blood	University of Michigan - Michigan Medical Genetics Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7TVLG</b>	XomeDx Express, Buccal Swab	University of Michigan - Michigan Medical Genetics Laboratories
<b>4JJWG</b>	XomeDx Fetal - Duo	GeneDx
<b>7DQWG</b>	XomeDx Plus - Duo	GeneDx
<b>77MHG</b>	XomeDx Plus - Proband	GeneDx
<b>77M2G</b>	XomeDx Plus - Trio	GeneDx
<b>7TV8G</b>	XomeDx Plus- Duo, blood	University of Michigan - Michigan Medical Genetics Laboratories
<b>7TV7G</b>	XomeDx Plus- Duo, buccal kit	University of Michigan - Michigan Medical Genetics Laboratories
<b>7TVBG</b>	XomeDx Plus- Proband, blood	University of Michigan - Michigan Medical Genetics Laboratories
<b>7TV9G</b>	XomeDx Plus- Trio, blood	University of Michigan - Michigan Medical Genetics Laboratories
<b>7TVCG</b>	XomeDX Plus-Trio, buccal kit	University of Michigan - Michigan Medical Genetics Laboratories
<b>2W48G</b>	XomeDx Prenatal - Comprehensive	GeneDx
<b>46N6G</b>	XomeDx Trio	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>4GZRG</b>	XomeDxFetal- Trio	GeneDx
<b>46J4G</b>	XomeDxPlus Proband	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>46V3G</b>	XomeDxPlus Trio	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>777MG</b>	XomeDxPlus WES + mtDNA Sequencing and Deletion Testing	University of Minnesota Physicians Outreach Laboratory

### Expanded Carrier Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7SSBG</b>	Beacon 787-Expanded Carrier Screening Panel (With X-linked Disorders) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7SSCG</b>	Beacon 787-Expanded Carrier Screening Panel (Without X-linked Disorders) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>86GSG</b>	Beacon ACMG Tier 3 Female Carrier Screening Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>86GTG</b>	Beacon ACMG Tier 3 Male Carrier Screening Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>6LMNG</b>	Beacon ACOG/ACMG Female Carrier Screening Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>5DRBG</b>	Beacon ACOG/ACMG Male Carrier Screening Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>78HAG</b>	Beacon Expanded Female Carrier Screening Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77ZHG</b>	Beacon Expanded Female Carrier Screening Plus Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77ZJG</b>	Beacon Expanded Male Carrier Screening Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7743G</b>	Beacon Expanded Male Carrier Screening Plus Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77K3G</b>	Beacon Focus Female Carrier Screening Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>78HBG</b>	Beacon Focus Male Carrier Screening Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7QH4G</b>	CARRIER SCREEN - CLASSIC, FEMALE (24 GENE PANEL)	Clinical Pathology Laboratories
<b>7QH5G</b>	CARRIER SCREEN - CLASSIC, MALE (22 GENE PANEL)	Clinical Pathology Laboratories
<b>7QGZG</b>	CARRIER SCREEN - COMPLETE, MALE (147 GENE PANEL)	Clinical Pathology Laboratories
<b>79SAG</b>	Carrier Screening (Horizon)	ProPath
<b>6U8PG</b>	Common Carrier Screening Panel	Connective Tissue Gene Tests
<b>7V3RG</b>	Comprehensive Carrier Screening Panel for Genetic Conditions	Genesys Diagnostics Inc
<b>78GTG</b>	Expanded Carrier Screen by Next Generation Sequencing with Fragile X	ARUP Laboratories
<b>7832G</b>	Expanded Carrier Screening	UCSF Molecular Diagnostics Laboratory
<b>86D7G</b>	Extended Carrier Screening Panel	Connective Tissue Gene Tests
<b>6RN4G</b>	Foresight Fundamental Plus panel	Myriad Genetics
<b>78HYG</b>	Foresight Universal Panel Carrier Screen	Myriad Genetics
<b>34P9G</b>	FUNDAMENTAL PLUS 2528 (COUNSYL)	Enzo Clinical Labs
<b>7V27G</b>	GeneAware - Complete Panel - Female	Baylor Genetics, LLC
<b>7V22G</b>	GeneAware - Complete Panel - Male	Baylor Genetics, LLC
<b>7V26G</b>	GeneAware™ ACMG & ACOG Panel (Female)	Baylor Genetics, LLC
<b>7UZZG</b>	GeneAware™ ACMG & ACOG Panel (Male)	Baylor Genetics, LLC
<b>7V24G</b>	GeneAware™ Expanded Panel (Female)	Baylor Genetics, LLC
<b>7UZXG</b>	GeneAware™ Expanded Panel (Male)	Baylor Genetics, LLC

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7VXPG</b>	GeneAware™ Expanded Plus Panel (Female)	Baylor Genetics, LLC
<b>7VXQG</b>	GeneAware™ Expanded Plus Panel (Male)	Baylor Genetics, LLC
<b>6343G</b>	GeneSeq PLUS	Integrated Genetics
<b>7SVAG</b>	GeneSeq PLUS	LabCorp
<b>3NY2G</b>	GeneSeq PLUS without VUS	Integrated Genetics
<b>4ZMUG</b>	GeneSeq PLUS without VUS, Prenatal	Integrated Genetics
<b>2AL5G</b>	GeneSeq PLUS, Prenatal	Integrated Genetics
<b>7759G</b>	Horizon 14 (14 disease panel)	Natera
<b>7758G</b>	Horizon 27 (27 disease Pan-ethnic Standard panel)	Natera
<b>7757G</b>	Horizon 274 (274 disease Pan-ethnic Extended panel)	Natera
<b>72ESG</b>	Horizon 421	Natera
<b>7A7WG</b>	Horizon ACMG Panel	Natera
<b>6V4QG</b>	INHERIGEN	GenPath Diagnostics
<b>6UWPG</b>	INHERIGEN PLUS	GenPath Diagnostics
<b>6Y5WG</b>	InheriGenTx	BioReference Laboratories
<b>7SUSG</b>	Inheritest 100 PLUS Panel	LabCorp
<b>7SUTG</b>	Inheritest 300 PLUS Panel	LabCorp
<b>77LMG</b>	Inheritest 500 PLUS Panel	Integrated Genetics
<b>77BTG</b>	Inheritest 500 PLUS Panel	LabCorp
<b>7SURG</b>	Inheritest High Frequency Panel	LabCorp
<b>7838G</b>	Inheritest 500 PLUS with Repro Partners Report	Integrated Genetics
<b>77ENG</b>	Inheritest Carrier Screen - Society Guided Panel (14 Genes)	Integrated Genetics
<b>7SRSG</b>	M Beacon Focus B, 14 Gene Panel	LabCorp
<b>7ZPEG</b>	MYRIAD FORESIGHT CARRIER SCREENING	University of Illinois at Chicago - Biochemical Genetics Laboratory
<b>784RG</b>	MYRIAD WOMEN\\'S HEALTH FAMILY PREP SCREEN 2	University of California Davis Health System - UCMDMC - Molecular and Cytogenetic Laboratory
<b>78NPG</b>	Natera One Panel	ProPath
<b>3EHWG</b>	NewbornGenID	Advanced Molecular Diagnostics, LLC
<b>2D58G</b>	NewbornGenID include Fragile X (Female Patients Only)	Advanced Molecular Diagnostics, LLC

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7SUAG</b>	QHerit; 381 Diseases, Male	Quest Diagnostics
<b>7VXWG</b>	QHerit; 400 Diseases, Male	Quest Diagnostics
<b>7SU9G</b>	QHerit; 421 Diseases, Female	Quest Diagnostics
<b>7VXXG</b>	QHerit; 445 Diseases, Female	Quest Diagnostics
<b>6UVXG</b>	QHerit; Expanded Carrier Screen	Quest Diagnostics
<b>7DHGG</b>	QHerit; Extended, Female	Quest Diagnostics
<b>7DHHG</b>	QHerit; Extended, Male	Quest Diagnostics
<b>7DHJG</b>	QHerit; Plus, Female	Quest Diagnostics
<b>7DHKG</b>	QHerit; Plus, Male	Quest Diagnostics
<b>4TGTG</b>	UNIVERSAL GENETIC TEST 2521 (COUNSYL)	Enzo Clinical Labs
<b>4B77G</b>	UNIVERSAL PANEL PLUS THROMBOPHILIAS 2524 (COUNSYL)	Enzo Clinical Labs
<b>4CK8G</b>	UNIVERSAL PLUS GENETICS TEST 2522 (COUNSYL)	Enzo Clinical Labs

### Hematologic Malignancy Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2FKWG</b>	Comprehensive Hematologic Cancer Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7SFFG</b>	Comprehensive HemeComplete Profile	PathGroup
<b>2DVSG</b>	Comprehensive Lymphoid Oncology (DNA and RNA analysis)	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>7SFEG</b>	Comprehensive Myeloid Profile	PathGroup
<b>5M87G</b>	FISH Panel : Myeloid Disorders	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>7VMHG</b>	FISH Panel: Myeloid Malignancy	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>6Z8DG</b>	FoundationOne Heme	Foundation Medicine
<b>33FRG</b>	Hematologic Cancer Fusion Analysis	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>45EWG</b>	Hematologic Cancer Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>73H3G</b>	Hematologic Neoplasm Mutation Panel	Ohio State University - Molecular Pathology Laboratory
<b>34YSG</b>	Hematological Malignancies GeneTrails Hematologic Malignancies 220 Gene Panel	Knight Molecular Diagnostic

GTU	Test Name	Laboratory Name
<b>34PNG</b>	HemaVision Leukemia Panel	University of Oklahoma Health Sciences Center - Molecular Pathology Laboratory
<b>7ATGG</b>	HEME DNA MUTATION PANEL (152 gene DNA NGS panel for mutations)	Texas Children's Hospital
<b>7ATHG</b>	HEME DNA/RNA COMBINED PANEL (Combined heme mutation and fusion panels)	Texas Children's Hospital
<b>6LTNG</b>	Heme Gene Panel by NGS	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>7SFCG</b>	HemeComplete NextGen Sequencing Assay	PathGroup
<b>72AHG</b>	IntelliGEN Myeloid	LabCorp
<b>7RJGG</b>	JMML Associated Exon Panel (JMML)	Rady Children's Institute for Genomic Medicine
<b>5PPVG</b>	Juvenile Myelomonocytic Leukemia Associated Exon Panel	UCSF Molecular Diagnostics Laboratory
<b>28E8G</b>	Leukemia Trial NGS Panel, Blood	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>28MEG</b>	Leukemia Trial NGS Panel, Bone Marrow	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>6LTPG</b>	Lymphoid Gene Panel by NGS	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>2DJPG</b>	Lymphoid Oncology DNA Analysis	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>6U9NG</b>	MayoComplete Myeloid Neoplasms, Comprehensive OncoHeme Next-Generation Sequencing, Varies	Mayo Clinic Laboratories
<b>7SBCG</b>	MayoComplete Plasma Cell Myeloma, Next-Generation Sequencing, Varies	Mayo Clinic Laboratories
<b>2XDBG</b>	Myeloid Complete Molecular Profile	Genetic Associates
<b>6VB9G</b>	Myeloid Extended Mutation Analysis Panel by Next Generation Sequencing	Molecular Pathology Laboratory Network
<b>6LTQG</b>	Myeloid Gene Panel by NGS	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>7UXCG</b>	Myeloid Malignancies Mutation and Copy Number Variation Panel by Next Generation Sequencing	ARUP Laboratories
<b>7ALDG</b>	MYELOID MALIGNANCIES MUTATION PANEL	Clinical Pathology Laboratories
<b>2XRDG</b>	Myeloid Malignancies Mutation Panel by Next Generation Sequencing	ACL Laboratories
<b>6VAZG</b>	Myeloid Malignancies Mutation Panel by Next Generation Sequencing	ARUP Laboratories

GTU	Test Name	Laboratory Name
<b>5TUPG</b>	Myeloid Malignancy Comprehensive NGS Panel	University of Minnesota Physicians Outreach Laboratory
<b>78N9G</b>	Myeloid Multigene Panel	UCSF Molecular Diagnostics Laboratory
<b>5CTBG</b>	Myeloid Mutation Panel- AML	University of North Carolina Hospitals - Molecular Genetics
<b>7Q2MG</b>	Myeloid Mutation Panel, Blood	Nebraska Medical Center - Molecular Diagnostic Laboratory
<b>7Q2HG</b>	Myeloid Mutation Panel, Other	Nebraska Medical Center - Molecular Diagnostic Laboratory
<b>8EFPG</b>	Myeloid Neoplasm Mutation Analysis Only	Medfusion
<b>7SFBG</b>	Myeloid NextGen Sequencing Assay	PathGroup
<b>7V4RG</b>	Myeloid NGS	Allina Health Laboratory
<b>7TV4G</b>	Myeloid NGS Panel	University of Michigan - Michigan Medical Genetics Laboratories
<b>7SQ8G</b>	Myeloid Panel NGS Bone Marrow	Cleveland Clinic Laboratories
<b>7SQ7G</b>	Myeloid Panel NGS Peripheral Blood	Cleveland Clinic Laboratories
<b>6UY4G</b>	Myeloid tumor panel - Somatic Mutation Analysis	Centogene
<b>7SPKG</b>	Neo Comprehensive - Myeloid Disorders	NeoGenomics Laboratories
<b>7URQG</b>	Neo Comprehensive™ - Heme Cancers	NeoGenomics Laboratories
<b>3ZSJG</b>	NeoTYPE AITL/Peripheral T-Cell Lymphoma Profile	NeoGenomics Laboratories
<b>86UPG</b>	NeoTYPE Lymphoid Disorders Profile	NeoGenomics Laboratories
<b>6VB7G</b>	NGS Hematologic Malignancy Mutation Panel	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<b>2LJDG</b>	NGS Hematolymphoid Panel (Lab Only)	Molecular Diagnosis - University of Rochester Medical Center - Strong Memorial Hospital
<b>5364G</b>	NGS_Myeloid 37 Genes Panel	CellNetix Pathology and Laboratories
<b>86LPG</b>	OnkoSight Advanced Chronic Lymphoid Neoplasm NGS Panel - PB/BM	BioReference Laboratories
<b>7C5AG</b>	OnkoSight Advanced NGS Myeloid Panel	BioReference Laboratories
<b>7T3WG</b>	OnkoSight Advanced Pan Heme Fusion NGS Panel	BioReference Laboratories
<b>7DN9G</b>	Paired Tumor / Normal – Comprehensive Hematologic Cancer Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7DN8G</b>	Paired Tumor / Normal - Hematologic Cancer Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7UJHG</b>	PennSeq Hematological Malignancies Panel	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>2733G</b>	T Large Granular Lymphocyte NGS Panel	University of Minnesota Physicians Outreach Laboratory

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7ZQTG</b>	Targeted Myeloid Panel (TMP)	Columbia University - Personalized Genomic Medicine
<b>86SCG</b>	Tempus xT: Targeted panel of 648 genes (Hematologic Malignancy)	Tempus AI, Inc.

### Hereditary GI/Colon Cancer Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7TP3G</b>	CancerNext Expanded Colon Lynch	Ambry Genetics
<b>7TNZG</b>	CancerNext: Colon/Lynch	Ambry Genetics
<b>2NBHG</b>	CentoColon - NGS Panel (CNV included)	Centogene
<b>76DFG</b>	COLARIS AP PLUS with Myriad myRisk Hereditary Cancer Update Test	Myriad Genetics
<b>76DEG</b>	COLARIS PLUS with Myriad myRisk Hereditary Cancer Update Test	Myriad Genetics
<b>76DUG</b>	ColoNext	Ambry Genetics
<b>6UQPG</b>	ColoNext with RNAinsight	Ambry Genetics
<b>3598G</b>	Colorectal Cancer Comprehensive Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>76E3G</b>	Colorectal Cancer Comprehensive Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>3ATVG</b>	Colorectal Cancer Comprehensive Panel (Sequencing Only)	Fulgent Genetics
<b>3D5AG</b>	Colorectal Cancer Focus Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>6U88G</b>	Colorectal Cancer Focus Panel (Germline)	NeoGenomics Laboratories
<b>76E4G</b>	Colorectal Cancer Focus Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>34KAG</b>	Colorectal Cancer Focus Panel (Sequencing Only)	Fulgent Genetics
<b>2L58G</b>	Colorectal Cancer Germline NGS Panel	University of Michigan - Michigan Medical Genetics Laboratories
<b>76DSG</b>	Colorectal Cancer Panel	ACL Laboratories
<b>76DWG</b>	Colorectal Cancer Panel	GeneDx
<b>76BGG</b>	Colorectal Cancer Panel	PerkinElmer Genomics
<b>3DW5G</b>	ColoSeq - Lynch and Polyposis Panel	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6V2QG</b>	Comprehensive Colon Cancer Panel (Seq & Del/Dup)	Genetics Center - Molecular and Cytogenetic Diagnostic Laboratories
<b>7TNXG</b>	Comprehensive Common Cancer Panel: Colon/Lynch	GeneDx
<b>6S5NG</b>	Hereditary cancer colon	Fairview Diagnostic Laboratories
<b>2YB4G</b>	Hereditary Colorectal Cancer Panel	Blueprint Genetics
<b>25LZG</b>	Hereditary Colorectal Cancer Panel	Quest Diagnostics
<b>76DZG</b>	Hereditary Colorectal Cancer Panel	The University of Chicago Genetic Services
<b>7AAAG</b>	Hereditary Colorectal Cancer Panel (20 Genes)	Palo Verde Laboratory - division of Sonora Quest Lab
<b>7V2BG</b>	Hereditary Colorectal/Gastrointestinal Cancer Panel	Baylor Genetics, LLC
<b>7SHSG</b>	Hereditary Gastrointestinal Cancer High-Risk Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>76BCG</b>	Hereditary Gastrointestinal Cancer Panel	Blueprint Genetics
<b>76DNG</b>	Hereditary Gastrointestinal Cancer Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>7DLGG</b>	Hereditary Gastrointestinal Cancer Panel, Varies	Mayo Clinic Laboratories
<b>6V9EG</b>	Hereditary High-Risk Colon Cancer Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7V2GG</b>	Hereditary High-Risk Colorectal Cancer Panel	Baylor Genetics, LLC
<b>45K7G</b>	Inherited Colon Cancer Panel	Knight Molecular Diagnostic
<b>778DG</b>	Invitae Hereditary Colorectal Cancer Guidelines-Based Panel	Invitae Corporation
<b>77QBG</b>	Invitae Hereditary Colorectal Cancer Guidelines-Based Panel-Add-on Preliminary-evidence Colorectal Cancer Guidelines Genes	Invitae Corporation
<b>76CYG</b>	Invitae Hereditary Colorectal Cancer Panel	Invitae Corporation
<b>76CZG</b>	Invitae Hereditary Colorectal Cancer Panel-Add-on Preliminary-evidence Genes for Colorectal Cancer	Invitae Corporation
<b>6DXSG</b>	Lynch/Colorectal High Risk Panel	GeneDx
<b>7TNYG</b>	Rest of Comprehensive Common Cancer Panel: Colon/Lynch	GeneDx
<b>7B7KG</b>	STAT Colorectal Cancer Panel	PerkinElmer Genomics
<b>7VPKG</b>	Tempus xG Common Hereditary Cancers 36 genes (hereditary colon cancer indications)	Tempus AI, Inc.
<b>7VPGG</b>	Tempus xG+ Extended Hereditary Cancers 77 genes (hereditary colon cancer indications)	Tempus AI, Inc.
<b>76BSG</b>	VistaSeq Colorectal Cancer Panel	Integrated Genetics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>76DAG</b>	VistaSeq Colorectal Cancer Panel	LabCorp
<b>76CBG</b>	VistaSeq Colorectal Cancer Panel	Labcorp   Oncology
<b>2F64G</b>	VistaSeq High Risk Colorectal Cancer Panel	Integrated Genetics
<b>58EYG</b>	VistaSeq High Risk Colorectal Cancer Panel	LabCorp
<b>4NLFG</b>	VistaSeq High Risk Colorectal Cancer Panel	Labcorp   Oncology

### Hereditary Polyposis Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7TQFG</b>	Adenomatous Polyposis Focus (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>5MJ8G</b>	APC & MUTYH seq and del/dup	Ambry Genetics
<b>6UQRG</b>	APC and MUTYH with RNAinsight	Ambry Genetics
<b>8734G</b>	APC- and MUTYH-Associated Polyposis Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>2AUYG</b>	COLARIS AP	Myriad Genetics
<b>86HDG</b>	ColoSeq Genes Sequenced	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>86HCG</b>	ColoSeq Polyposis	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>4QZBG</b>	Comprehensive Polyposis Syndrome Panel (Seq & Del/Dup Analysis)	Genetics Center - Molecular and Cytogenetic Diagnostic Laboratories
<b>4R4DG</b>	FAP: APC Seq, APC Del/Dup, MUTYH 2 Muts	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<b>6QVMG</b>	Hereditary Polyposis Panel	PreventionGenetics, part of Exact Sciences
<b>7PQHG</b>	Invitae Adenomatous Polyposis Panel	Invitae Corporation
<b>7TQGG</b>	Polyposis Comprehensive (Sequencing & Deletion/Duplication)	Fulgent Genetics

### Hypertrophic Cardiomyopathy (HCM) Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6S4CG</b>	Cardiomyopathy, familial hypertrophic	Fairview Diagnostic Laboratories
<b>5LVCG</b>	HCM Panel Deletion / Duplication	DDC Clinic Laboratory
<b>6PZ3G</b>	HCM Panel Next Generation Sequencing	DDC Clinic Laboratory

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6PZ4G</b>	HCM Panel NGS & Del Dup Comprehensive	DDC Clinic Laboratory
<b>2YZEG</b>	HCMNext	Ambry Genetics
<b>2YZAG</b>	HCMNext Reflex	Ambry Genetics
<b>5W3JG</b>	Hypertrophic Cardiomyopathy – HCM (41 genes) Deletion/Duplication Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>2FPZG</b>	Hypertrophic Cardiomyopathy – HCM (41 genes) NGS + Sanger fill-in Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>3XPUG</b>	Hypertrophic Cardiomyopathy – HCM (41 genes) NGS Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>2GNQG</b>	Hypertrophic Cardiomyopathy (HCM) Panel	Blueprint Genetics
<b>3CEJG</b>	Hypertrophic Cardiomyopathy (HCM) Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>6DXMG</b>	Hypertrophic Cardiomyopathy (HCM) Panel	GeneDx
<b>7DN2G</b>	Hypertrophic Cardiomyopathy (HCM) Panel	Washington University in St. Louis Genomics and Pathology Services
<b>7SBZG</b>	Hypertrophic Cardiomyopathy Gene Panel, Varies	Mayo Clinic Laboratories
<b>5N9HG</b>	Hypertrophic Cardiomyopathy NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>3XG9G</b>	Hypertrophic Cardiomyopathy NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>25E2G</b>	Hypertrophic Cardiomyopathy NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>3XPSG</b>	Hypertrophic Cardiomyopathy NGS Panel (Sequencing Only)	Fulgent Genetics
<b>3XQ7G</b>	Hypertrophic Cardiomyopathy Panel	Knight Molecular Diagnostic
<b>3XQBG</b>	Hypertrophic Cardiomyopathy Panel	PerkinElmer Genomics
<b>5CYCG</b>	Hypertrophic Cardiomyopathy Panel	PreventionGenetics, part of Exact Sciences
<b>7UVVG</b>	Hypertrophic Cardiomyopathy Panel	The University of Chicago Genetic Services
<b>7U3GG</b>	Hypertrophic Cardiomyopathy Panel (GeneDx)	University of Michigan - Michigan Medical Genetics Laboratories
<b>3XNZG</b>	Hypertrophic Cardiomyopathy Panel (PCPGM)	University of Michigan - Michigan Medical Genetics Laboratories
<b>3HV4G</b>	Hypertrophic Cardiomyopathy Panel (PCPGM), Known Mutation	University of Michigan - Michigan Medical Genetics Laboratories
<b>2KTCG</b>	Hypertrophic Cardiomyopathy Panel, Sequencing	ARUP Laboratories
<b>44WDG</b>	Invitae Hypertrophic Cardiomyopathy Panel	Invitae Corporation

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>42LUG</b>	Invitae Hypertrophic Cardiomyopathy Panel-Add-on Preliminary-evidence Genes for Hypertrophic Cardiomyopathy	Invitae Corporation
<b>7B79G</b>	STAT Hypertrophic Cardiomyopathy Panel	PerkinElmer Genomics

### Long QT Syndrome (LQTS) Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>5FWSG</b>	Invitae Long QT Syndrome Panel	Invitae Corporation
<b>786WG</b>	Invitae Long QT Syndrome Panel-Add-on Preliminary-evidence Genes for Long QT Syndrome	Invitae Corporation
<b>77MWG</b>	Long and Short QT Syndrome (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>6Q7RG</b>	Long QT Panel Deletion / Duplication	DDC Clinic Laboratory
<b>782LG</b>	Long QT Panel Next Generation Sequencing	DDC Clinic Laboratory
<b>782KG</b>	Long QT Panel NGS & Del Dup Comprehensive	DDC Clinic Laboratory
<b>2KQNG</b>	Long QT Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>784HG</b>	Long QT syndrome	Fairview Diagnostic Laboratories
<b>39DQG</b>	Long QT Syndrome – LQTS (15 genes) Deletion/Duplication Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>776UG</b>	Long QT Syndrome – LQTS (15 genes) NGS + Sanger fill-in Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>78A2G</b>	Long QT Syndrome – LQTS (15 genes) NGS Panel	Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory
<b>77S8G</b>	Long QT Syndrome (LQTS) Panel	Blueprint Genetics
<b>77BKG</b>	Long QT Syndrome (LQTS) Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>5XA8G</b>	Long QT Syndrome (LQTS) Panel Sequencing	Indiana University Molecular Genetics Diagnostic Laboratory - Indiana University Medical Center, School of Medicine
<b>27X3G</b>	Long QT Syndrome and Short QT Syndrome	Knight Molecular Diagnostic
<b>77C6G</b>	Long QT Syndrome by Next Generation Sequencing (KCNQ1, KCNH2, SCN5A, KCNE1, KCNE2, KCNJ2, CACNA1C, CAV3, SCN4B, AKAP9, SNTA1, ANK2, CALM1, CALM2, KCNJ5)	Medical Diagnostic Laboratories, LLC
<b>5X5SG</b>	Long QT Syndrome Gene Analysis	University of Minnesota Physicians Outreach Laboratory
<b>7SBNG</b>	Long QT Syndrome Gene Panel, Varies	Mayo Clinic Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>77L7G</b>	Long QT syndrome NGS Panel	Connective Tissue Gene Tests
<b>5N7XG</b>	Long QT syndrome NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>6MADG</b>	Long QT syndrome NGS Panel Deletion / Duplication	Connective Tissue Gene Tests
<b>774ZG</b>	Long QT syndrome NGS Panel Next Generation Sequencing	Connective Tissue Gene Tests
<b>76V5G</b>	Long QT Syndrome Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>785ZG</b>	Long QT Syndrome Panel	PerkinElmer Genomics
<b>78FTG</b>	Long QT Syndrome Panel	PreventionGenetics, part of Exact Sciences
<b>7VUUG</b>	Long QT Syndrome Panel	The University of Chicago Genetic Services
<b>7DN3G</b>	Long QT Syndromes panel	Washington University in St. Louis Genomics and Pathology Services
<b>772GG</b>	LongQTNext	Ambry Genetics
<b>783PG</b>	LQTS Panel	GeneDx
<b>2YB5G</b>	LQTS Seq/Del/Dup Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>7B7CG</b>	STAT Long QT Syndrome Panel	PerkinElmer Genomics

### Lung Cancer Diagnostic Algorithmic Tests <sup>1</sup>

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7V4VG</b>	CyPath® Lung	Precision Pathology Laboratory
<b>7S9CG</b>	Knodule ID	MagArray Inc
<b>7TRHG</b>	LungLB® Test	LungLife AI
<b>7RVWG</b>	Nodify CDT	Biodesix
<b>7RVVG</b>	Nodify Lung Nodule Risk Assessment Testing	Biodesix
<b>6USVG</b>	Nodify XL2	Biodesix
<b>7TRTG</b>	OncobiotaLUNGdetect	Micronoma
<b>24LHG</b>	Percepta® Lung Cancer Diagnostics	Veracyte
<b>7S9BG</b>	REVEAL Lung Nodule Characterization	MagArray Inc

## Lung Cancer Panel Tests

GTU	Test Name	Laboratory Name
<b>8EEQG</b>	50SEQ with MSI Panel	Medfusion
<b>5X7JG</b>	BRAF/EGFR NGS if EGFR neg rfx to ALK FISH if neg rfx ROS1	BioReference Laboratories
<b>2LVVG</b>	Comprehensive Lung Panel	University of Iowa Hospitals and Clinics - Department of Pathology
<b>7UQRG</b>	Comprehensive LungSEQPlus Panel	Medfusion
<b>7SNHG</b>	Comprehensive OnkoSight Advanced Lung Panel	BioReference Laboratories
<b>7TMTG</b>	Comprehensive OnkoSight Lung Panel + GeneStrat Liquid Biopsy	BioReference Laboratories
<b>5XBBG</b>	EGFR and KRAS, if both neg, reflex ALK, if neg reflex ROS1	BioReference Laboratories
<b>4R22G</b>	EGFR, if neg reflex to ALK, if neg reflex to ROS1	BioReference Laboratories
<b>3XRVG</b>	EGFR, KRAS, BRAF, PIK3CA mutation analysis, NSCLC panel	University of Pittsburgh Medical Center - Division of Molecular Diagnostics
<b>4NUWG</b>	FISH Panel : Non-Small Cell Lung Cancer	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>6LQSG</b>	Lung adenocarcinoma targeted NGS panel	Allina Health Laboratory
<b>7AFUG</b>	LUNG CANCER	Clinical Pathology Laboratories
<b>7Q8RG</b>	Lung Cancer Mutation Analysis	UCLA Diagnostic Molecular Pathology Laboratory
<b>6WW7G</b>	Lung Cancer Mutation Panel (EGFR, KRAS, ALK)	Quest Diagnostics
<b>42L6G</b>	Lung Cancer NGS Panel	University of Michigan - Michigan Medical Genetics Laboratories
<b>3ZETG</b>	Lung Cancer Panel by Next Generation Sequencing	Molecular Pathology Laboratory Network
<b>7TN5G</b>	Lung Carcinoma NGS Panel with Fusion	University of Minnesota Physicians Outreach Laboratory
<b>7R8HG</b>	LungSEQ Concurrent Panel, without PD-L1	Quest Diagnostics
<b>8EEXG</b>	LUNGSEQ Panel	Medfusion
<b>7AUEG</b>	LUNGSEQ Panel	Quest Diagnostics
<b>8EERG</b>	LUNGSEQ Panel Concurrent	Medfusion
<b>8EFKG</b>	LUNGSEQ Panel with FISH Concurrent - no PD-L1	Medfusion
<b>7RWLG</b>	MayoComplete Lung Cancer Mutations, Next-Generation Sequencing, Tumor	Mayo Clinic Laboratories
<b>7RWMG</b>	MayoComplete Lung Cancer-Targeted Gene Panel with Rearrangement, Tumor	Mayo Clinic Laboratories
<b>7RWKG</b>	MayoComplete Lung Rearrangements, Rapid Test, Tumor	Mayo Clinic Laboratories
<b>3ERKG</b>	MYRIAD MyPlan Lung Cancer	University of Minnesota Physicians Outreach Laboratory

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7BHGX</b>	NeoTYPE® DNA & RNA - Lung	NeoGenomics Laboratories
<b>6YZRG</b>	NeoTYPE® Lung Tumor Profile	NeoGenomics Laboratories
<b>262FG</b>	Non-Small-cell Lung Cancer (NSCLC) Therapeutic Profile II	LabCorp
<b>6ST7G</b>	Oncology FISH Analysis - Non-small Cell Lung Carcinoma Panel	Baylor Genetics, LLC
<b>86LVG</b>	OnkoSight Advanced Comprehensive Lung	BioReference Laboratories
<b>86LLG</b>	OnkoSight Advanced Lung Cancer NGS Panel	BioReference Laboratories
<b>6U25G</b>	OnkoSight Advanced Lung Cancer Panel, ALK and ROS1 by FISH	BioReference Laboratories
<b>7SSUG</b>	OnkoSight Advanced NGS Comprehensive Lung Panel	BioReference Laboratories
<b>2736G</b>	RespriDX™	Interpace Diagnostics
<b>26QHG</b>	Targeted Gene Panel with Fusions, Lung Cancer	Palo Verde Laboratory - division of Sonora Quest Lab

#### Lung Cancer Treatment Algorithmic Tests <sup>1</sup>

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6XS9G</b>	DetermaRX	Oncocyte
<b>7V98G</b>	LungOI	Imogene
<b>7VN6G</b>	PROphet NSCLC Test	OncoHost, Inc
<b>6UT9G</b>	VeriStrat	Biodesix

#### Lynch Syndrome / Hereditary Nonpolyposis Colorectal Cancer (HNPCC) Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6RPMG</b>	COLARIS	Myriad Genetics
<b>5HLGG</b>	Hereditary cancer Lynch syndrome	Fairview Diagnostic Laboratories
<b>2YYWG</b>	HNPCC concurrent	Ambry Genetics
<b>2FTJG</b>	HNPCC PANEL	Center for Human Genetics, Inc.
<b>7SNVG</b>	HNPCC/Lynch Deletion/Duplication	Ambry Genetics
<b>2FS8G</b>	HNPCC/Lynch Syndrome Del/Dup	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<b>2AWXG</b>	Invitae Lynch Syndrome Panel	Invitae Corporation
<b>2FSEG</b>	Lynch Syndrome - Hereditary (Germline) Testing	Labcorp   Oncology
<b>7QFCG</b>	Lynch Syndrome Focus (Sequencing & Deletion/Duplication)	Fulgent Genetics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2FRUG</b>	Lynch Syndrome Gene Panel: 5 Genes (EPCAM, MLH1, MSH2, MSH6, PMS2) by Gene Sequencing with Deletion/Duplication Analysis	Medical Diagnostic Laboratories, LLC
<b>7A2RG</b>	Lynch syndrome Panel	PerkinElmer Genomics
<b>57DNG</b>	Lynch syndrome Panel	PreventionGenetics, part of Exact Sciences
<b>2FRZG</b>	Lynch syndrome Panel	Quest Diagnostics
<b>5JURG</b>	Lynch syndrome Panel	The University of Chicago Genetic Services
<b>7AC3G</b>	Lynch Syndrome Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>7DL5G</b>	Lynch Syndrome Panel, Varies	Mayo Clinic Laboratories
<b>4JKFG</b>	Lynch Syndrome, MSH2 Sequencing and Deletion/Duplication (Including EPCAM)	Quest Diagnostics
<b>72H6G</b>	Lynch with RNAinsight	Ambry Genetics
<b>2FSKG</b>	MLH1 / MSH2 / MSH6 / PMS2 Comprehensive Analysis	Labcorp   Oncology
<b>2FSLG</b>	MLH1 / MSH2 / MSH6 Comprehensive Analysis	Labcorp   Oncology
<b>2FSNG</b>	MLH1 / MSH2 Comprehensive Analysis	Labcorp   Oncology
<b>2FSXG</b>	MLH1, MSH2, and MSH6 Sequencing and Deletion/Duplication Panel	Knight Molecular Diagnostic
<b>7DQLG</b>	MLH1/MSH2 Del/Dup Testing by MLPA	PerkinElmer Genomics
<b>2YYNG</b>	MSH2 seq & del/dup & EPCAM del/dup	Ambry Genetics
<b>7DQKG</b>	MSH6/MUTYH/EPCAM Del/Dup Testing by MLPA	PerkinElmer Genomics
<b>87B9G</b>	VistaSeq Lynch Syndrome Panel	LabCorp

#### Mitochondrial Disease (including Nuclear Genes) Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>5HL6G</b>	Cardioencephalomyopathy, fatal infantile, due to cytochrome c oxidase deficiency	Fairview Diagnostic Laboratories
<b>77W7G</b>	Cellular Energetics Ophthalmoplegia Syndromes (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
<b>5C94G</b>	Chronic Progressive External Ophthalmoplegia (CPEO/PEO) Panel	PreventionGenetics, part of Exact Sciences
<b>6DD7G</b>	Combined Mito Genome Plus Mito Focused Nuclear Gene Panel	GeneDx

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7R7VG</b>	Combined Mitochondrial Full Genome and Nuclear Gene Panel, Varies	Mayo Clinic Laboratories
<b>6RKZG</b>	Comprehensive Cellular Energetics Defects (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
<b>77WDG</b>	Comprehensive Mitochondrial Nuclear Gene Panel	PerkinElmer Genomics
<b>77SKG</b>	Comprehensive mtDNA Depletion Syndromes (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>76YBG</b>	Comprehensive Ophthalmoplegia Syndromes (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
<b>76YHG</b>	Cytochrome C Oxidase Deficiency (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
<b>788KG</b>	Dual Genome Leigh Disease Panel by Massively Parallel Sequencing	Baylor Genetics, LLC
<b>6LC5G</b>	Genomic Unity® Comprehensive Mitochondrial Disorders Analysis	Variantyx
<b>7V37G</b>	Genomic Unity® Mitochondrial Genome Deletions Analysis	Variantyx
<b>7V36G</b>	Genomic Unity® Nuclear Encoded Mitochondrial Gene Analysis	Variantyx
<b>4PU2G</b>	Initial Mitochondrial Evaluation	Athena Diagnostics Inc
<b>3XEDG</b>	Invitae Nuclear Mitochondrial Disorders Panel	Invitae Corporation
<b>77YFG</b>	Leigh and Leigh-Like Syndrome Panel (Nuclear Genes Only)	PreventionGenetics, part of Exact Sciences
<b>2YAYG</b>	Mito Disord Panel (mtDNA-108 Nuc Genes)	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<b>77FZG</b>	Mitochondrial complex I deficiency	Fairview Diagnostic Laboratories
<b>77GLG</b>	Mitochondrial Complex I Deficiency Panel (Nuclear Genes)	PreventionGenetics, part of Exact Sciences
<b>4J4CG</b>	Mitochondrial complex II deficiency	Fairview Diagnostic Laboratories
<b>6QVJG</b>	Mitochondrial Complex II Deficiency Panel	PreventionGenetics, part of Exact Sciences
<b>4HZRG</b>	Mitochondrial complex III deficiency	Fairview Diagnostic Laboratories
<b>386QG</b>	Mitochondrial Complex III Deficiency Panel (Nuclear Genes)	PreventionGenetics, part of Exact Sciences
<b>2CB4G</b>	Mitochondrial complex IV deficiency	Fairview Diagnostic Laboratories
<b>76ZNG</b>	Mitochondrial Complex IV Deficiency Panel (Nuclear Genes)	PreventionGenetics, part of Exact Sciences
<b>2CAVG</b>	Mitochondrial complex V (ATP synthase) deficiency	Fairview Diagnostic Laboratories
<b>5LQTG</b>	Mitochondrial Complex V Deficiency Panel (Nuclear Genes)	PreventionGenetics, part of Exact Sciences

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7U5TG</b>	Mitochondrial Disorders (mtDNA) Sequencing and Deletion Analysis by NGS	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>6LTZG</b>	Mitochondrial Disorders Panel (Nuclear Genes Only)	PreventionGenetics, part of Exact Sciences
<b>77N4G</b>	Mitochondrial DNA depletion syndrome	Fairview Diagnostic Laboratories
<b>2YB3G</b>	Mitochondrial DNA Depletion Syndrome Panel	Blueprint Genetics
<b>54EFG</b>	Mitochondrial Encephalomyopathic Evaluation	Athena Diagnostics Inc
<b>77FTG</b>	Mitochondrial Encephalopathy/Leigh Syndrome Nuclear Gene Panel	PerkinElmer Genomics
<b>772TG</b>	Mitochondrial Genome Maintenance/Integrity Nuclear Genes Panel	PreventionGenetics, part of Exact Sciences
<b>3WQPG</b>	Mitochondrial Hepatoencephalopathic Evaluation	Athena Diagnostics Inc
<b>3WSXG</b>	Mitochondrial Neurogastrointestinal Encephalopathy Evaluation (MNGIE)	Athena Diagnostics Inc
<b>4RDTG</b>	Mitochondrial Respiratory Chain Complex II Deficiency Panel by Massively Parallel Sequencing	Baylor Genetics, LLC
<b>77T2G</b>	Mitochondrial Respiratory Chain Complex I-V Nuclear Gene Deficiency Panel by Massively Parallel Sequencing	Baylor Genetics, LLC
<b>62P9G</b>	mtDNA Depletion Syndrome NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>77EVG</b>	mtDNA Depletion Syndrome NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77TJG</b>	mtDNA Depletion Syndrome NGS Panel (Sequencing Only)	Fulgent Genetics
<b>34PAG</b>	Multiple mitochondrial dysfunctions syndrome	Fairview Diagnostic Laboratories
<b>78DMG</b>	Non-mitochondrial Comprehensive Ophthalmoplegia Syndromes (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>76YUG</b>	Nuclear mitochondrial genes	Fairview Diagnostic Laboratories
<b>2TC7G</b>	Nuclear Panel by Massively Parallel Sequencing	Baylor Genetics, LLC
<b>2BKCG</b>	Nuclear-Mito NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>2BKFG</b>	Nuclear-Mito NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>2BKEG</b>	Nuclear-Mito NGS Panel (Sequencing Only)	Fulgent Genetics
<b>76USG</b>	Oxidative Phosphorylation (OXPHOS) Defects (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2BKHG</b>	Oxidative Phosphorylation Disorders NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>78H9G</b>	Oxidative Phosphorylation Disorders NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77K2G</b>	Oxidative Phosphorylation Disorders NGS Panel (Sequencing Only)	Fulgent Genetics
<b>6UV6G</b>	PEO Panel by Massively Parallel Sequencing	Baylor Genetics, LLC
<b>776AG</b>	PGXome Custom - Combined Oxidative Phosphorylation Deficiency	PreventionGenetics, part of Exact Sciences
<b>77KGG</b>	PGXome Custom - Comprehensive Cellular Energetics Defects	PreventionGenetics, part of Exact Sciences
<b>77GJG</b>	PGXome Custom - Mitochondrial Complex I Deficiency	PreventionGenetics, part of Exact Sciences
<b>4C5HG</b>	PGXome Custom - Mitochondrial Complex II Deficiency	PreventionGenetics, part of Exact Sciences
<b>6P7LG</b>	PGXome Custom - Mitochondrial Complex III Deficiency	PreventionGenetics, part of Exact Sciences
<b>78KNG</b>	PGXome Custom - Mitochondrial Complex IV Deficiency	PreventionGenetics, part of Exact Sciences
<b>6P7NG</b>	PGXome Custom - Mitochondrial Complex V Deficiency	PreventionGenetics, part of Exact Sciences
<b>77SZG</b>	Respiratory Chain Deficiency Panel	Knight Molecular Diagnostic
<b>7B35G</b>	STAT Comprehensive Mitochondrial Nuclear Gene Panel	PerkinElmer Genomics
<b>7B38G</b>	STAT Mitochondrial Encephalopathy/Leigh Syndrome Nuclear Gene Panel	PerkinElmer Genomics

### Mitochondrial DNA (mtDNA) Sequencing Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7SKYG</b>	Advanced mtDNA Point Mutations	LabCorp
<b>76WBG</b>	CentoMito® Comprehensive - NGS Panel (Mito Genome included)	Centogene
<b>79ASG</b>	CentoMito® Genome - Mito Genome	Centogene
<b>3KDTG</b>	CHOP MitoGenome Sequencing + Deletion Analysis	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>2Z9XG</b>	Complete Mitochondrial Sequencing (mtDNA)	Gene by Gene
<b>2DZ4G</b>	Comprehensive mtDNA Analysis by Massively Parallel Sequencing	Baylor Genetics, LLC
<b>79MYG</b>	Deafness, nonsyndromic, sensorineural, mitochondrial - Single Gene Sequencing by NGS	Centogene
<b>2LKVG</b>	Dual Genome Panel by Massively Parallel Sequencing	Baylor Genetics, LLC

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6UPNG</b>	Genomic Unity® Mitochondrial Genome Analysis	Variantyx
<b>7V38G</b>	Genomic Unity® Mitochondrial Genome Sequence Analysis	Variantyx
<b>39CMG</b>	Known mtDNA Variant(s) Testing by NGS-Urine-Test T822	GeneDx
<b>4AZZG</b>	Maternal MitoGenome Seq + Del	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>6U87G</b>	Maternal Relative Exome + MitoGenome Combined Test	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>6L9XG</b>	Mito Genome Sequencing & Deletion Testing	GeneDx
<b>7QBEG</b>	Mitochondrial Depletion NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>253UG</b>	Mitochondrial Disorders (mtDNA) Sequencing and Deletion Analysis by NGS	ARUP Laboratories
<b>72G9G</b>	Mitochondrial DNA Depletion Testing (Muscle)	LabCorp
<b>5D36G</b>	Mitochondrial DNA Sequencing	New Jersey Medical School - Institute of Genomic Medicine
<b>2G4RG</b>	Mitochondrial DNA Testing	Bionano Laboratories
<b>6R5TG</b>	Mitochondrial Full Genome Analysis, Next-Generation Sequencing (NGS), Varies	Mayo Clinic Laboratories
<b>4CH3G</b>	MITOCHONDRIAL GENOME SEQUENCING	Center for Human Genetics, Inc.
<b>7RXFG</b>	MITOCHONDRIAL GENOME SEQUENCING	LabCorp
<b>6R9BG</b>	MITOCHONDRIAL GENOME SEQUENCING	MNG Laboratories
<b>6R9DG</b>	Mitochondrial Genome Sequencing + Deletion Analysis	MNG Laboratories
<b>7RXEG</b>	Mitochondrial Genome Sequencing and Deletion Analysis	LabCorp
<b>2JZ5G</b>	Mitochondrial Genome Test	Blueprint Genetics
<b>28H4G</b>	Mitochondrial Myopathy mtDNA	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>2XYNG</b>	mtDNA Whole Genome Sequencing	Columbia University - Personalized Genomic Medicine
<b>27KAG</b>	MVL MitoSeq Panel	Molecular Vision Laboratory
<b>7R7XG</b>	Nuclear Mitochondrial Gene Panel, Next-Generation Sequencing, Varies	Mayo Clinic Laboratories
<b>6USKG</b>	PGmito - Mitochondrial Genome Sequencing	PreventionGenetics, part of Exact Sciences
<b>35JUG</b>	Rapid MitoGenome Seq + Del	Children's Hospital of Philadelphia - Division of Genomic Diagnostics

## Pan-Cancer Hereditary Cancer Panel Tests

GTU	Test Name	Laboratory Name
<b>7DQMG</b>	BRCA2/CHEK2 Del/Dup Testing by MLPA	PerkinElmer Genomics
<b>76DVG</b>	Cancer Panel	PreventionGenetics, part of Exact Sciences
<b>76DGG</b>	CancerNext	Ambry Genetics
<b>6UR6G</b>	CancerNext with RNAinsight	Ambry Genetics
<b>7TP4G</b>	CancerNext Expanded HBOC	Ambry Genetics
<b>7TP2G</b>	CancerNext: HBOC	Ambry Genetics
<b>76E2G</b>	CancerNext-Expanded	Ambry Genetics
<b>6UR7G</b>	CancerNext-Expanded with RNAinsight	Ambry Genetics
<b>5BTUG</b>	CentoCancer - NGS Panel (CNV included)	Centogene
<b>5APDG</b>	CentoCancer comprehensive panel - NGS Panel (CNV included)	Centogene
<b>7C5QG</b>	Color Extended	Color Genomics
<b>7C5RG</b>	Color Standard	Color Genomics
<b>76E5G</b>	Common Cancer Management Panel	GeneDx
<b>7V2AG</b>	Common Hereditary Cancer Panel	Baylor Genetics, LLC
<b>7RX8G</b>	Common Hereditary Cancer Screening Panel	PreventionGenetics, part of Exact Sciences
<b>76BTG</b>	COMP CANCER PANEL	ACL Laboratories
<b>76CKG</b>	Comprehensive Cancer Panel	PerkinElmer Genomics
<b>76DQG</b>	Comprehensive Common Cancer Panel	BioReference Laboratories
<b>76C3G</b>	Comprehensive Common Cancer Panel: HBOC	GeneDx
<b>5FV7G</b>	Comprehensive Hereditary Breast and Gynecologic Cancer Panel: 18 genes (BRCA1/2, High Risk Extended Panel and Lynch Syndrome genes) by Gene Sequencing with BRCA1/2, EPCAM, MLH1, MSH2, MSH6, PMS2 Deletion/Duplication Analysis	Medical Diagnostic Laboratories, LLC
<b>7V28G</b>	Comprehensive Hereditary Cancer Panel	Baylor Genetics, LLC
<b>76CDG</b>	Comprehensive Hereditary Cancer Panel	Blueprint Genetics
<b>5K7ZG</b>	Comprehensive Hereditary Cancer Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>262EG</b>	Comprehensive Hereditary Cancer Panel	Quest Diagnostics
<b>7BP4G</b>	Comprehensive Hereditary Cancer Panel	The University of Chicago Genetic Services

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>77HHG</b>	Comprehensive Pan-cancer analysis (DNA and RNA analysis)	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>76DHG</b>	CustomNext-Cancer	Ambry Genetics
<b>6UR2G</b>	CustomNext-Cancer with RNAinsight	Ambry Genetics
<b>76CAG</b>	Empower - Multi-cancer expanded Hereditary Cancer Test	Natera
<b>76C5G</b>	Empower - Multi-cancer Hereditary Cancer Test	Natera
<b>4M9NG</b>	Full Comprehensive Cancer Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>6U89G</b>	Full Comprehensive Cancer Panel (Germline)	NeoGenomics Laboratories
<b>76CUG</b>	Full Comprehensive Cancer Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>4U95G</b>	Full Comprehensive Cancer Panel (Sequencing Only)	Fulgent Genetics
<b>3HLKG</b>	Full Focus Cancer Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>6U8AG</b>	Full Focus Cancer Panel (Germline)	NeoGenomics Laboratories
<b>76DMG</b>	Full Focus Cancer Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>4CCHG</b>	Full Focus Cancer Panel (Sequencing Only)	Fulgent Genetics
<b>25XXG</b>	Guideline Based Hereditary Cancer Panel	Quest Diagnostics
<b>6URZG</b>	HC COMMON HEREDITARY CANCER PANEL	UCSF Molecular Diagnostics Laboratory
<b>76BWG</b>	Hereditary Breast and Ovarian Cancer - Expanded and Lynch Syndrome Panel	PreventionGenetics, part of Exact Sciences
<b>76DRG</b>	Hereditary Cancer	Color Genomics
<b>76CCG</b>	Hereditary Cancer High Risk Panel	Blueprint Genetics
<b>7VAGG</b>	Hereditary Cancer NGS Panel	AiLife Diagnostics
<b>7VAHG</b>	Hereditary Cancer NGS Panel Rapid	AiLife Diagnostics
<b>76DXG</b>	Hereditary Cancer Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>7DLHG</b>	Hereditary Common Cancer Panel, Varies	Mayo Clinic Laboratories
<b>7DKGG</b>	Hereditary Expanded Cancer Panel, Varies	Mayo Clinic Laboratories
<b>76CMG</b>	High/mod Risk Panel (20 genes)	ACL Laboratories
<b>7VVFG</b>	HopeSeq Germline Confirmatory Panel (Hope Hereditary Cancer Predisposition Panel)	City of Hope National Medical Center - Molecular Diagnostic Laboratory
<b>76DKG</b>	Inherited Cancer Panel	Knight Molecular Diagnostic

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>76CHG</b>	Integrated BRACAnalysis with Myriad myRisk Hereditary Cancer Update Test	Myriad Genetics
<b>77BJG</b>	Invitae Cancer Screen	Invitae Corporation
<b>7UZRG</b>	Invitae Common Hereditary Cancers + RNA Panel	Invitae Corporation
<b>7DGXG</b>	Invitae Common Hereditary Cancers + RNA Panel-Common Hereditary Cancers Genes Eligible for RNA Analysis	Invitae Corporation
<b>76BNG</b>	Invitae Common Hereditary Cancers Panel	Invitae Corporation
<b>7UZSG</b>	Invitae Multi-Cancer + RNA Panel	Invitae Corporation
<b>7AV8G</b>	Invitae Multi-Cancer + RNA Panel-Multi-Cancer Genes Eligible for RNA Analysis	Invitae Corporation
<b>76BMG</b>	Invitae Multi-Cancer Panel	Invitae Corporation
<b>76BUG</b>	Myriad myRisk	Myriad Genetics
<b>6RMUG</b>	myRisk Update Test	Myriad Genetics
<b>7DQGG</b>	NxGen Hereditary Cancer Panel	NxGen MDx
<b>76CWG</b>	OncoGene Dx Custom Panel	GeneDx
<b>7V3LG</b>	OnkoRisk Hereditary Oncology Guideline Panel (Non-NYS)	BioReference Laboratories
<b>7V3MG</b>	OnkoRisk Hereditary Oncology Management Panel (Non-NYS)	BioReference Laboratories
<b>7V3KG</b>	OnkoRisk Hereditary Oncology Plus Panel (Non-NYS)	BioReference Laboratories
<b>78FJG</b>	Pan-cancer DNA Analysis	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>2TQXG</b>	PrevenTest	Advanced Molecular Diagnostics, LLC
<b>74G2G</b>	Rest of Comprehensive Common Cancer Panel: HBOC	GeneDx
<b>7RVTG</b>	Riskguard	Exact Sciences Laboratories, LLC
<b>7B7FG</b>	STAT Comprehensive Cancer Panel	PerkinElmer Genomics
<b>7TS6G</b>	Targeted Oncology Panel Next Generation Sequencing Bone Marrow	Cleveland Clinic Laboratories
<b>7TS7G</b>	Targeted Oncology Panel Next Generation Sequencing Cytology	Cleveland Clinic Laboratories
<b>7TS9G</b>	Targeted Oncology Panel Next Generation Sequencing Other	Cleveland Clinic Laboratories
<b>7VPJG</b>	Tempus xG Common Hereditary Cancers 36 genes (all other hereditary cancer indications)	Tempus AI, Inc.
<b>7VPFG</b>	Tempus xG+ Extended Hereditary Cancers 77 genes (all other hereditary cancer indications)	Tempus AI, Inc.

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>76D8G</b>	VistaSeq Hereditary Cancer Panel	Integrated Genetics
<b>76BXG</b>	VistaSeq Hereditary Cancer Panel	LabCorp
<b>76BRG</b>	VistaSeq Hereditary Cancer Panel	Labcorp   Oncology

### Pharmacogenetic Neuropsychiatric Panel Tests <sup>1</sup>

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7T3YG</b>	Comprehensive Pharmacogenetics Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>2FGMG</b>	Drug Metabolizing Enzymes Panel (CYP2D6, CYP2C9, CYP2C19), DNA analysis	Shodair Children's Hospital - Genetics Laboratory
<b>7VN9G</b>	EffectiveRx Neuropsychiatric Panel	GENETWORx
<b>2AUXG</b>	GeneSight Psychotropic	Myriad Genetics
<b>5YRYG</b>	GeneSight Psychotropic	Myriad Neuroscience
<b>33Y2G</b>	Genetic Pharmacology Psychiatry Drug Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>6S83G</b>	Genomind PGx Test	Genomind
<b>7S9DG</b>	IDgenetix	Castle Biosciences Inc
<b>7URYG</b>	PGx: Mental Health Panel	Helix
<b>7QENG</b>	Pharmacogenetics Panel: Psychotropics	ARUP Laboratories
<b>7UNVG</b>	Pharmacogenetics Panel: Psychotropics, with GeneDose Access	ARUP Laboratories
<b>7PRTG</b>	Psych HealthPGx Panel	RPRD Diagnostics LLC
<b>6UAFG</b>	Psychotropic Pharmacogenomics Gene Panel, Varies	Mayo Clinic Laboratories
<b>3WG4G</b>	SureGene	Clinical Reference Laboratory
<b>3HHG</b>	Tempus nP	Tempus AI, Inc.

### Prostate Cancer Prognostic Algorithmic Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7BN5G</b>	Decipher Prostate Genomic Classifier	Veracyte
<b>7Q8UG</b>	Oncotype DX Genomic Prostate Score Assay	MDx Health
<b>4SVPG</b>	Oncotype DX Prostate Cancer Assay	University of Minnesota Physicians Outreach Laboratory
<b>6UUUG</b>	Polaris	Myriad Genetics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6UU7G</b>	Polaris Biopsy	Myriad Genetics

### Thyroid Cancer Diagnostic Algorithmic Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2YBRG</b>	Afirma Genomic Sequencing Classifier	Veracyte
<b>7S8XG</b>	Afirma Xpression Atlas	Veracyte
<b>6LJEG</b>	Cytology + Reflex to ThyGeNEXT only	Interpace Diagnostics
<b>6LJCG</b>	Cytology + Reflex to ThyGeNEXT w/Reflex to ThyraMIR	Interpace Diagnostics
<b>6UW7G</b>	ThyGeNEXT only	Interpace Diagnostics
<b>6LJDG</b>	ThyGeNEXT w/ Reflex to ThyraMIR	Interpace Diagnostics
<b>6UUFG</b>	ThyroSeq	CBLPath
<b>6UTYG</b>	ThyroSeq - Thyroid Cancer Next-Generation Sequencing Panel	University of Pittsburgh Medical Center - Division of Molecular Diagnostics

### TPMT and NUDT15 Typing Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>33HDG</b>	NT Panel (NUDT15, TPMT)	RPRD Diagnostics LLC
<b>48S9G</b>	PGX TPMT and NUDT15 Genotyping	Indiana University Molecular Genetics Diagnostic Laboratory - Indiana University Medical Center, School of Medicine
<b>6RA6G</b>	Thiopurine Methyltransferase (TPMT) and Nudix Hydrolase (NUDT15) Genotyping, Varies	Mayo Clinic Laboratories
<b>3ADCG</b>	TPMT and NUDT15	ACL Laboratories
<b>7C3XG</b>	TPMT and NUDT15	Allina Health Laboratory
<b>2533G</b>	TPMT and NUDT15	ARUP Laboratories
<b>7BEHG</b>	TPMT and NUDT15	Clinical Pathology Laboratories
<b>34EWG</b>	TPMT and NUDT15	University of California Davis Health System - UCDMC - Molecular and Cytogenetic Laboratory
<b>7U5AG</b>	TPMT and NUDT15	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>79YEG</b>	TPMT and NUDT15 Genotype	ProPath
<b>7TRSG</b>	TPMT and NUDT15 Genotyping	LabCorp
<b>34EKG</b>	TPMT and NUDT15 Genotyping	Stanford Clinical Laboratories - Biochemical Genetics Laboratory

GTU	Test Name	Laboratory Name
2KQGG	TPMT and NUDT15 Genotyping	University of North Carolina Hospitals - Molecular Genetics

### Tumor Specific BCR/ABL Quantitation and Breakpoint Analysis Tests

GTU	Test Name	Laboratory Name
2YTJG	BCR / ABL - t(9;22) by RT-PCR (Quantitative)	Labcorp   Oncology
3LU5G	BCR ABL P210 QUANT	SUNY Upstate Medical University
7BQPG	BCR/ABL by FISH	Allina Health Laboratory
6ND3G	BCR/ABL D-FISH	Center for Genetic Testing at Saint Francis
3KP8G	BCR/ABL qRT PCR, Major p210 and Minor p190 Mutations, Minimal Residual Disease	Molecular Pathology Laboratory Network
2V82G	BCR/ABL Quantification by RT-PCR	BioReference Laboratories
6YXMG	BCR/ABL t(9;22) major (p210) IS Quantitative	Molecular Diagnostic Laboratory - Barnes Jewish Hospital
6LE9G	BCR/ABL1 (T(9;22)) RNA Quantitative with Interpretation	University of Iowa Hospitals and Clinics - Department of Pathology
6UV4G	BCR/ABL1 (T(9;22)) RNA Quantitative with Interpretation	University of Iowa Hospitals and Clinics - Department of Pathology
7VVVG	BCR/ABL1 Kinase Mutation Analysis by NGS	City of Hope National Medical Center - Molecular Diagnostic Laboratory
5SUVG	BCR/ABL1 p190 Quantitative PCR Blood	Cleveland Clinic Laboratories
5UA8G	BCR/ABL1 p190 Quantitative PCR Bone Marrow	Cleveland Clinic Laboratories
7TH5G	BCR/ABL1 p210 and p190 Diagnostic PCR Blood	Cleveland Clinic Laboratories
7TH6G	BCR/ABL1 p210 and p190 Diagnostic PCR Bone Marrow	Cleveland Clinic Laboratories
5UAAAG	BCR/ABL1 p210 Quantitative PCR Blood	Cleveland Clinic Laboratories
6LQRG	BCR/ABL1 p210 Quantitative PCR Bone Marrow	Cleveland Clinic Laboratories
5PH3G	BCR-ABL Diagnostic, Blood	Johns Hopkins Medical Institutions - Pathology Laboratory
4GZ7G	BCR-ABL Diagnostic, Bone Marrow	Johns Hopkins Medical Institutions - Pathology Laboratory
5W36G	BCR-ABL Mutation	University of Illinois at Chicago - Biochemical Genetics Laboratory
5PGYG	BCR-ABL p190 Followup, Blood	Johns Hopkins Medical Institutions - Pathology Laboratory
4GLHG	BCR-ABL p190 Followup, Bone Marrow	Johns Hopkins Medical Institutions - Pathology Laboratory
4AGMG	BCR-ABL p210 Followup, Blood	Johns Hopkins Medical Institutions - Pathology Laboratory
3AUMG	BCR-ABL p210 Followup, Bone Marrow	Johns Hopkins Medical Institutions - Pathology Laboratory
723HG	BCR-ABL1 Gene Rearrangement, Quantitative, PCR	Quest Diagnostics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>8E7CG</b>	BCR-ABL1 Major (p210) by Quantitative RT-PCR for Monitoring	Medfusion
<b>8E7GG</b>	BCR-ABL1 Major (p210) by Quantitative RT-PCR with Reflex to Minor (p190) for Baseline Quantitation	Medfusion
<b>2ZTAG</b>	BCR-ABL1 Non-Standard p230	NeoGenomics Laboratories
<b>7BDCG</b>	BCR-ABL1 QUAL, RFLX QUANT, NEW DIAGNOSIS	Clinical Pathology Laboratories
<b>7AEXG</b>	BCR-ABL1 QUAL, RFLX QUANT, NEW DIAGNOSIS, WHOLE BLOOD	Clinical Pathology Laboratories
<b>7AF2G</b>	BCR-ABL1 QUANT, MAJOR, WHOLE BLOOD	Clinical Pathology Laboratories
<b>7AEZG</b>	BCR-ABL1 QUANT, MINOR, WHOLE BLOOD	Clinical Pathology Laboratories
<b>73BFG</b>	BCR-ABL1 Standard p210, p190	NeoGenomics Laboratories
<b>6X7DG</b>	BCR-ABL1 Transcript Detection for Chronic Myelogenous Leukemia (CML) and Acute Lymphocytic Leukemia (ALL), Quantitative	LabCorp
<b>86NEG</b>	BCR-ABL1, Major (p210), Quantitative	Molecular Diagnosis - University of Rochester Medical Center - Strong Memorial Hospital
<b>7BDEG</b>	BCR-ABL1, QUANT, CML MONITORING (p210)	Clinical Pathology Laboratories
<b>3GCEG</b>	BCR-ABL1: High Sensitivity Major Breakpoint p210	University of Minnesota Physicians Outreach Laboratory
<b>5YZ4G</b>	FISH (9_22)	SUNY Upstate Medical University
<b>6VXDG</b>	QUANTITATIVE DETECTION OF BCR-ABL1 MAJOR FORM (p210)	University of California Davis Health System - UC DMC - Molecular and Cytogenetic Laboratory

#### Tumor-Type Agnostic Solid Tumor Molecular Profiling Panel Tests (51 or more genes)

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>86SBG</b>	Altera Tumor Profiling	Natera
<b>3CQFG</b>	Columbia Combined Cancer Panel (CCCP)	Columbia University - Personalized Genomic Medicine
<b>2CV6G</b>	Comprehensive NGS Solid Tumor Mutation Panel	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<b>2D5CG</b>	Comprehensive Solid Tumor (DNA and RNA analysis)	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>72Y9G</b>	Endeavor Comprehensive Solid Tumor Profile	PathGroup
<b>6UVBG</b>	FoundationOne	University of Minnesota Physicians Outreach Laboratory
<b>6R4AG</b>	FoundationOne CDx	Foundation Medicine
<b>7UJGG</b>	Fusion Transcript Panel	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>3V7ZG</b>	GatorSeq (NGS Panel)	University of Florida - Health Pathology Laboratories
<b>72MCG</b>	GeneTrails Comprehensive Solid Tumor Panel	Knight Molecular Diagnostic
<b>6YSYG</b>	GeneTrails Solid Tumor Panel with Reflex to Solid Tumor Fusion Gene Panel	Knight Molecular Diagnostic
<b>86PJV</b>	Guardant360 TissueNext	Guardant Health
<b>6ZQMG</b>	Iowa Cancer Mutation and RNA Fusion Profile and Interpretation	University of Iowa Hospitals and Clinics - Department of Pathology
<b>7AWEG</b>	MayoComplete Solid Tumor Panel, Next-Generation Sequencing, Tumor	Mayo Clinic Laboratories
<b>6WUTG</b>	MI Tumor Seek Hybrid	Caris Life Sciences
<b>6UUMG</b>	MSK-IMPACT	Memorial Sloan Kettering Cancer Center
<b>7SSDG</b>	Neo Comprehensive - Solid Tumor	NeoGenomics Laboratories
<b>7529G</b>	NeXT Dx™	Personalis
<b>2Z49G</b>	Next Generation Sequencing Oncology Tumor Testing	University of Minnesota Physicians Outreach Laboratory
<b>34WBG</b>	NGS Solid Tumor Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>7V34G</b>	OncоАly Solid Tumor Analysis	Variantyx
<b>2YA6G</b>	OncоАxOne	Admera Health
<b>7BNDG</b>	Oncomap ExTra	Exact Sciences Laboratories, LLC
<b>76ETG</b>	Oncoplex Select Cancer Gene Panel	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>76EQG</b>	Oncoplex Select Director	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>76EYD</b>	Oncoplex Select Interpretation	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>76EZG</b>	Oncoplex Select Methods	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>76ESG</b>	Oncoplex Select Panel	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>76ERG</b>	Oncoplex Select Result	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>76EXG</b>	Oncoplex Select Tested Sample	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>7UUYG</b>	OnkoSight Advanced 523 Gene NGS Panel	BioReference Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7SSVG</b>	OnkoSight Advanced 523 Gene NGS Reanalysis with PierianDx	BioReference Laboratories
<b>7URPG</b>	OnkoSight Advanced 523 Gene NGS with PierianDx Interpretation	BioReference Laboratories
<b>2FLHG</b>	Paired Tumor/Normal – Comprehensive Solid Tumor Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>2FLGG</b>	Paired Tumor/Normal – Solid Tumor Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7TKLG</b>	Pan-Cancer Solid Tumor NGS Panel	UCLA Diagnostic Molecular Pathology Laboratory
<b>6V93G</b>	PCMP - Personalized Cancer Mutation Panel	University of Pittsburgh Medical Center - Division of Molecular Diagnostics
<b>7UJFG</b>	PennSeq Solid Tumor Panel	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>2K9WG</b>	PGDx elio tissue complete	Personal Genome Diagnostics
<b>7A5HG</b>	Precise Tumor	Myriad Genetics
<b>2DSFG</b>	RNA Fusion Analysis	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>7ATMG</b>	SOLID TUMOR COMPREHENSIVE PANEL (Combined solid tumor mutation and fusion panels)	Texas Children's Hospital
<b>2D5BG</b>	Solid Tumor DNA Analysis	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>7SVMG</b>	Solid Tumor Expanded Panel	Quest Diagnostics
<b>6QZNG</b>	Solid Tumor Fusion Analysis	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>2ARCG</b>	Solid Tumor Gene Set	Washington University in St. Louis Genomics and Pathology Services
<b>3CSSG</b>	Solid Tumor Molecular Profile	Fulgent Genetics
<b>2NSPG</b>	Solid Tumor NGS Panel	University of Michigan - Michigan Medical Genetics Laboratories
<b>2FRPG</b>	Solid Tumor Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>6UWNG</b>	Solid tumor panel - Somatic Mutation Analysis	Centogene
<b>2LEVG</b>	Solid Tumor Targeted Cancer Gene Panel by Next-Generation Sequencing, NGS	Medical College of Wisconsin - Human and Molecular Genetics Center
<b>7C3KG</b>	Solid tumor targeted NGS panel	Allina Health Laboratory
<b>7UQMG</b>	Solid TumorSEQ Expanded Panel	Medfusion
<b>2CPUG</b>	Strata Select	Strata Oncology
<b>7TXFG</b>	Tempus xT (tumor only), Paraffin Block	University of Michigan - Michigan Medical Genetics Laboratories
<b>7TXHG</b>	Tempus xT Targeted Panel (Tumor plus Blood)	University of Michigan - Michigan Medical Genetics Laboratories
<b>7TXJG</b>	Tempus xT Targeted Panel (Tumor plus Saliva)	University of Michigan - Michigan Medical Genetics Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7VPEG</b>	Tempus  xT CDx: FDA-approved Companion Diagnostic	Tempus AI, Inc.
<b>86SDG</b>	Tempus  xT: Targeted panel of 648 genes (Solid Tumor Only)	Tempus AI, Inc.
<b>6RWJG</b>	TempusTM Test	ACL Laboratories
<b>6LS8G</b>	UW OncoPlex Cancer Gene Panel	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>5LYGG</b>	UW OncoPlex Single Gene	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory

#### Tumor-Type Agnostic Solid Tumor Molecular Profiling Panel Tests (5-50 genes)

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2DWYG</b>	Cancer Mutation 50-Gene Panel, Next Generation Sequencing	Veripath Laboratories
<b>2LVRG</b>	Cancer Mutation Profiling and Interpretation	University of Iowa Hospitals and Clinics - Department of Pathology
<b>2LCCG</b>	Cancer Mutation Profiling Blood Paired Normal Testing	University of Iowa Hospitals and Clinics - Department of Pathology
<b>2XYRG</b>	Columbia Solid Tumor (CSTP) Subpanels	Columbia University - Personalized Genomic Medicine
<b>5BZJG</b>	Columbia Solid Tumor Panel (CSTP)	Columbia University - Personalized Genomic Medicine
<b>7VQWG</b>	Comprehensive Solid Tumor Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7BZJG</b>	GIST targeted NGS panel	Allina Health Laboratory
<b>5BRWG</b>	NGS Solid Tumor Hotspot Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>2V3FG</b>	OncoSeq - NGS Panel for Solid Tumors	University of Pittsburgh Medical Center - Division of Molecular Diagnostics
<b>86LTG</b>	OnkoSight Advanced Solid Tumor NGS Panel	BioReference Laboratories
<b>4ZPLG</b>	Solid Tumor 15 Genes Panel by NGS	Beaumont Laboratories - Molecular Pathology Lab
<b>2E5TG</b>	Solid Tumor Core Panel	Quest Diagnostics
<b>6VAPG</b>	Solid Tumor Mutation Panel	Ohio State University - Molecular Pathology Laboratory
<b>6V3PG</b>	Solid Tumor Mutation Panel 15	ACL Laboratories
<b>7DK3G</b>	Solid Tumor Mutation Panel, Sequencing	ARUP Laboratories
<b>7PVTG</b>	Solid Tumor Precision Panel	Nebraska Medical Center - Molecular Diagnostic Laboratory
<b>7UQNG</b>	Solid TumorSEQ Core Panel	Medfusion

## Tumor-Type Agnostic Molecular Profiling Panels Tests with IHC and Cytogenetic Analyses

GTU	Test Name	Laboratory Name
<b>7VVCG</b>	HopeSeq FNA Comprehensive Panel	City of Hope National Medical Center - Molecular Diagnostic Laboratory
<b>7VVDG</b>	HopeSeq Solid Tumors Comprehensive Panel	City of Hope National Medical Center - Molecular Diagnostic Laboratory
<b>7SS9G</b>	Lumera NGS Profile	Fulgent Genetics
<b>7SSAG</b>	Lumera Xpanded Profile	Fulgent Genetics
<b>7SPFG</b>	MI Tumor Seek Hybrid + IHCs and Other Tests by Tumor Type	Caris Life Sciences
<b>6WRQG</b>	NeoTYPE® Discovery Profile for Solid Tumors	NeoGenomics Laboratories
<b>74TLG</b>	NeoTYPE® Precision Profile for Solid Tumors	NeoGenomics Laboratories
<b>6Y7FG</b>	OmniSeq Advance Assay	Labcorp   Oncology
<b>87BHG</b>	OmniSeq INSIGHT	LabCorp
<b>78NMG</b>	OmniSeq INSIGHT	OmniSeq, Inc

## Uveal Melanoma Prognostic Algorithmic Tests

GTU	Test Name	Laboratory Name
<b>6UTUG</b>	DecisionDx-UM	Castle Biosciences Inc
<b>3BSZG</b>	Uveal Melanoma Prognostic	Cleveland Clinic Laboratories
<b>2BLLG</b>	Uveal Melanoma Prognostic Test	LabCorp

## X-Linked Intellectual Disability Panel Tests <sup>1</sup>

GTU	Test Name	Laboratory Name
<b>2FM3G</b>	90-gene XLID Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>4ABZG</b>	FG syndrome	Fairview Diagnostic Laboratories
<b>7V35G</b>	Genomic Unity® X-linked Intellectual Disability Plus Analysis	Variantyx
<b>77WRG</b>	Intellectual disability, X linked	Fairview Diagnostic Laboratories
<b>4XMCG</b>	PANEL 2 (ATPGAP2, OPHN1, SLC6A8, SYN1) SEQUENCING ONLY	Center for Human Genetics, Inc.
<b>4SMHG</b>	PANEL 2 MLPA (OPHN1, SLC6A8)	Center for Human Genetics, Inc.
<b>78KLG</b>	PGXome Custom - Intellectual Disability, X-linked	PreventionGenetics, part of Exact Sciences
<b>23XZG</b>	XLID (X-Linked Intellectual Disability) NGS Panel (Deletion/Duplication Only)	Fulgent Genetics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>5DSSG</b>	XLID (X-Linked Intellectual Disability) NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>34N9G</b>	XLID (X-Linked Intellectual Disability) NGS Panel (Sequencing Only)	Fulgent Genetics
<b>4S99G</b>	XLID TIER 1 SEQUENCING	Center for Human Genetics, Inc.
<b>3KWVG</b>	XLID TIER 2 MLPA	Center for Human Genetics, Inc.
<b>3KMHG</b>	XLID TIER 2 SEQUENCING	Center for Human Genetics, Inc.
<b>5YXXG</b>	XLID TIER 3 MLPA ONLY	Center for Human Genetics, Inc.
<b>2BSYG</b>	XLID TIER 3 SEQUENCING ONLY	Center for Human Genetics, Inc.
<b>5XPVG</b>	XLID TIER 4 MLPA	Center for Human Genetics, Inc.
<b>55YDG</b>	XLID TIER 4 SEQUENCING ONLY	Center for Human Genetics, Inc.
<b>2BYZG</b>	X-linked Intellectual Disability (NGS Panel and Copy Number Analysis + Fragile X Repeat Expansion & Methylation)	MNG Laboratories
<b>33HUG</b>	X-Linked Intellectual Disability (XLID) NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>785RG</b>	X-Linked Intellectual Disability Gene Analysis	University of Minnesota Physicians Outreach Laboratory
<b>2YACG</b>	X-linked Intellectual Disability Panel	Blueprint Genetics
<b>4YJGG</b>	X-linked Intellectual Disability Panel	PreventionGenetics, part of Exact Sciences
<b>7727G</b>	X-Linked Non-Specific Intellectual Disability Panel	The University of Chicago Genetic Services

<sup>1</sup>Category and tests therein are considered experimental and investigational (E/I).